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# Identification of key regulators for grapevine ripening by integrated genetic approaches 

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## Identification of key

## regulators for grapevine ripening by integrated genetic approaches

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## Summary

Grapevine is one of the most economically relevant fruit crops in the world. In the past few years climate change and in particular global warming have started to significantly impact agriculture and viticulture as well. Indeed, higher temperatures occurring early during the growing season may advance the date of the phenological stages like bud burst, flowering and veraison. As a consequence, the shift in veraison to earlier dates causes ripening to occur under higher temperatures which impairs wine quality. It is of great interest for viticulture to investigate the timing of veraison and disentangle its genetic control. Genetic variation between cultivars leads to differences in the expression of phenotypic traits such as veraison time. A deeper knowledge of the genes involved in the phenology processes is critical to select varieties that are well adapted for current and future climatic conditions.

The aim of this thesis was to identify the most promising candidate genes for the regulation of the transition to grapevine berry ripening through complementary genetic study. The thesis is divided in two chapters: in the first chapter an approach based on the integration of data from QTL studies has been developed, in order to narrow down the number of candidate genes that may be related with berry ripening. Meta-QTL analysis integrated with transcriptomic data led to the identification of 61 candidate genes related to veraison, including well known transcription factors and genes related to carbohydrate metabolism.

In the second part, we have characterized a wide germplasm collection of different grapevine accessions (from CREA-VIT Conegliano collection) with the aim to identify the best panels of grapevine varieties, representing the most diverse genotypes and phenotypes for the traits of interest, to be exploited to test the genotype-phenotype association. The Conegliano collection has been previously phenotyped for flowering time and veraison time in the last 50 years; our work focused on the phenotypic data of the last 13 years (data kindly made available by our partners). These phenotypic data have been firstly used to assist the construction of a core collection, including the most genetic diverse accessions and individuals with contrasting phenotypes. Approximately 600 CREA-VIT accessions were previously genotyped (Cipriani et al., 2010) using 45 microsatellite markers. The
genotypic data have been used to assess the genetic diversity and population structure of the Conegliano collection. The end of the second part was dedicated to performing the association analysis itself with two different approaches, firstly using the GrapeReSeq 18K Vitis genotyping chip, then with an innovative approach called XP-GWAS (Yang et al., 2015) where pool of individuals displaying extreme phenotypes are whole-genome resequenced and allele frequencies compared to a random pool are screened for enrichment.

The findings of this study provide indications of the genetic factors controlling or influencing veraison time in grapevine. The elucidation of the genetic network underlying the beginning of the berry ripening phase is of fundamental importance for the need to breed new grapevine varieties adapted to changing climatic conditions.

## Background

## Vitis vinifiera L.

Grapevine belongs to the family Vitaceae, a family of fourteen genera and about 910 known species (Christenhusz and Byng, 2016), covering mainly woody or herbaceous lianas (Mullins et al., 1992). Vitis is the most common genus of the family and contains 79 species. Vitis species are mainly spread in temperate zones of the Northern hemisphere and the same number of species is almost found between America and Asia. Only one Vitis species is native to Europe, Vitis vinifera L., which consists of two subspecies, Vitis vinifera L. subspecies sylvestris, the wild grapevine, and Vitis vinifera L. subspecies vinifera, the cultivated grapevine (Boss et al., 2003). The wild species Vitis vinifera sylvestris is the progenitor of the cultivated one, characterized by many large, sweet berries (Duchene, 2016). As a result, Vitis vinifera vinifera (Vitis vinifera from now on) is hermaphroditic while Vitis vinifera sylvestris is dioecious. Considering cultivation area (about 7.5 million ha in 2014) and economic value, grapevine is one of the major fruit crops of the world. The centre of domestication for grapevine is considered the near East between the Black Sea and Iran (Terral et al., 2010). From there, the first cultivated forms were diffused by humans probably throughout the middle and near East up to central Europe. To date, all the most known grapevine varieties, such as Pinot, Cabernet Sauvignon or Sangiovese, derives from Vitis vinifera. More than 74 million tons of grapes are produced annually, with the greatest portion in Europe. The usages of the grapes are various (fresh as table grapes, dried as raisins, juice, jelly, jam, ethanol, vinegar, grape seed oil, tartaric acid, and fertilizer), however, the most common product is wine ( $53 \%$ of the world's yield of grape). Vitis vinifera is hence one of the most important cultivated plants of the whole world.

## The genome of Vitis vinifera

Vitis vinifera is a diploid plant with 38 chromosomes ( $\mathrm{n}=19$ ). It has a genome size of approximately $475-500 \mathrm{Mb}$ (Jaillon et al., 2007) and about 32,000 genes (Vitulo et al., 2014). Since fertilization is occurring mainly by means of wind and insect
pollination, all grapevine varieties are highly heterozygous. Inbreeding depression is also a particular feature of grapevine, with sterility occurring often from the first to third generation of selfing. A reference genome sequence is available since 2007 from Jaillon et al.; this sequence is derived from a cultivated clone of Vitis vinifera Pinot Noir (PN40024), which from recurrent selfings has reached about a $93 \%$ of homozygosity. The Vitis genome sequence was firstly created from a Sanger sequencing assembly at an average 8 -fold coverage of the genome, later updated with a 12X assembly in 2010 when additional sequences were added to the pool (http://www.genoscope.cns.fr/externe/GenomeBrowser/Vitis/). The PN4002412 Xv 0 assembly, available on the website of the international Grape Genome Browser Genoscope, was improved and updated to the 12 Xv 2 assembly by the French-Italian Public Consortium (Canaguier et al., 2017) but the sequences of contigs and scaffolds remained unchanged compared to version 12Xv0. Several different gene annotations derived from different gene predictions are available at present. One gene set derived from PN40024-12Xv0 is available since 2009 and can be found on the Genoscope website. The gene prediction 12Xv1 is the result of the union of v0 and a gene prediction performed at the Centro di Ricerca Interdipartimentale per le Biotecnologie Innovative (CRIBI) in Padova, Italy and is available on the CRIBI website (http://genomes.cribi.unipd.it/grape/). In 2014 an improved gene annotation, called v2 was established on the 12 Xv 0 assembly version at the CRIBI (Vitulo et al., 2014). Recently, with the advent of new sequencing technologies based on the so-called "long reads", the area of the plant genomics has started to take advantage of these new approaches, and few grapevine genome announcements based on it have been made (Chin et al., 2016; Roach et al., 2018).

## The reproductive developmental cycle of grapevine

Vitis vinifera is a polycarpic perennial plant that cycles between periods of flowering and vegetative growth. Its entire reproductive developmental cycle is completed over two consecutive growing seasons, separated by dormancy (Carmona et al., 2008). Both vegetative and reproductive tissues are formed simultaneously on the same shoot by the shoot apical meristem. During the spring
of the first season, lateral meristems, called uncommitted primordia, are formed within latent buds in the axis of grapevine leaves, and do not burst until the second year after formation. These will develop into inflorescence primordia, while other uncommitted primordia formed on rapidly growing shoots will develop into tendrils. Inflorescence primordia enter dormancy in autumn of the first year when day length decreases then stay dormant throughout winter (Fennell et al., 2015). Bud growth is halted during bud dormancy to better tolerate low temperatures and frost. The second season is initiated in spring with bud burst when environmental conditions become permissive. Shortly before and during bud burst, flower initials are formed and flower parts become distinct. The conversion from inflorescence primordia to inflorescences begins in the spring. Inflorescence formation is regulated at the level of formation of uncommitted primordia and at the level of differentiation. Flowering occurs around May - June in the Northern hemisphere, after which berry formation and berry ripening take place.

## Grapevine veraison

Veraison is the phenological stage that marks the beginning of ripening of the grapes. Several events are initiated during this phase, for example the change in skin colour, berry softening, sugar accumulation, and organic acid decline (Coombe and Bishop, 1980; Uhlig and Clingeleffer, 1998). Grapevine berry ripening follows a biphasic growth, resembling a double sigmoid curve. In the first phase, fruit set is occurring, and little hard green berries develop to a medium size; then growth interrupts for a lag phase and starts again with a second stage of maturation where softening of the berries, sugar accumulation, acid decline and colour changes of the pericarp occur. During ripening, the berries develop the properties specific to the cultivar, and then complex physical and chemical processes take place gradually from veraison to physiological maturity. The polyphenolic compounds, in particular anthocyanins, are the group subject to the main changes during berry ripening; anthocyanins accumulate rapidly, while phenolic acids levels are lower than those of anthocyanins and show different alteration during ripening (Giovanelli and Brenna, 2007). Sugar and organic acid content also undergo significantly changes during berry ripening; while the concentration of glucose and
fructose increase rapidly, organic acids decline (Sadras and Mccarthy, 2007). Each of the compounds that is subject to variation during the ripening stage may differ among genotypes and may be different according to the stage of development of the berry. Veraison is usually evaluated by checking the berries for the beginning of softening or the start of anthocyanin pigmentation, the latter process that results more difficult for white-berried cultivars, that instead of depositing anthocyanins in the skin, lose chlorophyll and start to brighten. In the internationally recognized system of descriptors for grapevine phenotypic traits developed by OIV (Organisation Internationale de la Vigne et du Vin, OIV), veraison time is described by five classes 1, 3, 5, 7 and 9 (OIV descriptor 303), with class 1 representing cultivars displaying an early veraison time and class 9 represents the very late ones. Grapevine phenology stages are also defined with the BBCH code (Lorenz et al., 1994) (stage 81), or in the E-L System (Coombe, 1995) (stage 35). Veraison time is strictly correlated to the other phenological stages and is considered a predictor of maturity and harvest time (Sadras and Petrie, 2011). It may differ significantly between cultivars and has a strong genetic determination (from this thesis, broad sense heritability estimated is 0.424 ). Several QTL studies have identified unique and common genetic loci explaining variable portion of the phenotypic variance, for example on chromosome 16 (Costantini et al., 2008; Duchêne et al., 2012; Fischer et al., 2004; Zyprian et al., 2016). Grapevine breeders have recently started to apply marker-assisted selection, approach that will reveal helpful especially for phenotypic traits where manual evaluation is complex and automated scoring is still lacking, such as phenology traits. Genetic markers associated to veraison time would unveil useful tools for marker-assisted breeding of new cultivars adapted to climate change.

## Impact of climate change on viticulture

It is a truth universally acknowledged that climate change is occurring and is shaping the future of agriculture. Viticulture is one of the niches of agriculture that is suffering the most, given the tremendous differences in terms of wine quality that can result from even light variations in terms of precipitations and temperature during a single season, especially in cool-climate viticulture (Vasconcelos et al.,
2009). In fact, with respect to the weather conditions, yield can vary up to $32.5 \%$ between seasons, which makes grapevine by far the crop with the highest seasonal variation in yield (Boss et al., 2003; Lebon et al., 2008). Yields and quality are not only affected by events mediated all over the season, but also by extreme and precise damages occurring during key phenological stages like budburst and flowering. The major and more evident effect of climate change is global warming, that is leading to a tendency to higher temperatures at earlier dates during the growing season. This obviously results in an advancement of bud burst, flowering and veraison stages. The predicted shift in veraison to earlier dates may cause ripening to occur under higher temperatures (Delrot et al., 2013), thus affecting grape composition and wine quality, in particular by impairing the accumulation of anthocyanins in the berries, a condition that is thought unfavourable for wine quality (Duchene, 2016). Moreover, with earlier ripening a compressed time window for many grapes is observed, leading to a compressed harvesting period. Stronger temperatures or frost damage can also affect the number of flowers and thus the number of berries per plant. Few studies have shown that the higher the temperatures around bud burst, the lower the number of flowers per inflorescence (Keller et al., 2010; Petrie and Clingeleffer, 2005). Furthermore, increased temperatures lead to longer drought periods and water stress which also poses a strain to viticulture; water deficit during floral initiation can lead to a decreased number of inflorescences and can have a negative influence on berry weight, especially when applied after veraison (Niculcea et al., 2014). On the other hand, cold, rain and frost conditions are other problems affecting berries development; in particular cold or rainy weather around flowering can reduce the number of grape clusters formed; frost conditions if occurring during late spring can cause significant crop loss when showing up after bud burst (Mosedale et al., 2015; Mullins et al., 1992).

Among all the possible options that we have to contrast the negative effects of climate change in viticulture, in this thesis we discuss the genetic approach; that is, the possibility of studying and understanding the genetic of the berry ripening process in order to exploit the source of natural variation coming from different grapevine varieties and breed new cultivars adapted to a changing climate.

## Chapter 1

# Integration of meta-QTL analysis and transcriptomic data identifies candidate genes controlling veraison time in grapevine 


#### Abstract

High temperature impairs the quality of grapes and wines. Understanding the genetic control of grapevine phenology-related processes is crucial to successfully breed varieties more adapted to a changing climate. Veraison time, in particular, is a key factor for determining climatic conditions during ripening, in a sense that, depending on the period of the season when occurring, it will impact the time of the season when ripening time will occur. Quantitative trait loci (QTL) studies attempting to elucidate the genetic determinism of developmental stages in grapevine have identified genomic regions including large number of genes. Broad scale transcriptomic studies, by identifying sets of genes modulated along berry development and ripening, have also highlighted a huge number of putative candidates. With the final aim of providing a functional and integrated genomic overview for the genetic control of grapevine veraison time, and of prioritizing possible main genetic regulators, we have applied a meta-QTL analysis for grapevine phenology-related traits and checked for co-localization of transcriptional candidates. Twelve QTL studies were considered, including 174 QTLs related to phenology. By using the software BioMercator v4.2 a consensus genetic map including 3130 markers was compiled. QTLs were projected onto the consensus map and clustered into meta-QTLs. Anchoring to the grapevine genome assembly 12X.v2 allowed us to select positional candidates. We generated 17 metaQTLs from 69 QTLs for the traits flowering, veraison and ripening among which 4 specifically related to veraison time. Moreover 11 meta-QTLs for genomic regions generically affecting phenology were revealed. This approach allowed reducing the number of positional candidates by almost 4-fold. Expression data generated by transcriptomic studies during berry development performed on several grape varieties were mined with different approaches to select, among positional candidates, genes significantly modulated at veraison time.


## Introduction

High temperature impairs the quality of grapes and wines. Understanding the genetic control of grapevine phenology-related processes is crucial to successfully breed varieties more adapted to a changing climate. Veraison time, in particular, is a key factor for determining climatic conditions during ripening. Understanding the genetic control of phenological developmental stages (i.e. flowering, veraison, ripening, etc) would be helpful for the adaptation of local varieties to changing climate. In particular, delaying berry veraison and ripening is a desirable breeding target, since ripening occurring under very hot summer strongly affect and uncouple berry quality traits. By applying QTL analysis, segments of the genome carrying with higher probability genes involved in the traits of interest, and thus with a potential for breeding applications, can be identified. QTL studies allow to define the genetic control of phenotypic traits dissecting the phenotypic variation and determining the contribution of each QTLs. QTLs studies in grapevine have focused on different phenotypic traits. Quantitative trait loci (QTL) studies attempting to elucidate the genetic determinism of developmental stages in grapevine have identified genomic regions including large number of genes. According to these QTL studies, a complex inheritance would control phenology traits, with low contributions scarcely reproducible among studies, even though few reproducible contributions were found. An interesting opportunity to rationalize and interpret the plethora of QTL information, especially to the aim of providing relevant trait candidates, comes from QTL meta-analysis (Goffinet and Gerber, 2000; Veyrieras et al., 2007) a statistical framework to project QTLs on a consensus map which allows to identify and mine co-localizing QTLs among independent experiments. Indeed, QTLs detected independently and located in a given region of a chromosome could possibly represent several estimations of the position of one single QTL. This hypothesis can be tested by appropriate statistical tools which indicate the most likely number of 'real' QTLs underlying a pool of QTLs from independent experiments, providing alongside consensus positions for these narrowing down the QTLs confidence intervals. The resulting meta-QTLs are expected to better define the boundaries of the causative genomic intervals by integrating information from different studies. QTL meta-analyses have become
popular in literature where they are used both to summarize QTL information about one trait as well as to locally verify the co-location of QTLs between different populations as a first step towards QTL validation and/or prioritization of candidates. Chardon et al. (Chardon et al., 2004) first applied this approach to study flowering time in maize by summarizing several QTLs from different mapping populations into meta-QTLs. Subsequent positional cloning and association mapping analysis found in meta-QTL intervals two genes effectively involved in modulating flowering time (Ananiev et al., 2007; Salvi et al., 2011). These successful examples confirmed meta-analysis as very useful method for predicting candidate genes and for developing molecular markers for breeding. Lately, metaanalysis has been successfully used in studying QTL in different species like rice (Khowaja et al., 2009), cotton (Said et al., 2015), potato (Danan et al., 2011) and many others. So far, this approach has not been applied in grapevine.

Given this, it must be also noted that the meta-analysis of QTLs is a methodology that does not necessarily imply causation; that is, a newly defined meta-QTL may arise from different original input QTLs that share a common genetic determinant, but may also arise from a simple "physical" co-localization or proximity. Especially in not so dense genetic maps, QTL often spans several cM , resulting in easy overlapping of QTLs that do not have a real biological common meaning. Also when different "rounds" of meta-analysis are conducted with QTLs identified from different phenotypic traits (Danan et al., 2011), the risk of obtaining meta-QTLs that do not share the same genetic background is higher. Indeed, QTL colocalization can be due to tight-linkage of QTL/genes playing different functions, but could also arise from pleiotropy; when pleiotropy is likely, it would also justify meta-analysis across traits, to further reduce the number of candidates. Such a risk of "fake" co-localization may be contained if input QTLs are manually selected and curated, and maybe also if genes contained in the original QTLs intervals are screened in advance and checked for congruence. A great advancement in the metaQTL analysis would also come from the integration of original genotypic data instead of genetic maps markers' distance and name; operation that is limited by the lack of resources and standards to deposit original genotyping data. Indeed, by using original data to create consensus map, this will reflect more precisely the real
position and distance of the different markers, then the QTLs projection would be more accurate, avoiding false overlappings and moreover the information of the alleles of the parents would be available thus making it fundamental in the view of selecting the favourable allele for new breeding programs.

Transcriptomic experiments in grapevine identified a number of genes as candidates for phenology in particular for the transition from immature to mature stage, the so-called veraison (Fasoli et al., 2012; Massonnet et al., 2017; Palumbo et al., 2014). The number of this transcriptomic candidates and QTL positional candidates is very large. To prioritize candidate genes it would be useful to integrate meta-QTL results, in particular meta-QTL positional genes, with transcriptomic candidate genes to identify most important genes. Broad scale transcriptomic studies, by identifying sets of genes modulated along berry development and ripening, have also highlighted a huge number of putative candidates. With the final aim of providing a functional and integrated genomic overview for the genetic control of grapevine veraison time, and of prioritizing possible main genetic regulators, we have applied a meta-QTL analysis for grapevine phenology-related traits and checked for co-localization of transcriptional candidates. We have created the first grapevine consensus map, performed the first meta-QTL analysis in grapevine, anchored the consensus map to the updated assembly of grapevine genome, integrated data from transcriptomic studies and prioritized candidate genes for veraison control in grapevine. In this study, we report a candidate-genes prioritization approach based on two steps: in the first step we perform a metaanalysis of QTLs associated to grapevine phenology; in the second step we integrate the information of grapevine phenology-related hub-genes pinpointed in transcriptomic studies with the results of the QTL meta-analysis, in order to identify genes to be considered as key regulators for the veraison stage of grapevine. The meta-QTL analysis was conducted creating the first grapevine genetic consensus map, integrating 39 different mapping populations. The consensus map was then anchored to the grapevine genome using 1055 markers. To our knowledge, this is the first meta-QTL analysis conducted in grapevine.

## Methods

## Grapevine consensus map construction

The process of consensus map creation started with the collection of individual genetic linkage maps with information on marker names and position in cM . To make it possible to combine the individual maps into a consensus map, marker names must be consistent. We manually checked and curated the name of the markers in order to correct misspellings and find synonyms. The files were input in BioMercator V4.2 (input files available upon request) and the linkage groups were oriented according to the reference map of Doligez (Doligez et al., 2006). Linkage groups that did not share at least two markers with the corresponding linkage groups of other maps were removed from the analysis, leading to a different number of input maps in the construction of the consensus linkage groups. The command InfoMap was used to evaluate markers order and consistency between each pair of input maps; in case of inversions in markers order between maps, the occurrence of the inverted markers in all the maps was evaluated and only the marker less represented across all the maps was removed. After all the inversions were corrected, the command ConsMap was used to construct the consensus map in a single step, without using any map as reference.

## In silico mapping of GCM markers and anchoring to the grapevine genome

Sequences of the GCM markers were downloaded from the publications where they originally appeared and were blasted against the 12X.v2 assembly of the grapevine genome using the website https://urgi.versailles.inra.fr/blast/. An anchor map was thus created composed of the univocally mapping GCM markers and corresponding position in base pairs. The anchor map was uploaded to BioMercator and the option "New genome version" was used to anchor the GCM to the new structural annotation of the grapevine genome as a .gff3 file (https://urgi.versailles.inra.fr/Species/Vitis/Annotations). Physical positions of QTLs, meta-QTLs and candidate genes located in confidence intervals were then automatically calculated by BioMercator using an internal formula (Yannick De Oliveira, personal communication) and downloaded to be reported in the tables.

## QTL projection and meta-analysis

Before performing the meta-analysis, specific information of QTL data were collected, in particular start and end position of the confidence interval and peak of the QTL, its associated variance explained value $\left(R^{2}\right)$, and the size of the population that was used for mapping the QTL. These information were input in BioMercator and each QTL was associated to the genetic map where it was originally mapped. The command QTLProj was then applied in order to project the QTLs of the component maps to the consensus map; the command performs a homothetic projection of the original QTL to the consensus map only when flanking markers are found where the ratio of the distance of these markers to the confidence interval of the QTL that is being projected is not reduced by a factor greater than 0.25 . In our case default options were kept. The meta-analysis itself was then executed; we used the Veyrieras algorithm (Veyrieras et al., 2007) which in BioMercator is divided in two steps, the QTLClust and MQTLView. The QTLClust performs the clustering of the input QTLs belonging to the same trait and finds the real number of meta-QTLs, calculating as many models as the number of the input QTLs and five different criterion values, AIC (Akaike information criterion), AICc, AIC3, BIC (Bayesian information criterion) and AWE (average weight of evidence), for each one of the models. The best model was then selected as the model minimizing the criterion value. The second step, MQTLView, was then used to graphically represent the meta-QTLs identified according to the selected model.

## Transcriptomic data integration

RNA-Seq gene expression data (Fasoli et al., 2018) from 99 berry RNA samples from the cultivars Pinot Noir collected in triplicates in the years 2012, 2013 and 2014 around the time of veraison were analysed. To identify the most significant genes modulated across veraison, we developed a strategy that takes into account the entity of the modulation in the comparison of the time-point before veraison. We used a recently published dataset (Fasoli et al., 2018) reporting the results of the differential expression analysis between different time-points of the berry samples from the cultivars Pinot Noir and Cabernet Sauvignon, sampled in triplicated in the years 2012, 2013 and 2014. We selected the genes that, between
the time-point -T1 and veraison (T0), were significantly differentially expressed and displayed an absolute value of $\log 2$ of the fold change $(|\log 2 F C|)$ greater than all the other comparisons.

## Results and discussion

## Collection of grape QTLs studies for QTLs data integration

With the final aim of integrating available information about grapevine QTLs emerging from independent studies/populations, and especially QTLs related to veraison time control, literature was mined to identify all published grapevine QTL studies up to October 2018. Studies relying on genetic maps without enough shared markers with other maps (i.e. genetic maps including only AFLP or SNPs), or not providing information about genetic map version used as reference or all genetic information about detected QTLs, were not considered. This resulted in the selection of 42 publications, reporting 47 different QTL maps. A list of all these, including reference and information about the related genetic map, is provided in Table 1. These QTL studies exploited overall 24 different cross populations, constituted on average by 157 offsprings (number of offsprings ranging from 74 to 265). Cross population were mainly F1, with the only exception of two populations obtained by self-pollination and one obtained by selfing an F1 (Blasi et al., 2011; Duchêne et al., 2009; Fennell et al., 2019). Large number of cross populations (14) were derived by crossing Vitis vinifera with hybrids or other Vitis species, but a number of intra-vinifera cross was also represented. The selected QTL studies included 2093 QTLs for 354 different phenotypes scored. For each QTL study we focused on QTLs detected in the consensus map, when this was provided. QTLs mapped on parental maps were included only if genetic data for consensus maps were not available (see Table 1). All QTLs were considered, independently of their score thresholds, LOD/variance values or years of observation. A detailed list of all scored phenotypes, grouped for each study and including the QTL short name used in the relative reference as well as a short description, is provided as Table S1 (external file). More details about the phenotypic scoring for each QTL can be found in the original reference. Each measured phenotype/QTL was manually attributed to its most related trait, for which the score was considered to be a
descriptor, and traits were arbitrarily grouped in nine main trait categories. An overview about currently more characterized plant traits in grape, grouped according to the nine different trait categories, is given in Figure 1.

Table 1. List of QTL studies

| QTL study reference | Genetic map reference | Cross | Female Parent | Male Parent | $\begin{aligned} & \text { Pop } \\ & \text { Type } \end{aligned}$ | $\begin{aligned} & \text { Pop } \\ & \text { Size } \end{aligned}$ | QTL categories | Total number of used QTLs |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| Azuma et al. 2015 | Ban et al. 2014 | line 693 | 626-84 | Iku82 | F1 | 74 | berry metabolites | 31 |
| Ban et al. 2014 | Ban et al. 2014 | line 693 | 626-84 | Iku82 | F1 | 98 | berry metabolites | 10 |
| Ban et al. 2016 | Ban et al. 2014 | line 693 | 626-84 | Iku82 | F1 | 98 | berry morphology, phenology | 24 |
| Bayo Canha, PhD thesis 2015 | Bayo Canha, PhD thesis 2015 | MN x SY | Vv Monastrell | Vv Syrah | F1 | 229 | phenology, berry morphology, cluster related traits, berry metabolites | 40 |
| Bellin et al. 2009 | Bellin et al. 2009 (parental maps) | CHx BI | Vv Chardonnay | Bianca | F1 | 116 | pathogen resistance | 18 |
| Bert et al. 2013 | Bert et al. 2013 | CS x RGM1995-1 | Vv Cabernet Sauvignon | Vr Gloire de Montpellier | F1 | 138 | vegetative traits, abiotic stress response | 129 |
| Blasi et al. 2011 | Blasi et al. 2011 | V. amurensis 'Ruprecht' S1 | Va 'Ruprecht' | Va 'Ruprecht' | S1 | 232 | pathogen resistance | 8 |
| Cabezas et al. 2006 | Cabezas et al. 2006 | D x AS | Vv Dominga | Vv Autumn Seedless | F1 | 118 | berry morphology, seeds related traits | 20 |
| Carreño Ruiz, PhD thesis 2012 | Carreño Ruiz, PhD thesis 2012 | RS x M | Vv Ruby Seedless | Vv Moscatuel | F1 | 78 | phenology, cluster related traits, berry morphology, berry metabolites, seeds related traits | 72 |
| Correa et al. 2014 | Correa et al. 2014 | RS x S | Vv Ruby Seedless | Vv Thompson Seedless | F1 | 137 | cluster related traits | 19 |
| Correa et al. 2015 | Correa et al. 2015 | RS x S | Vv Ruby Seedless | Vv Thompson Seedless | F1 | 137 | berry morphology, seeds related traits | 40 |
| Correa et al. 2016 | Correa et al. 2016 | RS x S | Vv Ruby Seedless | Vv Thompson Seedless | F1 | 137 | berry morphology | 6 |
| Costantini et al. 2008 | Costantini et al. 2008 | I x BP | Vv Italia | Vv Big Perlon | F1 | 163 | berry morphology, seeds related traits, phenology | 25 |
| Costantini et al. 2015 | Costantini et al. 2015 | SY x PN | Vv Syrah | Vv Pinot Noir | F1 | 170 | berry metabolites | 554 |
| Coupel-Ledru et al. 2014 | Coupel-Ledru et al. 2014 | Sx G (and reverse) | Vv Syrah | Vv Grenache | F1 | 186 | vegetative traits, abiotic stress response | 29 |
| Coupel-Ledru et al. 2016 | Coupel-Ledru et al. $2014$ | S x G (and reverse) | Vv Syrah | Vv Grenache | F1 | 186 | vegetative traits, abiotic stress response | 100 |
| Doligez et al. 2010 | Doligez et al. 2010 | MTP3140 | Vv MTP2223-27 | Vv MTP2121-30 | F1 | 139 | cluster related traits | 2 |
|  | Doligez et al. 2010 | MTP3140 | Vv MTP2223-27 | Vv MTP2121-30 | F1 | 139 | berry morphology, seeds related traits | 55 |
| Doligez et al. 2013 | Coupel-Ledru et al. 2014 | S x G (and reverse) | Vv Syrah | Vv Grenache | F1 | 191 | berry morphology, seeds related traits | 77 |
| Duchêne et al. 2009 | Duchêne et al. 2009 | S1 MO | Vv Muscat Ottonel | Vv Muscat Ottonel | S1 | 121 | berry metabolites | 7 |
| Duchêne et al. 2012 | Duchêne et al. 2012 | RI x GW | Vv Riesling | Vv Gewürztraminer | F1 | 188 | phenology | 22 |
|  | Fechter et al. 2014 | V3125 x Börner | Vv V3125 | Börner | F1 | 202 | phenology | 22 |
| Fechter et al. 2014 | Zyprian et al. 2016 | Gf.Ga-47-42 x Villard blanc | Gf.Ga-47-42 | Villard blanc | F1 | 151 | phenology | 9 |
| Garris et al. 2009 | Garris et al. 2009 | Vitis riparia x Vitis hybrid Seyval | Vr PI 588289 | Seyve Villard 5-276 | F2 | 119 | vegetative traits | 17 |


| Grzeskowiak et al. 2013 | Costantini et al. 2015 | SY x PN | Vv Syrah | Vv Pinot Noir | F1 | 170 | phenology, cluster related traits | 27 |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| Guo et al. 2015 | Guo et al. 2015 | '87-1’ x ‘9-22' | 87-1 | 9-22 | F1 | 149 | berry metabolites | 30 |
| Herzog et al., 2015 | Zyprian et al. 2016 | Gf.Ga-47-42 x Villard blanc | Gf.Ga-47-42 | Villard blanc | F1 | 151 | pathogen resistance | 1 |
| Huang et al. 2012 | Huang et al. 2012 | SxG (and reverse) | Vv Syrah | Vv Grenache | F1 | 191 | berry metabolites | 114 |
| Huang et al. 2014 | Huang et al. 2012 | SxG (and reverse) | Vv Syrah | Vv Grenache | F1 | 191 | berry metabolites | 29 |
| Malacarne et al. 2015 | Costantini et al. 2015 | SY x PN | Vv Syrah | Vv Pinot Noir | F1 | 170 | berry metabolites | 200 |
| Marguerit et al. 2012 | Marguerit et al. 2012 (parental maps) | CS x RGM1995-1 | Vv Cabernet Sauvignon | Vr Gloire de Montpellier | F1 | 138 | vegetative traits, abiotic stress response | 22 |
| Mejía et al. 2007 | Mejía et al. 2007 | RS $\times$ S | Vv Ruby Seedless | Vv Thompson Seedless | F1 | 144 | berry morphology, phenology, seeds related traits | 26 |
| Mejía et al. 2011 | Mejía et al. 2011 | RS x S | Vv Ruby Seedless | Vv Thompson Seedless | F1 | 139 | berry morphology, seeds related traits | 6 |
| Moreira etal 2011 | Moreira et al. 2011 | $\mathrm{VN} \times \mathrm{RP}$ | Vv White Muscat | Vr Wr 63 | F1 | 174 | pathogen resistance | 26 |
| Moreiraetal. 201 | Moreira et al. 2011 | RTx AM | VRH3082 1-42 | SK77 5/3 | F1 | 94 | pathogen resistance | 6 |
| Ochssner et al. 2016 | Ochssner et al. 2016 | V3125 x Börner | Vv V3125 | Börner | F1 | 202 | pathogen resistance | 7 |
| Rex et al. 2014 | Zhang et al. 2009 | V3125 x Börner | Vv V3125 | Börner | F1 | 188 | pathogen resistance | 20 |
| Schwander et al. 2012 | Schwander et al. 2012 | Gf.Ga-52-42 x Solaris | Gf.Ga-52-42 | Solaris | F1 | 265 | pathogen resistance | 5 |
| VanHeerden et al. 2014 | VanHeerden et al. 2014 (Regent parental map) | Regent x Red Globe | Regent | Vv Red Globe | F1 | 186 | pathogen resistance | 7 |
| Viana et al. 2013 | Viana et al. 2013 | AT0023 | D8909-15 | Vv B90-116 | F1 | 203 | vegetative traits, phenology, cluster related traits, berry morphology, berry metabolites, seeds related traits | 41 |
| Welter et al. 2007 | Welter et al. 2007 | RxL | Regent | Vv Lemberger | F1 | 144 | pathogen resistance | 21 |
| Zhang et al. 2009 | Zhang et al. 2009 | V3125 x Börner | Vv V3125 | Börner | F1 | 188 | pathogen resistance | 2 |
| Zhao et al. 2015 | Zhao et al. 2015 | '87-1' x ‘9-22' | 87-1 | 9-22 | F1 | 149 | berry morphology, phenology, berry metabolites | 5 |
| Zhao et al. 2016 | Zhao et al. 2015 | '87-1' x '9-22' | 87-1 | 9-22 | F1 | 149 | phenology | 5 |
| Zyprian et al. 2016 | Zyprian et al. 2016 | Gf.Ga-47-42 x Villard blanc | Gf.Ga-47-42 | Villard blanc | F1 | 151 | phenology, pathogen resistance | 157 |

Number of QTLs for each trait as well as number of studies considering each trait are shown. The trait for which the highest number of QTLs is currently available in the literature is berry metabolites content. This is expected since high throughput metabolomic approaches can easily release large datasets. However, the overall most scored trait across independent studies was berry weight (scored in 12 independent studies), while categories most addressed by QTL studies so far have been phenology and pathogen resistance.


| Abiotic Stress Response | Berry Metabolites | Berry Morphology | Cluster Related Traits | Pathogen Resistance | Phenology | Seeds Related Traits | Vegetative Traits |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| Drought Stress | Anthocyanins | Berry Weight | Fertility | Downy Mildew Resistance | Ripening Time | Seed Number | Growth |
| Chlorosis | Terpenols | Berry Firmness | Berry Number | Powdery Mildew Resistance | Veraison Time | Seed Weight | Leaf Morphology |
|  | Seed And Skin Tannins | Berry Diameter | Peduncle Length | Phylloxera Resistance | Flowering Time | Seed Percent Dry Matter | Water Use Efficiency |
|  | Flavonol | Berry Volume | Cluster Architecture | Black Rot Resistance | Interval | Seed Response To Ga | Leaf Area |
|  |  | Berry Response To Ga | Cluster Weight | Botrytis Resistance | Budburst |  |  |
|  |  |  | Cluster Response To Ga |  |  |  |  |

Figure 1. Summary of phenology QTL studies

## Building a grapevine consensus genetic map

All the 35 reference genetic maps for the QTL studies were used as input for the construction of a consensus map in BioMercator 4.2 software (Sosnowski et al., 2012). Moreover a grapevine reference map, developed from the integration of 5 different genetic maps (Doligez et al., 2006), was also included, as well as few other available grapevine genetic maps (Costantini et al., 2008; Venuti et al., 2013). Common markers made it possible the construction of a consensus for each of the 19 grapevine chromosomes with no residual conflicts. The consensus map consisted of 19 linkage groups, corresponding to the 19 grapevine chromosomes, including 3130 markers with a total length of 1922 cM and an average number of markers and length per linkage group of 164 and 101 cM respectively. The number of markers shared by at least two maps was 1209 , corresponding to $38.63 \%$ of the total markers, with an average of 63 shared markers per linkage group (Table 2). The full map file and a graphical overview are provided as Table S2 and Figure S1. The number of maps used for the construction of each linkage group varied from 26 (LG 11) to 39 (LGs $1,2,4,5,10,12,17,18$ ), due to the different number of markers shared among maps (Table 2, Table S3 [external file]).

Table 2. Consensus genetic map features

| LG | $\mathbf{N}^{\circ}$ of <br> markers | $\mathbf{N}^{\circ}$ unique <br> markers | $\mathbf{N}^{\circ}$ markers in at least <br> two maps | Length <br> $(\mathbf{c M})$ | $\mathbf{N}^{\circ}$ of individual maps <br> integrated |
| :---: | :---: | :---: | :---: | :---: | :---: |
| I | 214 | 130 | 84 | 95.68 | 39 |
| II | 130 | 71 | 59 | 89.73 | 39 |
| III | 135 | 92 | 43 | 92.03 | 37 |
| IV | 161 | 101 | 60 | 93.36 | 39 |
| V | 206 | 150 | 56 | 70.64 | 39 |
| VI | 139 | 93 | 46 | 90.72 | 38 |
| VII | 204 | 124 | 80 | 82.09 | 38 |
| VIII | 167 | 88 | 79 | 95.72 | 37 |
| IX | 128 | 77 | 51 | 85.01 | 35 |
| X | 168 | 93 | 75 | 141.87 | 39 |
| XI | 90 | 38 | 52 | 72.64 | 26 |
| XII | 211 | 153 | 58 | 143.13 | 39 |
| XIII | 156 | 89 | 67 | 94.79 | 37 |
| XIV | 202 | 118 | 84 | 93.44 | 37 |
| XV | 128 | 86 | 42 | 68.7 | 35 |
| XVI | 126 | 74 | 52 | 104.47 | 39 |
| XVII | 130 | 76 | 54 | 148.93 | 39 |
| XVIII | 275 | 172 | 96 | 64 | 103 |

Marker density was not equally distributed along the consensus, with peaks in putative centromeric positions similarly as found in original maps and in agreement with the expected suppression of recombination rate in such regions. However, comparison of markers order between the single component maps and the consensus map revealed a high level of correlation. Spearman's rank correlation values of pairwise comparisons were significantly high for all maps but two, possibly due to the low number of shared markers (Figure S2). The consensus genetic map was connected to the reference genome through the use of an anchor file. Markers physical position was recovered as explained in the methods section. Upon removal of markers showing incongruent or not unique physical position, 713 markers (on average of 38 markers per LG) were finally physically mapped on the 12X.v2 assembly of the grapevine genome (Canaguier et al., 2017). Their physical coordinates are also included in the map file Table S3. Among these markers, 480 (67\%) were shared by at least two original maps, and the majority (513, 72\%) were microsatellite markers.

## Distribution of grapevine QTLs on the consensus genetic map

All QTLs from the 47 QTL studies (Table 1) were projected onto the consensus map to identify possible overlaps across populations. In total 1899 QTLs (91\%) could be successfully plotted while 194 QTLs could not be projected to the consensus map due to the lack of anchoring markers. QTLs reduction was comparable across the different QTL categories, ranging from $87 \%$ of successfully plotted QTLs for berry metabolism to $98 \%$ for abiotic stress. Only for the trait category "cluster related trait" the number of plotted QTLs was lower (79\%). Summary plots with distribution of projected QTLs for each trait category are provided as Figure 2. For each QTL consensus map, hotspots of overlapping QTLs across studies relying on independent populations have been highlighted. In summary we found independent overlapping QTLs for all the traits related to berry morphology (LG 1, 15, 18) and seeds related traits (LG 1, 10, 14 and 18), with only exception of traits measured in response to GA, for all vegetative traits $(1,4,10$, 13,18 ) and for all phenology related traits (LG $1,2,3,6,7,14,16,18$ ) except bud burst. Moreover, we also found overlapping QTLs for anthocyanins (LG 2), and
downy (LG $1,4,5,6,7,12,17,18$ ) or powdery mildew resistance (LG 15). No overlapping QTLs were found instead for traits belonging to the category abiotic stress and for traits related to clusters.


Figure 2. Summary graphic plots with distribution of projected QTLs for each trait category. Clockwise, from top left: abiotic stress response, cluster related traits, berry metabolites, berry morphology (Continue)


Figure 2. Continued. Summary graphic plots with distribution of projected QTLs for each trait category. Clockwise, from top left: pathogen resistance, phenology, inlcuding meta-QTLs on the inner circle, seed related traits, vegetative traits.

Interestingly, overlapping QTLs across different traits or categories possibly involved in the expression of more complex traits can also be scored in this dataset. As an example, overlapping QTLs from independent studies were found in LG 18 both for seed fresh weight and berry weight traits. Since berry weight is known to be also dependent on seed content (Doligez et al., 2013) these overlaps could hide a common genetic basis and could thus be exploited in a meta-analysis aiming to reduce, based on QTL co-locations, the number of underlying positional candidate genes. In the following we provide a detailed meta-analysis on phenology related traits aiming to identify candidate genes for veraison time.

## Narrow down of candidates for veraison time by meta QTL-analyses

The list of phenology related QTLs projected onto the consensus map (141 QTLs from 13 studies) was manually curated for the purpose of performing a metaanalysis on overlapping QTLs from independent studies. Redundant QTLs, that is, QTLs with same peak position from the same study, which could overestimate the effect of that QTL in the analysis (Danan et al., 2011), were pruned as explained in the material and methods section. For the meta-analysis we considered 35 veraison related QTLs derived from six studies (Bayo Canha, 2016; Carreño Ruiz, 2012; Costantini et al., 2008; Emanuelli et al., 2013; Fechter et al., 2014; Zyprian et al., 2016). Meta-analysis was performed if at least two QTLs from independent experiments were overlapping. For veraison, overlapping QTLs from independent studies were found on LG1 and LG2 (from two studies and three studies respectively). The optimal number of meta-QTLs explaining overlapping QTLs was statistically determined by choosing the most likely model, as computed by BioMercator V4.2 software by five different tests. Our meta-analysis resulted in the identification of 4 veraison meta-QTLs located respectively on LG1 (one meta QTL) and on LG2 (three meta QTLs) (Table 3, Figure 2F). More in detail veraison meta-QTLs on LG2 resulted by integration of at least five original co-located QTLs, while ver_1_1 on LG1 was derived from two original co-located QTLs. Average CI was 3.5 cM ranging from 1.2 cM for ver_2_3 to 5.1 cM per ver_2_1, which was the largest one. On LG1 the original CI covered by QTLs was reduced from 23.9 cM to 4.3 cM ( 5.6 times) by the meta-analysis. On LG2 the reduction of CI by meta-
analysis was overall of five times, with a strongest effect on the ver_2_3 meta QTL. $R^{2}$ values of meta QTLs were all higher that $10 \%$. In particular ver_2_2 was the most relevant, explaining up to $34 \%$ of total variance. A similar meta-QTL analysis was applied to overlapping QTLs for berry colour on LG2 to validate our procedure. Indeed, berry colour genetic control has been already elucidated and linked to a transposon insertion in the promoter region of the MybA1 gene located on Chr 2 (Fournier-Level et al., 2009; Kobayashi et al., 2004; Walker et al., 2007). The metaQTL analysis on 28 overlapping QTLs derived from five independent studies identified 7 meta-QTLs (Table S5). Interestingly the MybA1 gene was included in the list of the 125 genes underlying these meta-QTLs (Table S6). Given such a high number and considering the high percentage of variance explained by many of these original QTLs, the meta-analysis resulted in a high number of very small metaQTLs, which eventually can be considered as a single meta-QTL spanning a wider area. Here is a limitation of the meta-QTL approach when dealing with situations where the number of QTLs is high and the peaks of the original QTLs are very close to each other but not overlapping.

Inspection of the phenology QTL consensus map revealed extensive co-localization across traits for different developmental stages (i.e. co-location of veraison and ripening QTLs). Co-location of veraison QTLs with other phenology QTLs was indeed highly significant compared to a random distribution ( $\chi^{2}$-test, $\mathrm{p}<0.01$ ). Overlapping phenology QTLs could represent several estimates of a single QTL affecting more developmental stages, which would justify the attempt to identify consensus QTLs across different phenology traits (Bancroft et al., 2009). In agreement with such option a meta-analysis for veraison QTLs including overlapping QTLs for other phenology traits on LG1 and LG2 identified metaQTLs (pheno-QTLs) largely overlapping with previously reported meta-QTLs regions (Table S7). Therefore, with the final aim of reducing the number of candidate genes underlying veraison QTLs, we decided to apply also meta-analysis on veraison QTLs when overlapping QTLs from other phenological traits could be found in independent studies; then we have identified 13 further indicative metaQTLs regions (pheno-QTLs) (Table 4, Figure 2H). Among these, two meta-QTLs on LG16 were particularly relevant, explaining on average up to $35 \%$ and $38 \%$ of
total phenotypic variance. In conclusion, the number of candidate genes underlying original veraison QTLs was narrowed down by applying meta-analysis of veraison QTLs, by a factor of 3.7. Meta-QTL analysis including alternative phenology related traits allowed also to reduce ( 2.2 times) the number of positional candidates (Figure 3), however this should be considered cautiously since relevant candidates might be skipped. Lists of candidate genes in veraison meta-QTLs and pheno metaQTLs intervals, with the corresponding CRIBIv1 annotation (Vitulo et al., 2014), are given in Table S 8 and Table S 9 respectively.

Table 3. Results of the meta-QTL analysis on overlapping veraison QTLs. LG Linkage Group, Meta-QTL The name assigned to the meta-QTL identified, Peak Position (cM) Position in $c M$ of the peak of the meta-QTL, $\boldsymbol{R}^{2}$ Proportion of variance explained by the meta-QTL, Start (cM) Start position in cM of the meta-QTL, End (cM) End position in cM of the meta-QTL, Start (bp) Start position in bp of the meta-QTL, End (bp) End position in bp of the meta-QTL, Meta-QTL Positional Candidates The number of positional candidate genes within the metaQTL interval, Original QTLs co-located The number of input QTLs that produced the meta-QTL, QTL Studies (Populations) The number of grapevine populations that produced the cross where QTLs were identified, Traits Phenotypic traits studied for the original QTL analysis, Reference Citation of the original publication where QTLs were originally reported.

| LG | $\begin{aligned} & \text { Meta- } \\ & \text { QTL } \end{aligned}$ | Peak Position Position (cM) | $\mathbf{R}^{2}$ | $\begin{aligned} & \text { Start } \\ & (\mathrm{cM}) \end{aligned}$ | $\begin{aligned} & \text { End } \\ & \text { (cM) } \end{aligned}$ | Start <br> (bp) | End (bp) | Meta-QTL <br> Positional <br> Candidates | Original QTLs co-located | QTL Studies <br> (Populations) | Traits | Reference |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| I | ver_1_1 | 31.29 | 0.11 | 29.15 | 33.43 | 2510506 | 3254952 | 78 | 2 | 2 | VT | Fechter et al 2014, Zyprian et al 2016 |
| II | ver_2_1 | 31.34 | 0.17 | 28.79 | 33.89 | 4029921 | 5344816 | 147 | 7 | 2 | VB, Vr | Bayo Canha 2015, Grzeskowiak et al 2013 |
|  | ver_2_2 | 41.55 | 0.13 | 40.00 | 43.30 | 5717649 | 7154894 | 96 | 5 | 3 | $\begin{aligned} & \text { Vr, VB, } \\ & \text { VE, VT } \end{aligned}$ | Costantini et al 2008, Bayo Canha 2015, Grzeskowiak et al 2013 |
|  | ver_2_3 | 53.47 | 0.34 | 52.88 | 54.07 | 13336750 | 16677137 | 94 | 5 | 3 | $\begin{aligned} & \text { Vr, VE, } \\ & \text { VP } \end{aligned}$ | Costantini et al 2008, Bayo Canha 2015, Grzeskowiak et al 2013 |

Table 4. Results of the meta-QTL analysis on veraison QTLs overlapping with other phenology QTLs. LG Linkage Group, Meta-QTL The name assigned to the meta-QTL identified, Peak Position (cM) Position in cM of the peak of the meta-QTL, $\boldsymbol{R}^{2}$ Proportion of variance explained by the meta-QTL, Start (cM) Start position in cM of the meta-QTL, End (cM) End position in cM of the meta-QTL, Start (bp) Start position in bp of the meta-QTL, End (bp) End position in bp of the meta-QTL, Meta-QTL Positional Candidates The number of positional candidate genes within the meta-QTL interval, Original QTLs co-located The number of input QTLs that produced the meta-QTL, QTL Studies (Populations) The number of grapevine populations that produced the cross where QTLs were identified, Traits Phenotypic traits studied for the original QTL analysis, Reference Citation of the original publication where QTLs were originally reported.

| LG | Meta-QTL | Peak Position (cM) | $\mathbf{R}^{2}$ | $\begin{aligned} & \text { Start } \\ & \text { (cM) } \end{aligned}$ | $\begin{aligned} & \text { End } \\ & \text { (cM) } \end{aligned}$ | Start (bp) | End (bp) | Meta-QTL <br> Positional <br> Candidates | Original QTLs colocated | QTL Studies (Populations) | Traits | Reference |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| III | pheno_3_1 | 27.67 | 0.15 | 24.43 | 30.92 | 560404 | 1647064 | 138 | 5 | 3 | $\begin{gathered} \text { VT, SSC, } \\ \text { Bpc } \end{gathered}$ | Carreño Ruiz 2012, Viana et al 2013, Zhao et al 2015 |
|  | pheno_3_2 | 50.42 | 0.14 | 45.30 | 55.54 | 5903464 | 10894193 | 288 | 4 | 3 | VT, SSC, Bpc, BB | Carreño Ruiz 2012, Viana et al 2013, Zhao et al 2015 |
| V | pheno_5_1 | 50.97 | 0.09 | 49.77 | 52.18 | 16799689 | 19536797 | 111 | 3 | 2 | VT,F-V, Ma | Zyprian et al 2016, Bayo Canha 2015 |
| VII | pheno_7_1 | 9.59 | 0.16 | 7.58 | 11.60 | 1087707 | 1552842 | 59 | 2 | 2 | VT, Fw | Carreño Ruiz 2012, Bayo Canha 2015 |
| XI | pheno_11_1 | 16.15 | 0.11 | 15.01 | 17.30 | 2934932 | 3356851 | 50 | 4 | 2 | $\begin{aligned} & \text { FBL, FS, } \\ & \text { Tar/Ma, VT } \end{aligned}$ | Bayo Canha 2015, Fechter et al 2014 |
| XII | pheno_12_1 | 77.85 | 0.19 | 74.31 | 81.40 | 23793458 | 24155112 | 27 | 2 | 2 | VT, RT | Carreño Ruiz 2012, Zyprian et al 2016 |
| XIV | pheno_14_1 | 55.03 | 0.22 | 51.45 | 58.62 | 22441297 | 24645689 | 157 | 7 | 4 | $\underset{\mathbf{V T}}{\text { B-F, FS, FT, }}$ | Carreño Ruiz 2012, Zyprian et al 2016, Fechter et al 2014, Duchêne et al 2012 |


| XVI | pheno_16_1 | 34.70 | 0.31 | 32.53 | 36.88 | 14012548 | 16583139 | 126 | 4 | 2 | F-V, VT | Costantini et al 2008, Zyprian et al 2016 |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | pheno_16_2 | 38.49 | 0.38 | 36.49 | 40.50 | 16503904 | 17318604 | 51 | 5 | 2 | F-V, VT | Zyprian et al 2016, Duchêne et al 2012 |
| XVII | pheno_17_1 | 48.83 | 0.13 | 45.12 | 52.54 | 4969509 | 6401642 | 113 | 6 | 3 | $\begin{aligned} & \text { FBL, FS, } \\ & \text { RDA, VB } \end{aligned}$ | Fechter et al 2014, <br> Grzeskowiak et al 2013, Mejía et al 2007 |
|  | pheno_17_2 | 61.83 | 0.11 | 61.46 | 62.20 | 8920888 | 9063993 | 12 | 7 | 4 | $\begin{aligned} & \text { FBL, FS, } \\ & \text { RDA, VB, } \\ & \text { F-V } \end{aligned}$ | Fechter et al 2014, Grzeskowiak et al 2013, Mejía et al 2007, Zyprian et al 2016 |
| XVIII | pheno_18_1 | 34.68 | 0.17 | 28.21 | 41.15 | 1836848 | 5349350 | 322 | 2 | 2 | VT, FT | Carreño Ruiz 2012, Zyprian et al 2016 |
|  | pheno_18_2 | 66.33 | 0.13 | 60.57 | 72.10 | 10927035 | 15526564 | 330 | 4 | 3 | VT, FT, F-V | Carreño Ruiz 2012, Zyprian et al 2016, Duchêne et al 2012 |



Figure 3. Bar plot showing for each linkage group where meta-QTLs were identified, the magnitude of positional and transcriptional candidate genes reduction in terms of the absolute number of candidate genes for each linkage group and each class.

## Prioritization of candidates by transcriptomic data integration

Positional candidates underlying meta-QTLs were explored for their expression in different organs according to the grapevine expression atlas (Fasoli et al., 2012). Sixty-three genes among the positional candidates underlying veraison meta-QTLs ( $15.2 \%$ of the positional candidates) were never expressed either in berry, rachis or seed and were thus excluded from our candidate list. In a similar way 350 never expressed genes ( $19.6 \%$ of the positional candidates) were excluded from candidates underlying pheno meta-QTLs. Transcriptomic changes in berries along development and in particular across veraison have previously been widely explored, revealing that a massive transcriptomic change is associated to the veraison (Palumbo et al., 2014). More in detail, by comparing the expression profiles at four different berry developmental stages in 10 different grapevine varieties a first list of shared genes differentially expressed across veraison was found (Massonnet et al., 2017; Palumbo et al., 2014). Moreover, a recent RNA-Seq study has characterized weekly gene expression in Pinot Noir berries along development in the three years 2012, 2013 and 2014 (Fasoli et al., 2018). With the final aim of selecting genes putatively involved in veraison control based on their expression profile we decided to exploit such transcriptomic information. RNASeq data were inspected to spot in each year the expected massive transcriptional remodulation associated to the veraison transition (Table S10). The transition across which the highest number of genes was differentially expressed was considered as "molecular veraison". Genes which were mainly modulated in their profiles across this transition in at least two of the three years have been selected also as transcriptional candidates. Altogether a list of 2850 transcriptomic candidates was selected, among which 494 have been previously suggested as veraison main regulators according to profile or network analyses (Palumbo et al., 2014). Under veraison meta-QTLs (Chr 1 and 2) we found 61 of these transcriptomic candidates. These represent the genes most likely involved in the veraison genetic control mapped at these locations (Table 5). Heatmaps showing their expression along berry development are shown (Figure 4). Moreover, 12 of these were among the genes proposed as main regulators of berry veraison transition. Among the most notable candidate genes, comprised in the ver_2_1 and
ver_2_2 meta-QTL intervals, two pectin methylesterase inhibitor (PMEI) genes, previously reported to control pectin methylesterase activity in tomato (Di Matteo et al., 2005). Their function is supposedly to inhibit pectin methylesterase activity and then a premature berry softening related to pectin degradation (Lionetti et al., 2015). They may play a central role in the beginning of ripening by regulating some initial events at veraison, such as softening and loss of turgor (Gambetta et al., 2015). Another interesting candidate gene, again located within ver_2_2 meta-QTL interval, is the NAC (VvNAC13) transcription factor. This gene is believed to play an important role in the transition from the immature to the mature stage of grapevine; its role in development has been demonstrated in various plant species including grapevine (Wang et al., 2013). NAC transcription factors have been shown to regulate vegetative and reproductive development in Arabidopsis (Raman et al., 2008), tomato (Hendelman et al., 2013) and papaya, suggesting a significative role as hub genes for fruit ripening. Finally, a cluster of Myb genes within ver_2_3 meta-QTL interval, MYBA1, MYBA2, and MYBA3, support our approach; indeed, these genes have been previously extensively characterized for their role in the transition to berry ripening, by regulating the accumulation of anthocyanins in the berry skin (Kobayashi et al., 2004; Walker et al., 2007). Finally, a similar approach was also applied in order to try to prioritize genes located under veraison QTLs at other genetic locations. By applying the same strategy, the total number of original candidates underlying veraison QTLs was further reduced (8.2 times) allowing to build a priority list including 217 candidate genes (Figure 3, Table S11).

Table 5. Transcriptomic candidates underlying veraison meta QTLs. Meta-QTL The name assigned to the meta-QTL identified, Gene ID ID of the gene according to V1 annotation, Chr Chromosome, Start (bp) Start position in bp of the meta-QTL, End (bp) End position in bp of the meta-QTL, Annotation V1 Gene description according to V1 annotation, Transcriptomic candidate Citation of the original publication where transcriptomic data where taken to use for integration.

| MetaQTL | Gene ID | Chr | Start (bp) | End (bp) | Annotation_V1 | Transcriptomic candidate |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| ver_1_1 | VIT_01s0011g02840 | 1 | 2525946 | 2527902 | Cytochrome B561 | Palumbo et al 2014, <br> Massonet et al 2017 |
|  | VIT_01s0011g02880 | 1 | 2546007 | 2547946 | Cationic amino acid transporter 8 | Fasoli et al 2018 |
|  | VIT_01s0011g02950 | 1 | 2618690 | 2632669 | Zinc finger (C3HC4-type ring finger) | Fasoli et al 2018 |
|  | VIT_01s0011g03000 | 1 | 2688188 | 2688578 | No hit | Palumbo et al 2014, <br> Massonet et al 2017 |
|  | VIT_01s0011g03050 | 1 | 2717642 | 2719224 | Unknown protein | Palumbo et al 2014, <br> Massonet et al 2017 |
|  | VIT_01s0011g03070 | 1 | 2751566 | 2753036 | ERF/AP2 Gene Family (VvRAV1) | Palumbo et al 2014, <br> Massonet et al 2017 |
|  | VIT_01s0011g03180 | 1 | 2879352 | 2882108 | Lysine and histidine specific transporter | Palumbo et al 2014, <br> Massonet et al 2017 |
|  | VIT_01s0011g03210 | 1 | 2924829 | 2926924 | Aspartic Protease (VvAP1) | Palumbo et al 2014, <br> Massonet et al 2017 |
|  | VIT_01s0011g03360 | 1 | 3044557 | 3045849 | Unknown | Palumbo et al 2014, <br> Massonet et al 2017, <br> Fasoli et al 2018 |
|  | VIT_01s0011g03400 | 1 | 3076982 | 3080343 | Proton-dependent oligopeptide transport (POT) family protein | Palumbo et al 2014, <br> Massonet et al 2017 |


|  |  |  |  | Palumbo et al 2014, <br> Massonet et al 2017, <br> Fasoli et al 2018 |  |  |
| :--- | :--- | :--- | :--- | :--- | :--- | :--- |
|  | VIT_01s0011g03450 | 1 | 3127812 | 3132261 | Alpha-glucosidase | Fasoli et al 2018 |


|  | VIT_02s0154g00080 | 2 | 4813347 | 4818031 | Multi-copper oxidase (SKU5) | Palumbo et al 2014, <br> VIT_02s0154g00090 |
| :--- | :--- | :--- | :--- | :--- | :--- | :--- |
|  | 2 | 4824906 | 4827102 | Vacuolar invertase 2, GIN2 | Massonet et al 2017 |  |


|  |  |  |  |  | Palumbo et al 2014, <br> Massonet et al 2017, <br> Fasoli et al 2018 |  |
| :--- | :--- | :--- | :--- | :--- | :--- | :--- |
|  | VIT_02s0012g00500 | 2 | 6449814 | 6450360 | Invertase/pectin methylesterase inhibitor | Palumbo et al 2014, |
|  | VIT_02s0012g00550 | 2 | 6518911 | 6526733 | Inositol polyphosphate 5-phosphatase II | Massonet et al 2017 |


| VIT_02s0033g00450 | 2 | 14420525 | 14421283 | VvMybA3 | Palumbo et al 2014, <br> Massonet et al 2017, <br> Fasoli et al 2018 |
| :--- | :--- | :--- | :--- | :--- | :--- |
| VIT_02s0033g00700 | 2 | 15436012 | 15438008 | Nitrilase | Palumbo et al 2014, <br> Massonet et al 2017 |
| VIT_02s0033g00800 | 2 | 15606321 | 15608743 | Nitrilase 4 (NIT4) | Palumbo et al 2014, <br> Massonet et al 2017, <br> Fasoli et al 2018 |
| VIT_02s0033g01000 | 2 | 16093114 | 16095441 | Anthraniloyal-CoA: methanol anthraniloyal <br> transferase | Palumbo et al 2014, <br> Massonet et al 2017 |
| VIT_02s0033g01020 | 2 | 16190948 | 16193334 | Anthraniloyal-CoA: methanol anthraniloyal <br> transferase | Palumbo et al 2014, <br> Massonet et al 2017 |
| VIT_02s0033g01030 | 2 | 16252684 | 16254941 | Anthraniloyal-CoA: methanol anthraniloyal <br> transferase | Palumbo et al 2014, <br> Massonet et al 2017 |
| VIT_02s0033g01050 | 2 | 16299329 | 16301080 | Anthraniloyal-CoA: methanol anthraniloyal <br> transferase | Palumbo et al 2014, <br> Massonet et al 2017 |
|  |  |  |  |  |  |



Figure 4. Heatmaps of gene expression levels of the transcriptomics candidates included in the veraison and phenology meta-QTLs from the RNA-Seq datasets of Massonnett 2017 and Palumbo 2014. In this panel the veraison transcriptomic candidates in the Massonnett dataset.


Figure 4. Continued. Heatmaps of gene expression levels of the transcriptomics candidates included in the veraison and phenology meta-QTLs from the RNA-Seq datasets of Massonnett 2017 and Palumbo 2014. In this panel the veraison transcriptomic candidates in the Palumbo dataset.


Figure 4. Heatmaps of gene expression levels of the transcriptomics candidates included in the veraison and phenology meta-QTLs from the RNA-Seq datasets of Massonnett 2017 and Palumbo 2014. In this panel the phenology transcriptomic candidates in the Massonnett dataset.


Figure 4. Heatmaps of gene expression levels of the transcriptomics candidates included in the veraison and phenology meta-QTLs from the RNA-Seq datasets of Massonnett 2017 and Palumbo 2014. In this panel the phenology transcriptomic candidates in the Palumbo dataset.

## Discussion

A classical way to dissect the genetic determinism of grape phenology has been QTL studies. However, QTLs mapping often provides inconsistent results among studies, and huge genomic locations. A big advantage can derive from metaanalysis, which offers stronger evidence than individual studies, by revealing regions robustly associated with traits in multiple environments and genetic backgrounds. This approach has been already successfully exploited to improve and validate QTLs in several species, allowing insights into the genetic architecture of complex traits and paving the way for fine mapping and gene cloning. With this aim a genetic consensus map was built from 39 available SSR maps, including 3130 markers. By looking at marker distribution we observed they were not regularly spread along the chromosomes, but tended to concentrate in the middle regions, even though a good correlation was found with original maps. This is not surprising, reflecting a similar trend to original maps, due to suppression of recombination in centromeric regions. Other consensus maps already reported this drawback. Moreover, genetic positions of marker on the consensus arose from positions of shared markers according to the Biomercator software procedure, and were not based on recombination, since original genetic data are unfortunately not available from original maps. We fully agree that QTL meta-analysis would gain power and precision if raw genotypic and phenotypic data were made available. Recent advances in markers technology, with development of the NGS-based GBS technology in particular, have given a strong impulse to plant genotyping, and QTL studies now rely more on dense SNP maps. However, unshared markers do not allow for a direct genetic comparison of mapped QTLs, but require an indirect comparison through anchoring to the genome assembly. The distribution pattern of QTLs on chromosomes differs strongly between genetic and physical maps. Therefore, integration directly at genetic level could aid the improving of QTL location through co-location and meta-analysis, when feasible. Further comparisons can be then undertaken to newly generated QTLs relying on high throughput SNP maps, following anchoring to the genome assembly. Taking all this into account, we concluded that the consensus map we built constitutes a valuable reference, especially to the aim of integrating available genetic information, from
related QTL studies. Moreover, it will also provide a valuable instrument to enquire co-location with newly generated QTLs relying on dense SNP maps.

Taking advantage of this tool we have provided a compendium of all available QTL information that can be integrated at genetic level. Interestingly QTLs plotting revealed extensive co-locations across studies for each of the phenology related traits, besides downy mildew resistance, powdery mildew resistance, anthocyanin, drought stress, fertility, water use efficiency and growth, as well as for some berry and seeds related traits. However, studies addressing phenology are still few, negatively affecting the number of studies supporting each of the co-located QTLs. $R^{2}$ values of plotted QTLs, beside their distribution, suggest a highly polygenic nature for phenology related traits, with several QTLs involved, each of small effect, differently from other traits like pathogen resistance, seeds related traits and colour, all showing a more oligogenic architecture. More in detail, concerning veraison time four main regions located on LG1 and 2 have so far emerged consistently. Interestingly, plotting on a unique consensus map of QTLs also allows inspection of co-location across traits and categories, which is especially relevant for complex traits. In this way QTL meta-analysis also allows genetic correlation among traits to be investigated. In a previous work a second round meta-QTL analysis was proposed for seed yield QTLs and co-located yield associated QTLs in rapeseed, which allowed "indicator" meta-QTLs contributing to the complex trait crop yield to be defined. Indeed, QTL co-localization can be due to tight-linkage of QTL/genes playing different functions, but could also arise from pleiotropism. When pleiotropy is likely, it would also justify meta-analysis across traits, to further reduce the number of candidates. Veraison time is expected to be strictly related to other phenological stages. Accordingly, tests on the previously mentioned regions on LG1 and LG2 confirmed that, at least in some cases, comparable results are achieved when only veraison or all co-located phenology related traits are considered for the meta-analysis (see ver_2_1/pheno_2_1 as an example). We therefore also attempted a similar approach for veraison QTLs co-located with other phenology QTLs, finally identifying a number of regions, of which the most relevant were those located on LGs 14,16 and 18 . However, we are aware that these
rely on the pleiotropic assumption, which could be not always satisfied. A recent QTL study based on a GBS SNP map also addressed the mapping of veraison time. That study mainly aimed to discover and map stable QTLs across environments. A veraison QTL mapping on LG16 between 5 and 24 cM , which corresponded to the region between 2 and 16 Mbp , was found, but was not consistent across environments. Interestingly, it partially overlapped the pheno_16_2 meta-QTL we derived here starting from a veraison meta-QTL and its co-location to a floweringveraison interval QTL. Beside the detailed analysis of phenology traits, we have undertaken, our compendium now provides a useful tool for the inspection of colocation and meta-analysis for further traits in a similar way.

Transcriptomic studies have been also widely applied to characterize molecular changes associated to the onset of ripening, revealing, first of all, a massive transcriptomic rearrangement at veraison time. Among others, genes triggering such transition are expected to modulate their expression at this stage, although alternative regulative mechanisms cannot be excluded. We thus mined available transcriptomic profiles to i) identify the timing of such massive change, ii) select genes differentially expressed during this time in more varieties. Then, beside inspection of positional candidates underlying meta-QTLs, we propose to also integrate information about differential expression at veraison time, in order to prioritize candidates.

On LG1 a veraison time QTL was previously mapped. A more recent study also mapped a QTL for veraison at this location, which allowed us to define the ver_1_1 meta-QTL. Flowering QTLs consistently overlapped at same location suggesting a possible control of veraison time through regulation of flowering time. Accordingly, candidates for the flowering time control mapped under this metaQTL, like the PFT1 (phytochrome and flowering time 1) gene or a CONSTANS-like gene both controlling the photoperiodic flowering pathway in A. thaliana. However, integration of transcriptomic data allowed to pinpoint 14 candidates, among which the VvRAV1 transcription factor, belonging to the plant-specific $R A V$ (RELATED TO ABI3 AND VP1) family, was included. In Arabidopsis RAV1 was shown to act as negative regulator of both development and flowering, probably in
complexes with other co-repressors. Interestingly, some members of this gene family were shown to modulate developmental transitions, especially in response to temperature. Moreover RAV1 was also shown to be negatively regulated by brassinsteroid and abscisic acid, both hormones modulated at the onset of veraison time.

On LG2 meta-QTL analysis of overlapping veraison QTLs allowed 3 main regions to be spotted. In the first of these regions flowering QTLs were also plotted, again supporting a possible regulation of veraison time through flowering, even though no genes controlling flowering time where found under this locus. Interestingly, the orthologous of the Arabidopsis YABBY1/FIL transcription factor, which directly activate the AtMYB75, a key regulator of anthocyanin biosynthesis, was found among candidates selected by the integration of expression data. Moreover, by looking at other functional categories possibly related to veraison time, a gene encoding for a vacuolar invertase 2, key enzyme of sugar metabolism in fruits during ripening, a stay-green protein 1 gene related to a gene shown to be involved in ripening in tomato, beside two pectin methylesterase inhibitor (PMEI) genes, were found as differentially expressed. These last belong to a gene family previously characterized in grape. Their function is supposed to inhibit pectin methylesterase activity in pectin degradation and may play a role in the beginning of ripening by regulating initial events such as softening and loss of turgor. Interestingly, network analysis of gene expression profiles during berry ripening revealed PMEI among genes likely involved in triggering the major transcriptome reprogramming that occurs at veraison. Within ver_2_2 meta-QTL, the most notable candidate considering both positional and expression data was the VvNAC13 transcription factor. This gene belongs to a wide family of transcription factors in grapevine. Interestingly members of this family in tomato are involved in ethylene biosynthesis, reception and signalling during ripening. Moreover, they were also already suggested as playing a crucial role in berry transcriptome modulation associated to veraison, according to network analysis of berry expression profiles. However, in the same region, a gene encoding an atypical pseudo-response regulator (APRR2), involved in the circadian clock mechanism and contributing to fruit pigmentation and ripening in tomato, as well as two 1-
aminocyclopropane-1-carboxylate oxidases, taking part in ethylene biosynthesis and ripening were also selected by our approach and represent promising candidates. Lastly, a cluster of Myb genes locates within ver_2_3 meta-QTL interval. These genes have previously been extensively characterized for their involvement in the transition to berry ripening, by regulating the accumulation of anthocyanins in the berry skin. This finding, thus, supports our approach, even though these genes are unlikely to be themselves the early trigger of ripening onset. Other genomic regions were also proposed by previous studies for the genetic control of veraison time, among which the most relevant were mapping on LG 14, 16 and 18. By considering overlapping with other phenology related QTLs, followed by integration of transcriptomic data, we also selected candidates for these regions. The pheno_14_3 meta-QTL was computed from overlapping veraison QTL and flowering QTLs, and was accordingly highly enriched in candidates playing a role in the flowering transition control or fruit ripening, among which the most notable are Constans 2 (COL2), the feronia receptor-like kinase, a gene encoding a Brassinosteroid-6 oxidase, a gene encoding a COBRA protein and the putative MADS-box FRUITFULL 2. Interestingly this last gene was recently shown to also contribute to modulate the onset of ripening in tomato at early fruit development, beside its involvement at later ripening stages. Instead, a QTL previously mapped on LG 16, and explaining a large part of the genetic variance in the corresponding mapping population, partially co-localized to QTLs for the derived trait flowering-veraison interval, and with the genomic region involved in veraison recently identified by a SNP map and previously discussed. According to our strategy, the original interval was reduced to two regions of about 3.3 Mbp overall, including 15 transcriptomic candidates. Interestingly, more recently, the SSR marker UDV052, mapping under the pheno_16_3 meta-QTL close to the two candidates ABC transporter and an ERF transcription factor (19.1 Kbp and 56.9 Kbp respectively), was shown to be significantly associated to the early phenotype in a collection of different varieties, thus supporting our approach (Zyprian et al., 2018). Lastly, three different veraison QTLs were mapped on LG18. Two of them partially co-located with flowering QTLs from an independent study, and one of them was overlapping also with a QTL for the flowering-veraison interval. Under the derived
meta-QTLs, pheno_18_1 and pheno_18_2, spanning a still large region, we selected 74 transcriptomic candidates among which 19 were encoding proteins involved in regulation of gene expression, signalling or development. Candidates involved in carbohydrate metabolism, including especially a hexose (HT2) and a sucrose transporter (SUT2-2), putatively modulating sucrose signalling, or candidates encoding for genes for cell wall degradation (like a glucanase and a galactosidase, as examples), were also among those selected.

## Conclusions

By building a grape consensus genetic map anchored to the genome assembly a comprehensive overview about genomic distribution of several QTLs from published studies and their co-location both inside traits as well as across related traits was provided. Extensive co-localization was evident especially for phenology related traits. Four veraison meta-QTLs located on LG1 and 2, and several phenology meta-QTLs among which most relevant on LG 14, 16 and 18 were derived from 141 phenology related QTLs. Integration of meta-QTLs with expression data from prior transcriptomic studies allowed to select a set of 272 candidate genes for the genetic control of the veraison transition, reducing by about 20 and 10 times the genes proposed so far by either only genetic or transcriptomic approaches. Among these candidates 78 genes were involved in regulation of gene expression, signal transduction or development. Specific relevant candidates according to their annotation have been discussed. Further studies will now test and eventually validate the putative involvement of these candidates in the genetic control of the veraison transition during berry development.

## Chapter 2

## Exploit the genetic diversity of a grapevine collection for genetic association studies


#### Abstract

In plant organisms, genetic association studies still represent a valuable tool to uncover the genetic loci underlying specific quantitative traits. For grapevine phenology, such studies are still poor because of different reasons: the complex genetic architecture of the trait, the high levels of heterozygosity of the species, and its genomic characteristics (i.e. extent of linkage disequilibrium). By performing two complementary GWAS approaches on a grapevine germplasm collection, and by comparing the results with the results of the meta-analysis approach, we were able to identify and prioritize with high confidence genomic regions that could be regions of interest carrying the genes responsible for the beginning of the berry ripening, the veraison stage. In the first part of the chapter, we describe the genetic and phenotypic characterization of a wide germplasm collection of different grapevine accessions (CREA-VIT Conegliano collection) with the aim to identify the best panels of grapevine varieties, representing the most diverse genotypes and phenotypes for the traits of interest to be exploited to test the genotype-phenotype association. The second part of the chapter is dedicated to describe the association analysis itself with two different approaches, firstly using the GrapeReSeq 18K Vitis genotyping chip, then with an innovative approach called XP-GWAS (Yang et al., 2015) where pool of individuals displaying extreme phenotypes are wholegenome resequenced and allele frequencies compared to a random pool are screened for enrichment.


## Methods

## Plant material

The CREA-VIT Conegliano collection (hereafter CCC) is a grapevine germplasm collection comprising more than 2,000 grapevine accessions, among wine, table
cultivars, hybrids and rootstocks. The CCC has been extensively phenotyped for over 50 years for phenology traits (flowering time, veraison time, ripening time) and to a less extent for other various phenotypic traits like cluster architecture and wine acidity. The set of grapevine cultivars representing the object of our work consists in 617 unique genotypes (corresponding to 937 cultivars) for which phenotypic and genotypic data were complete and available. The list of the 617 genotypes is given in Supplementary Table 12.

## Genetic data and genetic diversity characterization

The genotypic data provided by our partners consisted in the genetic profiles of the 617 grapevine unique genotypes genotyped with 45 microsatellites (SSR) markers; of these, 32 were already published (Cipriani et al., 2010), while the remaining 13 have been developed and run subsequently. This additional set of microsatellite markers is composed of the 9 SSR markers used internationally for grapevine varietal identification (Maul et al., 2012) plus 4 additional markers developed by the researchers of the CREA-VIT to perform variety identification service (the ISV named markers and the VMCNG4B9). The protocols describing the generation of the SSR data are available from Cipriani. SSR markers were chosen to cover the whole genome of grapevine (19 chromosomes) with at least two markers per chromosome. To give an overview of the genetic diversity of the CCC different indexes commonly used in population genetics, such as expected heterozygosity, were calculated with the software Genalex v6.5.1 (Peakall and Smouse, 2012). Accessions displaying more than $20 \%$ of missing data were discarded, so the number of accessions used for the genetic diversity characterization and subsequent analysis was 530 .

## Population structure analysis

To explore the genetic variability of the collection and the population structure, the first method employed was PCA (principal component analysis) which helps to visualize the entire variability of the data in only two dimensions. PCA was performed on the $530 \times 45$ data set with the dudi.pca function of the $R$ package ade4, after centring and scaling, and replacing missing values with column mean.

Using this method, the alleles sizes are firstly converted to absence/presence ( $0 / 1$ ), so the PCA is not calculated directly on the allele sizes but on allele frequencies. The genetic diversity of the collection represented by the PCA was matched with the geographical origins (each cultivar was classified according to its geographic origin or region of cultivation as reported in the VIVC website http://www.vivc.de/index.php or http://catalogoviti.politicheagricole.it/catalogo.php) and grapes usage (wine vs table) of the individuals in the collection. The second method employed to explore population structure is the one included in the software STRUCTURE v2.3.4 which uses a Bayesian approach to assign a posterior probability to cluster individuals in sub-populations (Pritchard et al., 2000). Ten independent runs for K values ranging from 1 to 20 were performed with a burn-in length of $1,000,000$ followed by $1,500,000$ iterations. The admixture model was applied, and no prior population information was used. The best K was chosen based on the estimated membership coefficients $(\mathrm{Q})$ for each individual in each cluster. The optimal subpopulation model was investigated by applying the informal pointers (i.e. the plateau criterion) proposed by Pritchard et al.; individuals with a proportional membership $\mathrm{Q}>0.8$ were considered members of the group, while the remaining were considered admixed individuals. To validate the results from STRUCTURE software a nonparametric clustering approach called DAPC (Discriminant Analysis of Principal Components) implemented in the $R$ package adegenet 2.0.1 (Jombart and Collins, 2015) was also applied. This non-parametric approach makes different assumptions about the Hardy-Weinberg equilibrium of the genetic loci, so that it can be applied to very different ranges of populations. Prior clusters were identified by a sequential K-means clustering algorithm (find.clusters function) after data transformation by PCA. Then, a discriminant analysis (DA) used part of the principal components (PCs) to describe the clusters. K-means was ran with K varying from 1 to 20 and to ensure convergence we increased the number of starting points to 400 . The number of clusters was chosen based on the Bayesian Information Criterion (BIC).

## Phenotypic data

Phenotypic data consist of the flowering beginning and veraison beginning date in the last 13 years (2004-2016), with the value recorded as Julian day. Flowering beginning corresponds to the phenological stage when $10 \%$ of flowerhoods of a cluster have fallen, while veraison beginning corresponds to the stage when $10 \%$ of the berries of a cluster changed the colour from green to the particular colour of the cultivar, and this stage is usually considered the beginning of ripening. The interval from flowering to veraison, calculated as the difference between the two traits in number of days, was also included in the analysis. For each accession in the collection five plants are present, and all the plants are grafted on SO4 rootstock. The single phenotypic value per accession per year is obtained as the mean of the five plants. Beside raw data analysis, we also tried to understand the effect of temperature on phenological stages; we thus transformed raw flowering and veraison date values to a new value based on the GDD index. GDD (Growing Degree-Days) is an index used in agriculture that measures the heat accumulation throughout the year. The GDD value was calculated for each day of each year (2004-2016) for the Conegliano region and replaced the raw values (weather data source http://www.arpa.veneto.it/bollettini/storico/Mappa_2019_TEMP.htm). In this way the effect of temperature is normalized among years making the phenotypes more comparable. GDD is calculated as $G D D=\max \left(\frac{T_{\max }+T_{\text {min }}}{2}-T_{\text {base }}, 0\right)$, were temperatures below $10^{\circ} \mathrm{C}$ and above 30 ${ }^{\circ} \mathrm{C}$ are set to $10^{\circ} \mathrm{C}$ and $30^{\circ} \mathrm{C}$ respectively. GDD values were calculated for each day and then summed to obtain a value for each accession. Historical weather data were retrieved from ARPAV website. ARPAV is the regional agency for the environment protection of Veneto region. Data were recorded at a station located in the same area where the experimental vineyard is located. Phenotipic data are recorded every year according to the OIV descriptors. Phenological observations were recorded by visual inspection. All statistical analysis were performed with the statistical programming language $R$, version 3.4.4 (Team and R Development Core Team, 2016).

## Retrieval of published SNPchip data

We downloaded genotyping data publicly available from a recent publication (Laucou et al., 2018) from a French grapevine germplasm collection (Vassal collection) to be used as starting point for our GWAS study. Indeed, Laucou et al. released genotyping data for more than 700 grapevine accessions. Among these, 95 were in common, that is they had the same name, with the accessions of the CREAVIT collection. We obtained these data and explored the genetic diversity of this 95 accessions subset in our collection. More in detail, released data consisted of genotyping data of 10,207 solid SNPs with no missing data, obtained with the GrapeReSeq 18K Vitis genotyping chip.

## Core collection construction

Core collections construction was performed using the SSR data (530 accessions by 45 markers) with the software Core Hunter 3 (De Beukelaer et al., 2018). The software can select the smallest number of individuals maximizing different indexes. We applied a strategy of genetic diversity maximization, implemented in the software CoreHunter 3 (De Beukelaer et al., 2018), which allows to maximize allelic diversity of a collection, starting from a number of fixed entries. Core Hunter 3 can construct cores based on genetic marker data, phenotypic traits or precomputed distance matrices, optimizing one of many provided evaluation measures depending on the precise purpose of the core (e.g. high diversity, representativeness, or allelic richness). We decide to perform a maximization using allele coverage, i.e. the percentage of marker alleles observed in the full collection that are retained in the core, keeping as fixed the 95 cultivars in common with the GrapeReSeq dataset (Paslier M-C et al., 2013). Core Hunter was run 100 times to obtain the minimum number of unique individuals maximizing allele coverage. After obtaining this number, the individuals were ranked by occurrence and the most present individuals were selected to constitute the core collection. The $R$ package poppr (Kamvar et al., 2014) was used to evaluate allelic diversity of the core and compare it to the entire collection.

The DNA of the cultivars chosen with the core collection construction method was extracted with Qiagen DNeasy Plant Mini Kit (Qiagen, Hilden, Germany), according to the manufacturer's instructions from frozen leaf tissue. DNA was quantified with a NanoDrop. DNA was purified with Agencourt AMPure XP (Beckman Coulter) and sent for hybridization to Fondazione Edmund Mach. Genotype data were scored and validated from the GrapeReSeq 18K Vitis genotyping chip raw data using GenomeStudio Data Analysis v2011.1 (Illumina Inc, San Diego, CA, USA). Genotypes were called and for the subsequent analysis only the 10,207 solid SNPs comprised in the Vassal dataset were kept.

## Association analysis SNPchip

The association test was performed on the three phenotypic traits independently, flowering, veraison and flowering-veraison interval, both on mean value across years and on each independent year. Three different software were used: EMMAX (Kang et al., 2010), GAPIT (Lipka et al., 2012) and QTCAT (Klasen et al., 2016). For calculating the association with EMMAX, both no population structure correction and a correction based on kinship were tested, the latter obtained using all the 10K SNPs for the 132 individuals using the command \$kinship plink $-s-v$. Regarding GAPIT, a correction based on population structure was applied; the population structure included this time was the Q-matrix calculated with the 45 SSR markers and the software STRUCTURE. Regarding QTCAT, the software includes a new method that does not need population structure correction, so no other parameters other than genotypes and phenotypes were included. Linkage disequilibrium was evaluated on the 132 cultivars panel using the $R$ package LDcorSV (Mangin et al., 2012) that takes into account population structure and relatedness. The complete set of 10,207 SNPs was used for the evaluation.

## XP-GWAS pool design and DNA extraction

With this approach the panel selected for the association study from a germplasm collection is selected based on the phenotypic and not the genotypic diversity. Indeed, the distribution of the trait of interest was explored and the cultivars displaying extreme phenotypes were selected from the tails of the distribution. The
extreme phenotype GWAS approach (XP-GWAS) (Yang et al., 2015) expects that three different pools of individuals from a germplasm collection or a natural population are created based on the phenotypic distributions of the trait under study. In particular, the more the sample size and the standard deviation of the pools are similar the better. The three pools, hereafter called low, high and random pool, were designed from the flowering-veraison interval trait since it was the trait that displayed a more normal distribution. Starting from the 530 cultivars of the CCC, we sampled 48 and 47 cultivars from the low and high tail of the distribution respectively, and we randomly picked 38 cultivars from the entire collection. The distributions and statistics of the three pools were studied with the software $R$ 3.4.4. For each accession of the 3 pools, frozen leaves tissue was grinded and equal amounts of frozen powder were pooled together. DNA was extracted with Qiagen DNeasy Plant Mini Kit (Qiagen, Hilden, Germany), according to the manufacturer's instructions. Extracted DNA was resuspended in water and sent to the genomics facility for quality control and sequencing. Genomic DNA samples were quantified with the Qubit dsDNA HS Assay kit (Life Technologies). DNA purity and integrity were assessed at the Nanodrop 1000 spectrophotometer (Thermo Scientific) and by capillary electrophoresis on a 2200 TapeStation (Agilent Technologies), respectively. For each pool, 500ng DNA was sheared into 350bp fragments by sonication using a S220 Covaris Ultrasonicator. The fragmented DNA was then used for the preparation of Whole Genome libraries following the Kapa Hyper Prep PCR-free workflow (KAPA Biosystems) according to manufacturer's instructions with minor modifications. To avoid index-hopping during sequencing, libraries were treated with Illumina Free Adapter Blocking Reagent (Illumina) prior to further processing. DNAseq libraries were pooled at equimolar concentration and sequenced over 3 lanes of an Illumina HiSeqX sequencer using 150bp paired-end reads.

## Results and discussion

## CCC genetic diversity characterization and population structure description

The CREA-VIT Conegliano collection (CCC) is a wide grapevine germplasm collection including more than 2,000 grapevine accessions. The cultivars for which
complete phenotypic information on phenology and genotypic information at 45 microsatellite markers was available (Cipriani et al., 2010) were 617. We first used the latter information to study the genetic diversity and the population structure of this collection subset. Full information about country of origin, cultivation, berry colour and usage, retrieved either from http://www.vivc.de/index.php or http://catalogoviti.politicheagricole.it/catalogo.php are provided in Supplementary Table 11 and Figure 5.


Figure 5. First panel, top: country of origin of the 530 grapevine accessions, when not available, replaced with the area of major cultivation. Second panel, bottom: destination usage of the grape varieties.

The majority of cultivars of the CCC are of Italian origin, especially from north and centre of Italy, followed by cultivars of western Europe origin. Concerning the destination usage of the grapes, the majority ( $77 \%$ ) are cultivars designated to wine production. The 530 distinct genotypes characterized with 45 SSR markers (Cipriani et al., 2010 and personal communication) revealed a large number of alleles detected (Table 7). The number of different alleles (A) for the SSRs was 369 and ranged from 2 to 16 per locus, with an average of 8.2. The observed $(H o)$ and
expected heterozygosities $(\mathrm{He})$ were 0.658 and 0.75 , respectively, and the former is in line with the value in other grapevine collections (Nicolas et al., 2016).

Table 7. Summary statistics of genetic variation at 45 SSR loci in the 530 germplasm cultivars. In the column locus, in green the previously reported SSR markers (Cipriani et al., 2010), in yellow the new set of markers. $\boldsymbol{N}$ (Sample Size), Na (No. Alleles), Ne (No. Effective Alleles), I (Information Index), Ho (Observed Heterozygosity), He (Expected Heterozygosity), uHe (Unbiased Expected Heterozygosity), F (Fixation Index).

| Locus | N | Na | Ne | I | Ho | He | uHe | F |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| VChr1a | 495 | 9.000 | 2.737 | 1.400 | 0.655 | 0.635 | 0.635 | -0.031 |
| VChr1b | 526 | 4.000 | 2.728 | 1.156 | 0.671 | 0.633 | 0.634 | -0.060 |
| VChr1c | 528 | 3.000 | 1.973 | 0.724 | 0.523 | 0.493 | 0.494 | -0.060 |
| VChr2a | 525 | 4.000 | 1.859 | 0.771 | 0.520 | 0.462 | 0.462 | -0.126 |
| VChr2b | 497 | 4.000 | 1.748 | 0.772 | 0.443 | 0.428 | 0.428 | -0.035 |
| VChr3a | 487 | 12.000 | 6.275 | 2.026 | 0.786 | 0.841 | 0.841 | 0.064 |
| VChr4a | 475 | 6.000 | 2.448 | 1.135 | 0.589 | 0.592 | 0.592 | 0.003 |
| VChr5a | 520 | 11.000 | 5.149 | 1.880 | 0.806 | 0.806 | 0.807 | 0.000 |
| VChr5b | 513 | 9.000 | 3.704 | 1.517 | 0.754 | 0.730 | 0.731 | -0.033 |
| VChr5c | 450 | 7.000 | 4.215 | 1.623 | 0.760 | 0.763 | 0.764 | 0.004 |
| VChr6a | 509 | 3.000 | 2.037 | 0.846 | 0.417 | 0.509 | 0.510 | 0.182 |
| Vchr7a | 524 | 3.000 | 1.982 | 0.723 | 0.508 | 0.495 | 0.496 | -0.025 |
| VChr7b | 469 | 4.000 | 3.287 | 1.277 | 0.736 | 0.696 | 0.697 | -0.057 |
| VChr8a | 498 | 12.000 | 5.702 | 1.918 | 0.588 | 0.825 | 0.825 | 0.287 |
| VChr9a | 448 | 7.000 | 5.059 | 1.741 | 0.850 | 0.802 | 0.803 | -0.060 |
| VChr9b | 395 | 13.000 | 4.385 | 1.760 | 0.324 | 0.772 | 0.773 | 0.580 |
| Vchr10b | 512 | 4.000 | 2.831 | 1.083 | 0.670 | 0.647 | 0.647 | -0.036 |
| Vchri1a | 511 | 5.000 | 2.039 | 0.955 | 0.485 | 0.510 | 0.510 | 0.048 |
| Vchri2a | 525 | 8.000 | 2.566 | 1.204 | 0.547 | 0.610 | 0.611 | 0.104 |
| Vchr12b | 482 | 3.000 | 1.385 | 0.465 | 0.295 | 0.278 | 0.278 | -0.060 |
| Vchri3a | 504 | 7.000 | 3.281 | 1.479 | 0.704 | 0.695 | 0.696 | -0.013 |
| Vchr 13b | 511 | 8.000 | 2.558 | 1.342 | 0.614 | 0.609 | 0.610 | -0.009 |
| Vchri3c | 513 | 4.000 | 3.141 | 1.258 | 0.700 | 0.682 | 0.682 | -0.027 |
| Vchr14a | 403 | 5.000 | 2.294 | 0.956 | 0.600 | 0.564 | 0.565 | -0.065 |
| Vchr15a | 511 | 7.000 | 3.387 | 1.455 | 0.663 | 0.705 | 0.705 | 0.059 |
| Vchr16a | 525 | 8.000 | 1.507 | 0.734 | 0.250 | 0.336 | 0.337 | 0.258 |
| Vchr16b | 523 | 8.000 | 2.086 | 1.144 | 0.530 | 0.521 | 0.521 | -0.017 |
| Vchri7a | 488 | 2.000 | 1.781 | 0.630 | 0.281 | 0.439 | 0.439 | 0.360 |
| Vchr18a | 519 | 9.000 | 3.458 | 1.553 | 0.570 | 0.711 | 0.712 | 0.198 |
| Vchr 18 b | 529 | 5.000 | 2.829 | 1.146 | 0.681 | 0.647 | 0.647 | -0.053 |
| Vchr19a | 520 | 9.000 | 3.061 | 1.471 | 0.673 | 0.673 | 0.674 | 0.000 |
| Vchr 19 b | 523 | 5.000 | 3.281 | 1.306 | 0.562 | 0.695 | 0.696 | 0.191 |
| VVS2 | 529 | 13.000 | 5.504 | 1.964 | 0.822 | 0.818 | 0.819 | -0.005 |
| VVMD27 | 529 | 9.000 | 5.615 | 1.827 | 0.824 | 0.822 | 0.823 | -0.003 |
| VVMD7 | 529 | 14.000 | 5.320 | 1.907 | 0.822 | 0.812 | 0.813 | -0.013 |
| ISV2 | 525 | 16.000 | 6.641 | 2.061 | 0.891 | 0.849 | 0.850 | -0.049 |
| VrZAG62 | 530 | 10.000 | 6.383 | 1.979 | 0.860 | 0.843 | 0.844 | -0.020 |
| VVMD5 | 524 | 10.000 | 6.098 | 1.949 | 0.819 | 0.836 | 0.837 | 0.021 |
| VMCNG4B9 | 524 | 15.000 | 6.231 | 2.139 | 0.853 | 0.840 | 0.840 | -0.016 |
| VrZAG79 | 527 | 11.000 | 5.667 | 2.018 | 0.827 | 0.824 | 0.824 | -0.005 |
| ISV3 | 488 | 10.000 | 3.119 | 1.361 | 0.836 | 0.679 | 0.680 | -0.231 |
| ISV4 | 520 | 10.000 | 4.682 | 1.693 | 0.754 | 0.786 | 0.787 | 0.041 |
| VVMD28 | 526 | 14.000 | 7.751 | 2.207 | 0.882 | 0.871 | 0.872 | -0.013 |
| VVMD25 | 443 | 14.000 | 4.434 | 1.669 | 0.786 | 0.774 | 0.775 | -0.014 |
| VVMD32 | 428 | 15.000 | 5.864 | 1.962 | 0.862 | 0.829 | 0.830 | -0.039 |
| Mean | 501.8 | 8.200 | 3.780 | 1.426 | 0.658 | 0.675 | 0.676 | 0.027 |
| SE | 5.088 | 0.580 | 0.252 | 0.071 | 0.026 | 0.022 | 0.022 | 0.020 |

To explore the genetic variability of the collection and population structure, the first method employed was the PCA. The genetic diversity of the collection represented by the PCA was matched with the geographical origins and grapes usage of the cultivars in the collection. Clustering was weak concerning the geographic subdivision (weak gradient right to left, eastern Europe to north Italy/center-north Europe), while the clustering based on usage was more evident (Figure 6), displaying a sort of gradient from right to left (table to wine cultivars), with the wine/table cultivars in the middle of the distribution, similar to what has been reported before for other collections (Migicovsky et al., 2017).



Figure 6. Principal component analysis plots from SSR data. Each dot represents one of the 530 cultivars. Top plot represents geographic origin/cultivation, while bottom is grape usage destination. Top panel abbreviations: AM America, CI Center Italy, CNE Center North Europe, EE Eastern Europe, IS Islands (Sardinia, Sicily), IT Italy, NI North Italy, SI South Italy, UKN Unknown origin, WE Western Europe. Bottom panel abbreviations: T/R Table/Raisin, UKN Unknown origin, W/T Wine/Table, W/T/R Wine/Table/Raisin.

Population structure analysis performed with the software STRUCTURE on the SSR data set suggested as the most likely number of clusters (K), evaluated with the plateau criterion proposed by Pritchard et al. and the $\Delta \mathrm{K}$ method (Evanno et al., 2005) a maximum probability for $K=2$ and a smaller probability for $K=6$. Indeed, the $\Delta \mathrm{K}$ criterion gave the highest value at $\mathrm{K}=2$, and a small peak at $\mathrm{K}=6$. Instead, the plateau criterion, highlighted by the mean log-likelihood curve, revealed a maximum value with no standard deviation associated at $\mathrm{K}=6$, beyond that, a sort of plateau is reached, and the standard deviations associated with the remaining estimates increase. When the cultivars in the PCA are coloured according to the membership of the groups identified by STRUCTURE, the 2 groups subdivision is very evident (Figure 8). Table 8 gives the genetic diversity statistics associated to each one of the 2 subgroups and the admixed one, compared to the values (already reported in Table 7) obtained considering the entire collection.


Figure 7. Estimated number of clusters obtained with STRUCTURE for $K$ values from 1 to 20. Graphical representation of its derivative statistics $\Delta K$ (top) and estimated mean $L(K)$ (middle). Bottom figure represents plot generated based on the Q-matrix. Each cultivars is represented by a single vertical line, which is divided in coloured segments in proportion to the estimated membership in the two subgroups. On the y-axes is the likelihood of assignment to any given cluster $K$.


Figure 8. Principal component analysis plot from SSR data. Each dot represents one of the 530 cultivars. The colors represent the 2 groups subdivision identified by STRUCTURE.

Table 8. Summary statistics of genetic variation at 45 SSR loci in the 530 germplasm cultivars, subdivided into 2 groups as identified by STRUCTURE, compared to the values obtained considering the entire collection.

| Pop |  | $\mathbf{N}$ | $\mathbf{N a}$ | $\mathbf{N e}$ | $\mathbf{I}$ | $\mathbf{H o}$ | $\mathbf{H e}$ | $\mathbf{u H e}$ | $\mathbf{F}$ |
| :--- | :--- | ---: | ---: | ---: | ---: | ---: | ---: | ---: | ---: |
| A-STR | Mean | 148.778 | 7.378 | 3.736 | 1.409 | 0.655 | 0.671 | 0.673 | 0.025 |
|  | SE | 1.925 | 0.488 | 0.252 | 0.070 | 0.026 | 0.022 | 0.023 | 0.022 |
|  |  |  |  |  |  |  |  |  |  |
| B-STR | Mean | 131.644 | 7.378 | 3.745 | 1.420 | 0.663 | 0.675 | 0.677 | 0.017 |
|  | SE | 1.239 | 0.501 | 0.249 | 0.071 | 0.026 | 0.022 | 0.022 | 0.020 |
|  |  |  |  |  |  |  |  |  |  |
|  |  |  |  |  |  |  |  |  |  |
|  | Mean | 221.356 | 7.467 | 3.757 | 1.419 | 0.656 | 0.674 | 0.676 | 0.029 |
|  | SE | 2.027 | 0.481 | 0.246 | 0.070 | 0.026 | 0.023 | 0.023 | 0.021 |
|  |  |  |  |  |  |  |  |  |  |
| CCC | Mean | 501.8 | 8.2 | 3.78 | 1.426 | 0.658 | 0.675 | 0.676 | 0.027 |
|  | SE | 5.088 | 0.58 | 0.252 | 0.071 | 0.026 | 0.022 | 0.022 | 0.02 |

Homogeneity of genetic diversity between the three subgroups is appreciable, and the values are comparable to the ones of the entire collection; however, pairwise Fst values (fixation index or inbreeding coefficient), which describe the genetic differentiation between groups or population, indicates a very low level of differentiation between the three subgroups ( 0.0026 between A-STR and B-STR, 0.0019 between A-STR and ADMIXED, and 0.0021 between B-STR and
admixed), although very similar to values previously reported separating sativa cultivars of among Western, Central and Eastern Europe (Myles et al., 2011). We tried to characterize the subgroups to find specific features of the two, so we looked at enrichment in terms of geographic origins, and found a significative overrepresentation of cultivars from the east (mostly table grapes), the orientalis group (proles) as defined in the works of Negrul and Troshin (Negrul et al., 1946; Troshin LP et al., 1990), in the group A-STR, and a significant underrepresentation of cultivars from the North Italy group (Table 9). Regarding group B-STR, we found a significant overrepresentation of cultivars belonging to the North Italy group and a slight enrichment for cultivars belonging to the Center-North Europe group ( $p=0.061$ two-tailed, $p=0.0554$ one-tailed Fisher's Exact Test) comprised in the pontica occidentalis proles. On the other hand, in this group we observed very few cultivars belonging to groups overrepresented in group A-STR (AM, EE, IS [2 cultivars]).

Table 9. Composition of the two groups identified by STRUCTURE plus the admixed cultivars. Top table shows contingency table for country of origin, bottom table for colour. In red overrepresented groups, in blue underrepresented ones. In parenthesis, in italics are reported p-values from Fisher's Exact Test two-tailed. Abbreviations: AM America, CI Center Italy, CNE Center North Europe, EE Eastern Europe, IS Islands (Sardinia, Sicily), IT Italy, NI North Italy, SI South Italy, UKN Unknown origin, WE Western Europe.

| COUNTRY | A-STR | B-STR | ADMIXED | CCC |
| :--- | ---: | ---: | ---: | ---: |
| AM | $\mathbf{8}(0.09445)$ | 0 | $3(0.5692)$ | 11 |
| CI | $42(0.8417)$ | $34(1)$ | $58(1)$ | 134 |
| CNE | 0 | $8(0.06159)$ | $5(1)$ | 13 |
| EE | $\mathbf{1 3}(0.0285)$ | 0 | $5(0.4912)$ | 18 |
| IS | $\mathbf{1 5 ( 0 . 0 2 7 4 )}$ | $2(0.1956)$ | $5(0.2069)$ | 22 |
| IT | $27(0.7169)$ | $\mathbf{8}(0.006577)$ | $48(0.1853)$ | 83 |
| NI | $\mathbf{6 ( 3 . 4 3 1 e - 0 8 )}$ | $\mathbf{6 9}(9.71 e-05)$ | $56(0.9294)$ | 131 |
| SI | $20(0.4707)$ | 0 | $34(0.1449)$ | 54 |
| UKN | $3(0.7049)$ | $2(1)$ | $2(0.73)$ | 7 |
| WE | $25(0.1696)$ | $15(1)$ | $17(0.1862)$ | 57 |
|  | $\mathbf{1 5 9}$ | $\mathbf{1 3 8}$ | $\mathbf{2 3 3}$ | $\mathbf{5 3 0}$ |


| COLOR | A-STR | B-STR | ADMIXED | CCC |
| :--- | ---: | ---: | ---: | ---: |
| BLANC | $86(0.1308)$ | $\mathbf{3 8}(0.03239)$ | $100(0.9427)$ | 224 |
| GRIS | 0 | 0 | 2 | 2 |
| NOIR | $\mathbf{5 1}(0.009774)$ | $\mathbf{9 6}(0.04294)$ | $121(0.8398)$ | 268 |
| RED | 2 | 0 | 1 | 3 |
| ROSE | 2 | 0 | 1 | 3 |
| ROUGE | 4 | 3 | 1 | 8 |
| UKN | 14 | 1 | 7 | 22 |
|  | $\mathbf{1 5 9}$ | $\mathbf{1 3 8}$ | $\mathbf{2 3 3}$ | $\mathbf{5 3 0}$ |

When the same reasoning is applied to the colour characteristic, we observe an underrepresentation of black-skinned (noir) cultivars in the A-STR, coupled with an overrepresentation of the same in the B-STR.

To support the results obtained by STRUCTURE we also applied a non-parametric approach called DAPC to the same SSR dataset (530 individuals, 45 SSRs). To identify the optimal number of clusters the Bayesian Information Criterion (BIC) method was employed (Jombart et al., 2010). The greatest delta between two points is found between $\mathrm{K}=1$ and $\mathrm{K}=2$, while the lowest value is reached at $\mathrm{K}=7$. Both $K$ values ( 2 and 7 ) were explored and compared to the results obtained previously with STRUCTURE (Figure 9).


Figure 9. Bayesian Information Criterion (BIC) according to the number of inferred clusters ( $K=0-20$ ). 300 principal components were kept and the number of starting points was set at 100 . Number of iterations was 1000000 and the chosen number of clusters were $K=2$ and $K=7$.

When the individuals belonging to the subgroups identified by STRUCTURE are matched with the individuals belonging to the subgroups identified by DAPC there is a perfect match (Figure 10, top). The individuals classified as admixed by STRUCTURE (group c in the figure) are split between the 2 groups identified by DAPC, which usually identifies a very small number of admixed individuals, in this case just 3 (not shown). Moreover, the two DAPC groups separates very good in the PCA plot, meaning that their genetic diversity is high (Figure 10, bottom).


Figure 10. Top figure. Representation of contingency table by square sizes to compare prior group assignments using $K$-means clustering to the groups identified by STRUCTURE. Group a is $A$-STR, $b$ is $B$-STR and $c$ is admixed. Inferred clusters 1 and 2 are the two clusters identified by DAPC approach. Bottom panel is the discriminant analysis component plot for the two inferred clusters by DAPC.

Another way of assessing the results of the clustering method we employed is with phylogenetic trees and dendrograms. Using the $R$ package phangorn $v 2.1 .1$ we built a Neighbour-Joining tree based on Euclidean distances and a dendrogram based on Manhattan distances (Figure 11). In both, the 2 subgroups identified by STRUCTURE and DAPC (data not shown) and the groups founds by the trees were in agreement except for few individuals incorrectly placed, thus reinforcing the two groups subdivision identified initially by STRUCTURE.


Figure 11. Top panel, Neighbor-joining tree based on euclidean distances. In red, green and blue group ASTR, B-STR and admixed cultivars respectively. Bottom panel, dendrogram based on manhattan distances, Ward method. Red labels represent group A-STR, black labels are cultivars from group B-STR. Admixed individuals are not displayed.

We then tried to explore the additional subdivision of the population in 6 subgroups as previously suggested by STRUCTURE (Figure 7). When the PCA of the 530 cultivars is coloured according to the 6 subgroups (Figure 12), except for groups $d$
and $e$, which are slightly overlapping, the remaining 4 groups are well separated. Admixed cultivars are the majority $(316, \sim 60 \%)$ and are dispersed all over the points, but mostly concentrate in the middle of the plot. The smallest group is group $c(\mathrm{n}=10)$, while group $f$ is the largest, with 67 cultivars. In Table 10 summary statistics of the genetic diversity of the 6 subgroups are reported. Values are again comparable to each other's and to the entire collection. Pairwise fixation index ranged from 0.019 between $d$ and $f$ and 0.096 between $b$ and $c$ subpopulations (Table 11). Considering the small sample size compared to other grapevine germplasm collections (Emanuelli et al., 2013; Nicolas et al., 2016), these values, with an average value of 0.051 , may reflect statistically supported subpopulations. Also in this case we tried to characterize the subpopulations from a geographic (Table 12) and grape usage point of view. We notice some over- and underrepresentation: for example group $a$ is enriched in cultivars of the proles orientalis, very similarly to the previous group A-STR, while at the same time is scarce of cultivars from the north of Italy; group $b$ is very small ( $\mathrm{n}=12$ ), but being very close to group $a$ is again slightly enriched for cultivars from classified as America (Cardinal, Early Muscat, July Muscat, mainly table grape cultivars). Group $c$ is the smallest $(\mathrm{n}=10)$ composed only of wine cultivars and not enriched in cultivar from any geographic area; group $d$ is composed mainly by wine cultivars and is enriched in cultivars of north Italian origin and, though not significant, the other representatives are only cultivars from centre of Italy. Group $e$, similar to group $d$, is enriched in cultivars of centre-Italy origin (though significant only at one-tail), and even if almost overlapping to group $d$, it also displays overrepresentation of western European cultivars (in particular 7 wine French cultivars: Becuet, Pinot Blanc, Malbech, Merlot, Sauvignon, Semillon, Chenin). Group $f$ is significantly enriched for centre and south Italian cultivars, while it comprises only one cultivar classified as North Italian, and, according to the PCA plot, it is the only subgroup with no cultivars overlapping to any other subgroups. By building a Neighbor-joining phylogenetic tree without considering the admixed individuals identified by STRUCTURE, the 6 subgroups identified both by STRUCTURE separates well supporting again the hypothesis of the 6 groups subdivision (Figure 13 B).


Figure 12. Principal component analysis plot from SSR data. Each dot represents one of the 530 cultivars. The colors represent the 6 groups subdivision identified by STRUCTURE. Group $g$ is admixed cultivars, colored in light grey.

Table 10. Summary statistics of genetic variation at 45 SSR loci in the 530 germplasm cultivars, subdivided into 6 groups as identified by STRUCTURE, compared to the values obtained considering the entire collection. Group $g$ is admixed cultivars.

| Pop |  | N | Na | Ne | I | Ho | He | uHe | F |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| a | Mean | 40.556 | 6.489 | 3.649 | 1.375 | 0.667 | 0.661 | 0.669 | -0.009 |
|  | SE | 0.548 | 0.434 | 0.255 | 0.071 | 0.026 | 0.023 | 0.023 | 0.023 |
| b | Mean | 11.222 | 4.822 | 3.421 | 1.275 | 0.648 | 0.653 | 0.685 | 0.013 |
|  | SE | 0.220 | 0.278 | 0.220 | 0.062 | 0.032 | 0.022 | 0.023 | 0.033 |
| c | Mean | 9.400 | 4.556 | 3.252 | 1.230 | 0.664 | 0.634 | 0.670 | -0.035 |
|  | SE | 0.140 | 0.257 | 0.197 | 0.062 | 0.035 | 0.023 | 0.025 | 0.038 |
| d | Mean | 52.511 | 6.667 | 3.765 | 1.404 | 0.662 | 0.674 | 0.680 | 0.013 |
|  | SE | 0.564 | 0.402 | 0.253 | 0.069 | 0.026 | 0.022 | 0.023 | 0.023 |
| e | Mean | 23.911 | 5.844 | 3.481 | 1.323 | 0.618 | 0.647 | 0.661 | 0.046 |
|  | SE | 0.439 | 0.358 | 0.238 | 0.070 | 0.032 | 0.024 | 0.025 | 0.034 |
| f | Mean | 63.133 | 6.844 | 3.719 | 1.401 | 0.662 | 0.672 | 0.677 | 0.015 |
|  | SE | 0.795 | 0.452 | 0.248 | 0.070 | 0.026 | 0.022 | 0.022 | 0.023 |
| g | Mean | 301.044 | 7.778 | 3.767 | 1.425 | 0.658 | 0.675 | 0.676 | 0.027 |
|  | SE | 2.664 | 0.512 | 0.247 | 0.070 | 0.026 | 0.022 | 0.023 | 0.021 |
| CCC | Mean | 501.8 | 8.2 | 3.78 | 1.426 | 0.658 | 0.675 | 0.676 | 0.027 |
|  | SE | 5.088 | 0.58 | 0.252 | 0.071 | 0.026 | 0.022 | 0.022 | 0.02 |

Table 11. Pairwise $F_{\text {st }}$ index (calculated as $1-(\mathrm{Ho} / \mathrm{He})$ ) between the six subgroups subdivision as identified by STRUCTURE. Highest and lowest values highlighted in red and blue respectively, excluding admixed individuals.

| Pop | a | b | c | d | e | f |
| :--- | ---: | ---: | ---: | ---: | ---: | ---: |
| b | 0.055 |  |  |  |  |  |
| c | 0.066 | $\mathbf{0 . 0 9 6}$ |  |  |  |  |
| d | 0.022 | 0.052 | 0.055 |  |  |  |
| e | 0.036 | 0.067 | 0.081 | 0.037 |  |  |
| f | 0.023 | 0.045 | 0.066 | 0.019 | 0.037 |  |
| g | 0.016 | 0.045 | 0.060 | 0.013 | 0.032 | 0.010 |

Table 12. Contingency table of geographic composition of the six groups identified by STRUCTURE plus the admixed cultivars. In red overrepresented groups, in blue underrepresented ones. In parenthesis, in italics are reported p-values from Fisher's Exact Test two-tailed.


Also in this case, to validate STRUCTURE results, we tried the DAPC approach which suggested a possibility having a 7 subgroups subdivision (Figure 9). When we tried to match the composition of individuals identified by both the methods, this time the match is not perfect but is still significant. In particular for groups $a$, $b$, and $e$, the match is almost perfect with DAPC clusters 3,6 and 7 respectively (Figure 13 C ). The plot of the discriminant components based on the DAPC approach shows that not all the 7 groups were clearly separated from each other (Figure 13 D ).


D


Figure 13. A. Plot generated based on the Q-matrix. Each cultivar is represented by a single vertical line, which is divided in coloured segments in proportion to the estimated membership in the six subgroups identified by STRUCTURE. On the y-axes is the likelihood of assignment to any given cluster K. B. Neighbor-joining tree based on euclidean distances. Each colours represent a different subgroups of the 6 identified by STRUCTURE. C. Contingency table by square sizes to compare prior group assignments using K-means clustering to the groups identified by STRUCTURE. Groups a to $g$ are 6 groups identified by STRUCTURE, while $g$ are admixed cultivars. Inferred clusters 1 to 7 are the 7 clusters identified by DAPC approach. D. Scatter plot of the first two discriminant components as evaluated by the dapc function of the adegenet $R$ package. Black crosses indicate the centre of each one of the inferred clusters.

## Phenotypic data

Phenotypic data consisting of Julian days (days from the beginning of the year) for the beginning of flowering (FB) and the beginning of veraison (VB) were recorded for the whole CCC for over 50 years. Flowering-veraison interval trait (F-V) was also considered and obtained as the raw difference between the time of veraison beginning and flowering beginning. We focused our analysis on the data of seasons 2002 to 2016, given the observation (Tomasi et al., 2011) that a significant breakpoint in the grapevine phenology in the Conegliano region occurred around 20 years ago. Complete phenotypic data were available for 13 years for FB trait, while for VB , and consequently also $\mathrm{F}-\mathrm{V}$, for 11 years.

When we checked the single years distribution of flowering, many years showed a bimodal distribution, as well as the all-years distribution (Figure 14, Panel 1). We then checked the distribution of the traits with histograms and quantile-quantile plots (QQ plots). Flowering suggests a bimodal distribution while veraison approximates well a normal distribution. Raw phenotypic values were then plotted for each year as boxplot in order to check the variability for the three traits among the 13 years. Variability among years looks greater for flowering than for veraison, though quite significant for both (p-value $\ll 0.01$, ANOVA), showing a slightly decreasing trend over the years. The inter-years variability looks more levelled when the trait flowering-veraison interval is considered, though again still significant when an ANOVA test is performed to compare all the years together (pvalue $\ll 0.01$ ). Since it is important to consider the effect of temperature on phenological stages of plants, and in particular in grapevine (De Cortázar-Atauri et al., 2017; Duchêne et al., 2010; Williams et al., 1985) we wanted to verify the effect of temperature on the two traits, performing a transformation of the raw flowering and veraison date values to a new value based on the GDD index. GDD (Growing Degree-Days) is an index used in agriculture that measures the heat accumulation throughout the year. After plotting the newly GDD- transformed values the distribution of flowering changed from bimodal to a good normal distribution, while the veraison distribution remained unchanged (Figure 14, Panel 2); especially from the QQ plots it can be observed that when using the GDD values almost all
the points fall on the theoretical quantile distribution except for few points at the flowering tails. The same effect can be seen on the flowering-veraison trait, and it can be noted that when considering the GDD values, the entire distribution profile of the F-V interval trait is completely determined by the VB trait profile. Such a decrease in variability among years when using the GDD corrected traits was also confirmed by a comparison of the $p$-values intensities from a pairwise $t$-test between every year (Figure 15). Overall, the $p$-values of the comparison of the means lose significance when considering GDD values, both for flowering and for veraison. Finally, we plotted the traits (both as raw date and GDD-corrected dates) for each genotype as boxplot (Data not shown) to check the intra- and intergenotypic variability and could observe that both decreased after the GDD correction for the flowering trait while remained similar for veraison.



| Flowering |  |  |  |  |
| :--- | :---: | :---: | :---: | :---: |
| YEAR | Count | Mean | Median | SD |
| $\mathbf{2 0 0 4}$ | 263 | 163.21 | 164 | 2.33 |
| $\mathbf{2 0 0 5}$ | 268 | 153.64 | 154 | 3.33 |
| $\mathbf{2 0 0 6}$ | 368 | 157.85 | 158 | 3.19 |
| $\mathbf{2 0 0 7}$ | 288 | 137.96 | 138 | 3.00 |
| $\mathbf{2 0 0 8}$ | 564 | 155.27 | 155 | 2.46 |
| $\mathbf{2 0 0 9}$ | 506 | 144.17 | 144 | 2.10 |
| $\mathbf{2 0 1 0}$ | 570 | 154.92 | 155 | 2.27 |
| $\mathbf{2 0 1 1}$ | 506 | 143.35 | 143 | 2.07 |
| $\mathbf{2 0 1 2}$ | 526 | 153.80 | 154 | 3.44 |
| $\mathbf{2 0 1 3}$ | 595 | 159.50 | 159 | 3.21 |
| $\mathbf{2 0 1 4}$ | 568 | 146.72 | 147 | 3.41 |
| $\mathbf{2 0 1 5}$ | 583 | 150.33 | 151 | 3.46 |
| $\mathbf{2 0 1 6}$ | 524 | 156.56 | 156 | 3.62 |
| $\mathbf{A L L}$ | 6129 | 152.08 | 154 | 6.999 |





| Veraison |  |  |  |  |
| :--- | ---: | ---: | ---: | ---: |
| YEAR | Count | Mean | Median | SD |
| $\mathbf{2 0 0 4}$ | 263 | 226.29 | 227 | 6.64 |
| $\mathbf{2 0 0 5}$ | 268 | 222.46 | 222 | 8.01 |
| $\mathbf{2 0 0 6}$ | 368 | 224.94 | 226 | 7.45 |
| $\mathbf{2 0 0 7}$ | 288 | 208.70 | 207.5 | 8.22 |
| $\mathbf{2 0 0 8}$ | 564 | 225.98 | 226 | 10.46 |
| $\mathbf{2 0 1 0}$ | 570 | 218.87 | 219 | 7.05 |
| $\mathbf{2 0 1 1}$ | 506 | 208.13 | 209 | 7.09 |
| $\mathbf{2 0 1 3}$ | 595 | 225.41 | 226 | 8.95 |
| $\mathbf{2 0 1 4}$ | 568 | 221.05 | 223 | 9.85 |
| $\mathbf{2 0 1 5}$ | 583 | 216.52 | 217 | 7.93 |
| $\mathbf{2 0 1 6}$ | 524 | 227.07 | 227 | 7.57 |
| $\mathbf{A L L}$ | 5623 | 220.52 | 221 | 10.42 |



| Flowering-Veraison |  |  |  |  |
| :--- | ---: | ---: | ---: | ---: |
| YEAR | Count | Mean | Median | SD |
| $\mathbf{2 0 0 4}$ | 263 | 63 | 63 | 6.64 |
| $\mathbf{2 0 0 5}$ | 268 | 68.9 | 68 | 7.79 |
| $\mathbf{2 0 0 6}$ | 369 | 67.1 | 68 | 7.76 |
| $\mathbf{2 0 0 7}$ | 288 | 70.8 | 70 | 7.64 |
| $\mathbf{2 0 0 8}$ | 567 | 70.8 | 71 | 10.2 |
| $\mathbf{2 0 1 0}$ | 568 | 64 | 64 | 7.16 |
| $\mathbf{2 0 1 1}$ | 500 | 64.7 | 66 | 7.17 |
| $\mathbf{2 0 1 3}$ | 536 | 65.9 | 67 | 8.75 |
| $\mathbf{2 0 1 4}$ | 564 | 74.4 | 76 | 9.95 |
| $\mathbf{2 0 1 5}$ | 583 | 66.2 | 67 | 7.86 |
| $\mathbf{2 0 1 6}$ | 521 | 70.5 | 71 | 7.37 |
| ALL | 5392 | 67.85 | 68.27 | 8.03 |

Figure 14. Panel 1. On previous page, panel showing different plots and tables regarding raw phenotypic values; flowering, veraison and flowering-veraison interval in orange, purple and blue respectively. For images, from top left, clockwise: density plots showing trait distribution over different years; histogram of overall distribution across all years; QQ plot across all years; boxplot showing inter-years differences.



| Veraison GDD |  |  |  |  |
| :--- | ---: | ---: | ---: | ---: |
| YEAR | Count | Mean | Median | SD |
| $\mathbf{2 0 0 4}$ | 264 | 1123 | 1134 | 87.3 |
| $\mathbf{2 0 0 5}$ | 269 | 1239 | 1232 | 86.3 |
| $\mathbf{2 0 0 6}$ | 372 | 1407 | 1415 | 92 |
| $\mathbf{2 0 0 7}$ | 289 | 1400 | 1394 | 114 |
| $\mathbf{2 0 0 8}$ | 567 | 1458 | 1465 | 153 |
| $\mathbf{2 0 1 0}$ | 569 | 1322 | 1320 | 90.9 |
| $\mathbf{2 0 1 1}$ | 507 | 1267 | 1271 | 84.9 |
| $\mathbf{2 0 1 3}$ | 538 | 1392 | 1407 | 130 |
| $\mathbf{2 0 1 4}$ | 565 | 1394 | 1428 | 124 |
| $\mathbf{2 0 1 5}$ | 583 | 1439 | 1439 | 121 |
| $\mathbf{2 0 1 6}$ | 521 | 1386 | 1376 | 88.8 |
| ALL | 5044 | 1347.91 | 1352.82 | 106.6 |

Flowering-Veraison GDD

| YEAR | Count | Mean | Median | SD |
| :--- | ---: | ---: | ---: | ---: |
| $\mathbf{2 0 0 4}$ | 263 | 643 | 641 | 86.3 |
| $\mathbf{2 0 0 5}$ | 268 | 769 | 762 | 83.8 |
| $\mathbf{2 0 0 6}$ | 371 | 937 | 947 | 92.7 |
| $\mathbf{2 0 0 7}$ | 288 | 897 | 893 | 106 |
| $\mathbf{2 0 0 8}$ | 567 | 987 | 993 | 149 |
| $\mathbf{2 0 1 0}$ | 568 | 855 | 851 | 91.7 |
| $\mathbf{2 0 1 1}$ | 500 | 778 | 783 | 86.7 |
| $\mathbf{2 0 1 3}$ | 536 | 945 | 967 | 127 |
| $\mathbf{2 0 1 4}$ | 564 | 893 | 922 | 123 |
| $\mathbf{2 0 1 5}$ | 583 | 934 | 943 | 118 |
| $\mathbf{2 0 1 6}$ | 521 | 894 | 887 | 86.1 |
| $\mathbf{A L L}$ | 5029 | 866.545 | 871.727 | 104.6 |

Figure 14. Panel 2. On previous page, panel showing different plots and tables regarding GDD phenotypic values; flowering, veraison and flowering-veraison interval in orange, purple and blue respectively. For images, from top left, clockwise: density plots showing trait distribution over different years; histogram of overall distribution across all years; QQ plot across all years; boxplot showing inter-years differences.


Figure 15. Pairwise t-test p-values intensity for flowering (upper panel) and veraison (lower panel) traits; in each panel lower triangle is raw values while upper triangle displays values for GDD values. Scales are from red to green, where red values represent more lower p-values while yellow to green values represent higher pvalues.

## Diversity panel construction

Since our aim was to perform a GWAS on the CCC, we wanted to utilize a subset diversity panel (core collection) representing the entire allelic diversity of the CCC. We downloaded GrapeReSeq 18K Vitis genotyping chip data from a recent publication (Laucou et al., 2018) consisting in a matrix of 10,207 SNPs x 783 unique grapevine individuals with no missing values. The number of cultivars in common between the 530 from CCC and the 783 was 95 (Supplementary Table 13). When highlighted in the PCA plot based on SSR data, these 95 cultivars show a sort of good coverage of the entire plot space (Figure 16).


Figure 16. PCA plot based on SSR data of the CCC. In blue, the 95 cultivars in common with the French collection are highlighted. In orange, the rest of the cultivars.

When considering the average number of alleles (allelic diversity) of the 95 cultivars in common between the CCC and the French collection, the value reached is 6.95 , compared to the one of the entire CCC which is 8.2 . To obtain the minimum number of cultivars covering all the loci of the entire collection, we used the approach of the core collection construction, using the $R$ package corehunter (De Beukelaer et al., 2018). This software is able to select the minimum number of representatives from larger collection, with least redundancy, while maximizing
different indexes, like allelic diversity or expected heterozygosity. Given as input the SSR genotypic data matrix from the CCC ( $45 \mathrm{SSR} \times 530$ cultivars), and as main option, to keep the 95 cultivars in common as fixed (that is, always included in the resulting core collection), we evaluated the lowest number of cultivars needed to reach the allelic diversity average value of 8.2 . As shown in Table 13, the number of cultivars needed to reach this value is 132, that is, the 95 fixed ones plus 37 additional cultivars from the CCC.

Table 13. Values of allele coverage (CV, i.e. the percentage of marker alleles observed in the full collection that are retained in the core) and allelic diversity ( $A D$ ) obtained from the software corehunter, starting from the 95 cultivars fixed, adding one cultivar at the time, until reaching the complete coverage of the CCC.

| $\mathbf{N}^{\circ}$ of entries | $\mathbf{C V}$ | $\mathbf{A D}$ |
| :--- | ---: | ---: |
| 95 | 0.8482385 | 6.9555557 |
| 96 | 0.8617886 | 7.06666652 |
| 97 | 0.8753388 | 7.17777816 |
| 98 | 0.8834688 | 7.24444416 |
| 99 | 0.8915989 | 7.31111098 |
| 100 | 0.899729 | 7.3777778 |
| 101 | 0.9051491 | 7.42222262 |
| 102 | 0.9105691 | 7.46666662 |
| 103 | 0.9159892 | 7.51111144 |
| 104 | 0.9214092 | 7.55555544 |
| 105 | 0.9268293 | 7.60000026 |
| 106 | 0.9295393 | 7.62222226 |
| 107 | 0.9322493 | 7.64444426 |
| 108 | 0.9349593 | 7.66666626 |
| 109 | 0.9376694 | 7.68888908 |
| 110 | 0.9403794 | 7.71111108 |
| 111 | 0.9430894 | 7.73333308 |
| 112 | 0.9457995 | 7.7555559 |
| 113 | 0.9485095 | 7.7777779 |
| 114 | 0.9512195 | 7.7999999 |
| 115 | 0.9539295 | 7.8222219 |
| 116 | 0.9566396 | 7.84444472 |
| 117 | 0.9593496 | 7.86666672 |
| 118 | 0.9620596 | 7.88888872 |
| 119 | 0.9647696 | 7.91111072 |
| 120 | 0.9674797 | 7.93333354 |
| 121 | 0.9701897 | 7.95555554 |
| 122 | 0.9728997 | 7.97777754 |
| 123 | 0.9756098 | 8.00000036 |
| 124 | 0.9783198 | 8.02222236 |
| 125 | 0.9810298 | 8.04444436 |
| 126 | 0.9837398 | 8.06666636 |
| 127 | 0.9864499 | 8.08888918 |
| 128 | 0.9891599 | 8.11111118 |
| 129 | 0.9918699 | 8.13333318 |
| 130 | 0.9945799 | 8.15555518 |
| 131 | 0.99729 | 8.177778 |
| 132 | 1 | 8.2 |
|  |  |  |

Since, among the 435 remaining cultivars ( $530-95=435$ ), the number of possible combinations to choose the 37 additional one was high, we simulated 100 core collections of 132 cultivars, and selected, among the 37 , the cultivars that were more present in the 100 simulations. We found 26 cultivars being always present in all the simulations; to choose the remaining 11 , we counted the frequency of appearance of the remaining 11 cultivars in all the 100 core collections and selected the most frequent ones. Figure 17 shows the PCA plot where the resulting diversity panel is highlighted. The 37 additional cultivars that resulted necessary to reach the allelic diversity maximization are depicted in red in Figure 17. From their position in the PCA plot, it can be noted that they get to cover quite uniformly the remaining areas that were not fully covered with just the individuals from the French collection (in blue).


Figure 17. PCA plot based on SSR data of the CCC. In blue, the 95 cultivars in common with the French collection are highlighted. In red, the 37 cultivars that together with the 95, maximize the allelic diversity of the CCC. In orange, the rest of the cultivars.

Summary statistics of the genetic diversity of the core collection compared to the CCC are given in Table 14. The average number of observed alleles is, as requested before, the same of the CCC, while both Shannon diversity Index and expected
heterozygosity are higher than in the CCC. The index of evenness, a measure of the distribution of genotype abundances, is highly comparable between the two.

Table 14. Summary genetic diversity statistics of the core collection and the CCC.

|  | $\mathbf{N a}$ | $\mathbf{I}$ | He | Evenness |
| :--- | ---: | ---: | ---: | ---: |
| CORE | 8.2 | 1.4570 | 0.6842 | 0.7526 |
| CCC | 8.2 | 1.4264 | 0.6759 | 0.7572 |

When the composition of the diversity panel from the geographical origin point of view is compared with the entire germplasm collection, the unique geographic group that shows an altered composition is the one of the cultivars from western Europe, which is enriched in the core collection compared to the entire collection (Table 15 A ). When the same comparisons are made for grapes usage, no enrichment is observed (Table 15 B ), and the same can be said regarding the numerosity of the cultivars of the core belonging to the subgroups identified by STRUCTURE, both when considering the 3 and the 6 groups subdivision (Table 15 C and D ), suggesting a homogenous composition of the diversity panel from different perspectives. Supplementary table 14 describes the 132 cultivars included in the diversity panel.

Table 15. A. Contingency table of geographic composition of the core collection of 132 cultivars identified with corehunter. B. Contingency table of grapes usage of the core collection. C. Contingency table of the composition of the 2 subgroups subdivision identified by structure compared between the core collection and the CCC. D. Contingency table of the composition of the 6 subgroups subdivision identified by structure compared between the core collection and the CCC. In red overrepresented groups. In parenthesis, in italics are reported p-values from Fisher's Exact Test two-tailed.

A | COUNTRY | CORE | CCC |
| :--- | :--- | ---: | :--- |
| AM | $6(0.1296)$ | 11 |

| CI | $31(0.827)$ | 134 |
| :--- | ---: | ---: |
| CNE | $4(0.7583)$ | 13 |

EE $\quad 9(0.1397) \quad 18$

| IS | $5(1)$ | 22 |
| :--- | ---: | ---: |
| IT | $14(0.2171)$ | 83 |

NI $\quad 24(0.2142) \quad 131$

| SI | $12(0.8714)$ | 54 |
| :--- | ---: | ---: |
| UKN | $1(1)$ | 7 |


| WE | $26(0.022 I)$ |
| ---: | ---: |
| 132 | 530 |

B

| USAGE | CORE | CCC |
| :--- | ---: | ---: |
| TABLE | $18(0.2755)$ | 52 |
| TABLE/RAISIN | $1(0.4880)$ | 2 |
| UKN | 0 | 3 |
| WINE | $89(0.4063)$ | 409 |
| WINE/TABLE | $23(0.1176)$ | 61 |
| WINE/TABLE/RAISIN | $1(1)$ | 3 |
|  | 132 | 530 |


| STR-6 | CORE | ALL |
| :--- | ---: | ---: |
| a | $18(0.0973)$ | 43 |
| b | $5(0.3576)$ | 12 |
| c | $2(1)$ | 10 |
| d | $20(0.2298)$ | 56 |
| e | $3(0.2404)$ | 26 |
| f | $9(0.0923)$ | 67 |
| g (admixed) | $75(0.8099)$ | 316 |
|  | 132 | 530 |

## Phenotypic diversity of the $\mathbf{1 3 2}$ cultivars of the GWAS diversity panel

We studied the phenotypic data of the 132 individuals included in the GWAS diversity panel, and compared them to the data of the entire collection, to understand if the approach of maximizing the number of alleles with the minimum number of individuals would retain a phenotypic distribution similar to the CCC one. Figure 16 shows the phenotypic distribution of the cultivars included in the
panel overlaid with the distribution of the entire collection. As can be seen from the density plots, the phenotypic values of the diversity panel precisely reflect the values of the entire collection, except for very few values in the tails.


Figure 16. Density plots of the phenotypic traits flowering beginning (FB), veraison beginning (VB) and flowering-veraison interval $\left(F_{-} V\right)$ from the diversity panel, transparent colour, and from the entire collection, normal colour. On the x-axis the phenotypic value expressed in days of year. On the y-axis the density distribution.

## Integration of published SNPchip data: DNA extraction, hybridization and SNP data extraction

DNA of the 37 additional samples identified to reach the 132 individuals diversity panel was extracted with the Qiagen DNeasy Plant Mini Kit (Qiagen, Hilden, Germany) and quantified with the NanoDrop. Supplementary Table 13 reports the concentration and quality ratio values. Approximately $1 \mu \mathrm{~L}$ for each sample at a concentration ranging from $40-60 \mathrm{ng} / \mu \mathrm{L}$ was used to perform the GrapeReSeq 18K Vitis genotyping chip hybridization. Hybridizations evaluation and SNP calling were performed using the standard workflow of GenomeStudio Data Analysis v2011.1 (Illumina Inc, San Diego, CA, USA). After exporting the genotypic matrix for the 37 samples for performing the subsequent GWAS analysis we subset it to the 10,207 solid SNPs identified by Laucou et al. Among these

10,207 SNPs, we found only 97 SNPs that did not meet quality criteria, so that the resulting matrix used for GWAS was 10,110 SNPs x 37 individuals. We then merged the data matrix obtained from Laucou et al. with the data matrix produced by us and checked the consistency of both data sets in terms of genotypes scoring.

## Association analysis SNPchip

Before running the association tests, we filtered SNPs having a minor allele frequency less than 0.05 using TASSEL, obtaining 9536 SNPs in total. In order to have a sort of confirmation of the design of our diversity panel, we performed a preliminary association test using the skin colour as phenotype. Using EMMAX (Kang et al., 2010) and running an association test correcting for kinship, inputting a kinship matrix as calculated by EMMAX itself on the same data matrix, we found as the most significant region associated to the colour phenotype, a region on chromosome 2 very close and overlapping to the extensively characterized colour locus (Fournier-Level et al., 2009) (Figure 17).

Skin color


Figure 17. Manhattan plot showing the results of the GWAS association test using grapes skin colour as phenotype. Most significantly associated SNPs reside in the region of chromosome 2 around 15 Mbp . In this case GWAS was performed with EMMAX.

The most significantly associated SNP, Vv_12X_V2_Chr2_15524738, is also the same most significantly associated SNP found when the association test is run using QTCAT, in this way giving a significant comparison between the approaches. A
similar test with grape skin colour was performed also by Laucou et al. and they identified the same region as the most significantly associated one.

We then proceeded with performing GWAS association on the phenology phenotypes with three different software, of which, two, use population structure or kinship as factor of correction, EMMAX and Gapit, while QTCAT works without using any other file for correction. Different runs were performed leading to many combinations (Supplementary Table 17), whose most significant results for the phenotypic values averaged across all the years are reported in Table 16. In this case the most significant results come from the raw phenotypic values and not the GDD ones (results not shown). Regarding EMMAX and Gapit, to make the results more comparable, the model used for both was the one corrected only by kinship. No significant results were found for the trait flowering beginning, while different significant SNPs were found for the other 2 traits (Table 16). Regarding veraison beginning, 5 significantly associated SNPs were found but only by the QTCAT approach, that were located respectively on chromosomes 3, 12, 13, 16 and 18. Concerning flowering-veraison interval, the three approaches identified altogether 13 signals of association, located on chromosomes 4, 8, 11, 12, 14, 16 and 18. Importantly 3 SNPs were significantly associated both to the veraison beginning trait and to flowering-veraison interval, thus suggesting a relevance of these regions in controlling grape veraison time and as consequence the floweringveraison interval length. As far as the use of different software is concerned, QTCAT identified the highest number of significantly associated SNPs (5 for veraison beginning and 10 for flowering-veraison interval) while. Gapit identified three SNPs and EMMAX only one. Interestingly two of the three SNPs associated to both traits also emerged by applying different softwares.

Table 16. The table reports the most significantly associated SNPs for the phenological traits averaged across the years. No significant results were found for the beginning of flowering. * $F D R<0.05$; ${ }^{* *} F D R<0.01$ as obtained from the approach under which the association was found. For QTCAT, the concept of FDR does not stand, so the stars are given according to the p-value.

|  | Trait | Software | Significance | Multiple evidence |
| ---: | ---: | ---: | ---: | ---: |
| $\boldsymbol{C h r} 3 \_379409$ | VB | QTCAT | $*$ | No |


| Chr4_6106210 | F-V | QTCAT | ** | No |
| :---: | :---: | :---: | :---: | :---: |
| Chr8_3847595 | F-V | Gapit | * | No |
| Chr8_7340158 | F-V | QTCAT | * | No |
| Chr8_8216855 | F-V | Gapit | ** | No |
| Chr11_963022 | F-V | QTCAT | * | No |
| Chr11_1343142 | F-V | QTCAT | * | No |
| Chr12_21100465 | VB, F-V | QTCAT, EMMAX | **, $p<0.1$ | Yes |
| Chr13_26217739 | VB | QTCAT | ** | No |
| Chr14_23248717 | F-V | QTCAT | ** | No |
| Chr14_29502138 | F-V | QTCAT | ** | No |
| Chr16_348557 | VB, F-V, F-V | QTCAT, Gapit, QTCAT | **, *, ** | Yes |
| Chr16_20303373 | F-V | QTCAT | ** | No |
| Chr18_3381571 | VB, F-V | QTCAT, QTCAT | **, ** | Yes |
| Chr18_22124028 | F-V | QTCAT | ** | No |

For further discussion we decided to focus on the SNPs associated to both traits veraison time and flowering veraison interval. The SNP $V v_{-} 12 X_{-} V 2 \_C h r 12 \_21100465$ that resulted significant $(p=0.00162)$ with the software QTCAT for the veraison beginning trait, and almost significant for the flowering-veraison interval trait with the software EMMAX (FDR $=0.052$ ). This SNP resides in the coding sequence of the gene VIT_12s0035g00280, that is annotated as an endomembrane protein 70 . The second SNP that resulted in common fir both traits and was revealed by between more than one approach was Vv_12X_V2_Chr16_348557. This SNP resulted significant with the software Gapit for the trait F-V $(\mathrm{FDR}=0.044)$ and with QTCAT both for VB $(p=0.00226)$ and F-V ( $p=0.00356$ ). This SNP resides in the coding sequence of the gene VIT_16s0039g00700, that is annotated as a maltose transporter (RCP1 in $A$. thaliana), thus a gene involved in the carbohydrate metabolic processes. The third and last SNP pointed out as in common for both traits is $V v_{-} 12 X_{-} V 2 \_C h r 18 \_3381571$. This SNP resulted significant only with the software QTCAT but with the two phenotypic traits VB $(p=0.00318)$ and F-V $(p=0.00294)$. Also this SNP resides in the coding sequence of a gene, VIT_18s0001g03540, whose function is described as an auxin transporter protein 4 (LAX2 in A. thaliana). Moreover, this same gene is included in a meta-QTL interval that we identified in chapter one, in particular within pheno_18_1, a minor meta-QTL originating from
two QTLs, one from a mapped from flowering time trait (Carreño Ruiz, 2012) and the other from veraison time trait (Zyprian et al., 2016).

Table 17. Table listing the upstream and downstream flanking genes to the most significant SNPs associated to VB and F-V traits. In the first column, highlighted in italics and bold are the genes where the significantly associated SNP falls. Only highlighted in italics are the genes considered interesting for explaining the phenotype according to their function.

| Flanking genes | Chr | Start | End | Function | Gene Ontology Slim |
| :---: | :---: | :---: | :---: | :---: | :---: |
| VIT_12s0035g00200 | 12 | 20991910 | 21013685 | Phospholipase D | lipid metabolic process; catabolic process; cellular process |
| VIT_12s0035g00210 | 12 | 21019518 | 21020261 | MAPKKK19 | cellular protein modification process |
| VIT_12s0035g00220 | 12 | 21046133 | 21050594 | No hit |  |
| VIT_12s0035g00240 | 12 | 21059265 | 21062347 | DAG protein, chloroplast precursor |  |
| VIT_12s0035g00250 | 12 | 21064129 | 21066252 | EMB2758 |  |
| VIT_12s0035g00260 | 12 | 21070623 | 21081261 | Mlo4 | response to stress; cell death; response to biotic stimulus |
| VIT_12s0035g00270 | 12 | 21081724 | 21082383 | Ferredoxin, chloroplast (PETF) | generation of precursor metabolites and energy; transport; cellular process; protein metabolic process |
| VIT_12s0035g00280 | 12 | 21083032 | 21100612 | Endomembrane protein 70 |  |
| VIT_12s0035g00290 | 12 | 21118063 | 21120870 | Sugar transporter ERD6-like 6 |  |
| VIT_12s0035g00300 | 12 | 21130316 | 21134875 | Carbohydrate transmembrane transporter SFP1 | transport; cellular process |
| VIT_12s0035g00310 | 12 | 21148330 | 21154433 | Protein kinase SPK-3 ASK1 (SnRK-6) | cellular protein modification process; response to stress; signal transduction; response to abiotic stimulus; response to endogenous stimulus |
| VIT_12s0035g00320 | 12 | 21159933 | 21169655 | Phosphomethylpyrimidine kinase; thiamin-phosphate pyrophosphorylase | biosynthetic process; cellular process |
|  |  |  |  |  |  |
| VIT_16s0039g00570 | 16 | 283917 | 286597 | 10-deacetylbaccatin III 10-O-acetyltransferase | cellular protein modification process |
| VIT_16s0039g00610 | 16 | 316837 | 332541 | Serine/threonine-protein kinase ppk15 |  |
| VIT_16s0039g00660 | 16 | 332542 | 337573 | RNA polymerase nonessential primary-like sigma factor SIGA |  |


| VIT_16s0039g00690 | 16 | 341722 | 344560 | D111/G-patch domain-containing protein | carbohydrate metabolic process; <br> transport; catabolic process; response to biotic stimulus; cellular process |
| :---: | :---: | :---: | :---: | :---: | :---: |
| VIT_16s0039g00700 | 16 | 346728 | 353020 | RCP1 (root cap 1) |  |
| VIT_16s0039g00720 | 16 | 357505 | 361097 | Folate-biopterin transporter | DNA metabolic process; biosynthetic process |
| VIT_16s0039g00730 | 16 | 361349 | 363172 | DNA polymerase delta, subunit D | cellular process; cellular component organization |
| VIT_16s0039g00740 | 16 | 370309 | 400930 | Histone-lysine N-methyltransferase ASHH3 | metabolic process |
| VIT_16s0039g00760 | 16 | 401533 | 403436 | CYP89A2 |  |
|  |  |  |  |  |  |
| VIT_18s0001g03390 | 18 | 3298924 | 3306685 | S-receptor kinase | carbohydrate metabolic process; cellular protein modification process; biosynthetic process; pollen-pistil interaction; cellular process |
| VIT_18s0001g03420 | 18 | 3307440 | 3307550 | No hit |  |
| VIT_18s0001g03430 | 18 | 3309917 | 3311940 | Flavonol synthase | > metabolic process; biosynthetic process; cellular process; secondary metabolic process |
| VIT_18s0001g03440 | 18 | 3313991 | 3315438 | No hit |  |
| VIT_18s0001g03450 | 18 | 3320975 | 3328281 | Glycine-rich protein |  |
| VIT_18s0001g03470 | 18 | 3338337 | 3340959 | Flavonol synthase | metabolic process; biosynthetic process; cellular process; secondary metabolic process |


| VIT_18s0001g03490 | 18 | 3349225 | 3350592 | Flavonol synthase | metabolic process; biosynthetic process; cellular process; secondary metabolic process |
| :---: | :---: | :---: | :---: | :---: | :---: |
| VIT_18s0001g03510 | 18 | 3354104 | 3359003 | Flavonol synthase XM_002284374.1 | > metabolic process; biosynthetic process; cellular process; secondary metabolic process |
| VIT_18s0001g03520 | 18 | 3359393 | 3369207 | Camphor resistance CrcB |  |
| VIT_18s0001g03540 | 18 | 3380455 | 3383288 | Auxin transporter protein 4 | transport; signal transduction; response to endogenous stimulus |
| VIT_18s0001g03570 | 18 | 3387900 | 3389335 | Thaumatin ATLP-1 | response to stress |
| VIT_18s0001g03580 | 18 | 3389546 | 3393993 | Ubiquitin-fold modifier 1 precursor |  |
| VIT_18s0001g03610 | 18 | 3401893 | 3411144 | Auxin-independent growth promoter |  |
| VIT_18s0001g03630 | 18 | 3412345 | 3414417 | Pentatricopeptide (PPR) repeat-containing protein | biological_process; catabolic process; cellular process |
| VIT_18s0001g03640 | 18 | 3417193 | 3418012 | No hit |  |
| VIT_18s0001g03650 | 18 | 3418650 | 3419295 | No hit |  |
| VIT_18s0001g03670 | 18 | 3422279 | 3424214 | Zinc finger (C2H2 type) family |  |
| VIT_18s0001g03680 | 18 | 3433101 | 3445210 | Protein kinase | cellular protein modification process |
| VIT_18s0001g03720 | 18 | 3445760 | 3445939 | No hit |  |
| VIT_18s0001g03730 | 18 | 3445960 | 3490992 | SET Domain group 37 |  |

Since in this diversity panel we estimated a linkage disequilibrium decay at 0.2 of around 77 Kbp (averaged across all the chromosomes and corrected by kinship with the $R$ package $L D \operatorname{corSV}$ ) showed in Supplementary Table 16, it is clear that we reported only the gene closest to the significant SNP but other genes may be linked to the studied trait. Looking at the Manhattan plot and the QQ plot of expected versus observed $p$-values for the F-V GWAS performed with EMMAX, it is clear to observe that, on the contrary with the grapes skin colour phenotype, it is not possible to identify a typical dense column of SNPs with low $p$-values around the most significant one. Indeed, the most significant SNP looks like a false positive, maybe due to population structure stratification, but if we consider that for chromosome 12, LD decay at 0.2 is around 50 Kbp and after filtering the closest downstream and upstream SNPs are 20 Kbp and 48 Kbp away respectively, that does not look so much like a strange signal of association. Considering the other two significantly associated SNPs that we selected, the same idea can be applied, and we see that for SNP Vv_12X_V2_Chr16_348557, the closest SNPs are at 25 Kbp (down) and 22 Kbp (up), where LD decay for chromosome 16 is around 37 Kbp. For the SNP identified with QTCAT, this reasoning is more difficult to discuss since the method does not imply any kind of correction and every SNP is tested individually within blocks of similar SNPs.

A complete list of the genes flanking the three most significant selected SNPs, in the range of LD decay for that chromosome, is given in Table 17. Few interesting candidate genes are highlighted and here discussed. On chromosome 12 two downstream genes of $V v_{-} 12 X_{-} V 2_{-}$Chr12_21100465 SNP, that is VIT_12s0035g00290 and VIT_12s0035g00300, appeared as functionally related to the phenotype. The first gene is annotated as a sugar transporter ERD6-like 6, a subgroup of the monosaccharide transporters family firstly described in Arabidopsis (Büttner, 2007). ERD stands for early-responsive to dehydration, and the gene is in fact induced upon dehydration (Kiyosue et al., 1998). Interestingly, dehydration is a characteristic step of the berry ripening process after veraison (Keller, 2010), and different transcriptomic studies have previously identified several sugar accumulation related genes as induced after veraison in the grapevine berry (Fasoli et al., 2012, 2018; Fontana et al., 2007; Massonnet et al., 2017;

Palumbo et al., 2014). The other gene is annotated as a carbohydrate transmembrane transporter SFP1, a sugar-porter family protein closely related to ERD6-like genes. SFP1 has a homolog, SFP2, and they are found as tandem genes. SFP1 was shown to be senescence-induced, which is paralleled by an accumulation of monosaccharides in the Arabidopsis leaves (Quirino et al., 2001). Regarding SNP Vv_12X_V2_Chr16_348557 potential candidates, we highlight the gene VIT_16s0039g00570, annotated as a 10-deacetylbaccatin III 10-Oacetyltransferase, an enzyme belonging to the family of acyltransferases that catalyses the conversion of acetyl-CoA and 10-deacetylbaccatin III to CoA and baccatin III (Walker and Croteau, 2000). This enzyme participates in diterpenoid biosynthesis. Terpenoids are well known as major components for wine flavour and aroma (Lund and Bohlmann, 2006; Martin et al., 2010; Wen et al., 2015), and have been found to accumulate during the last phase of berry development, when most of the secondary metabolites accumulate (Lücker et al., 2004). Different transcriptomic studies have identified diverse genes encoding for enzymes responsible for the biosynthesis of secondary metabolites to increase their expression right before the last of the three steps of berry development (Deluc et al., 2007), but few if none have focused on this specific candidate, making it interesting for further explorations. The last one of the three candidate SNPs, Vv_12X_V2_Chr18_3381571, is flanked upstream by a family of flavonol synthase encoding genes. Flavonol synthase, from the family of oxidoreductases, is an enzyme that catalyses the production of flavonol. Flavonols are a predominant class of flavonoids, secondary metabolites widely present in plants, that are involved in different functions like response to biotic and abiotic stresses. In grapevine berries they accumulate in the cell wall and the vacuole of the skin cells and are almost absent in the pulp. Like anthocyanins, the determinants of the red colour in red grapevine cultivars, flavonols are a product of the phenylpropanoid pathway. Regarding the content of flavonols in the berry skin during development, it has been found to start accumulation right after veraison, and then reach a maximum at the initial stage of berry development (Downey et al., 2003). Other genes encoding for flavonol synthase or for different enzymes involved in the phenylpropanoid pathway have been also recently emerged as candidates from few transcriptomics
experiments (Fasoli et al., 2012; Palumbo et al., 2014); even more recently a gene encoding for a 2'-hydroxy isoflavone/dihydroflavonol reductase (from the anthocyanin biosynthesis) has been defined, amongst others, as a marker transition gene, representing a gene that marks the transition to the late ripening stages of the berry, in agreement to the observation on the accumulation of anthocyanins (Downey et al., 2003).

When we looked at the phenotypic differences for the three allelic combinations of each one of the three most interesting SNPs, selected for resulting as significantly associated across more than one trait or more than one software, in almost all cases we observe quite a significant difference between the heterozygotes and the homozygotes cultivars for those SNPs, as shown in Figure 17.


Figure 17. Boxplots showing phenotypic values of veraison beginning and flowering-veraison interval phenotypes against genotypes of the three selected SNPs significantly associated to the traits. The three SNPs Vv_12X_V2_Chr12_21100465, Vv_12X_V2_Chr16_348557 and Vv_12X_V2_Chr18_3381571 are in $A, B$ and $C$ respectively.

In particular for SNP $V v_{-} 12 X_{-} V 2_{-} C h r 16_{-} 348557$ where the cultivars seem more equally distributed in the three genotypic classes, the differences are quite significant and an average of around ten days diversity in the veraison date and flowering-veraison time interval is observed between the two homozygotes (Figure 17 B).

Overall, the results suggest that probably a bigger sample size is needed to catch all the rare alleles. Indeed phenology in general, and in particular veraison time, is considered a polygenic complex trait, with many loci contributing to small variations, and many authors have previously failed to identify strong signals of association for such complex traits (Marrano et al., 2018; Migicovsky et al., 2017; Myles et al., 2011). Moreover, the grapevine SNPchip, with its 10K SNPs, may not be an adequate technology to identify all the small variance-contributors loci for such complex traits. In particular because linkage disequilibrium extends longer for regions subject to selection but, on the contrary to berry size or grapes skin colour, which are typical traits selected over many generations, the time of veraison and phenology in general are traits that have not been selected since long times.

To overcome such limitations, we opted for a different experimental design that takes into account the phenotypic diversity of a germplasm collection and measures allele frequencies in the extreme pools, enabling discovery of associations between genetic variants and traits of interest, without testing the association of the single individual to the phenotype. Indeed, the XP-GWAS approach overcome the limitations of a SNPchip GWAS by allowing a bigger sample size at a reduced cost, enriching for rare alleles and augment allele effects via extreme phenotypic selection, and using a whole-genome resequencing approach, opposed to genotyping chip where density of coverage of the genome is way lower than in a resequencing. Moreover, XP-GWAS, by considering the average phenotypic value of a pool of individuals, tolerates a degree of inaccuracy in the phenotyping data, which is desirable for traits difficult to phenotype like flowering and veraison time
or phenology in general, where the trait is not easily quantifiable by human eyes. We created three pools of individuals, named low, high and random pool by observing the distribution of the flowering-veraison interval trait and after DNA pooling, we are currently subjecting them to resequencing.

## Conclusions

Grapevine veraison is a stage that is critical for determining the final quality of the grape, and ultimately the wine. The transition from the immature to the mature stage of the grapevine berry has been demonstrated to be a critical point where environmental conditions can impact significantly the entire cycle of formation of the berry and its components. In particular, in a scenario where climate change will impact the temperature with a forecasted increase of around 2 degrees in the next 50 years in many areas including the most famous viticulture area (Jones et al., 2005), this situation may disclose different aspects to be taken into account when considering viticulture and the winemaking process. For example, it has been demonstrated that higher temperatures impact the formation of typical compounds of the grape berry skin, like flavonoids and anthocyanins in particular (Mori et al., 2007; Pastore et al., 2017; Yamane et al., 2006) and this will affect the assembly of other compounds that represent the specificity of a wine, like tannins. The increased temperature of the last years led to and will lead to an advance of the phenological stages (Tomasi et al., 2011), and this advance could bring to a more rapid ripening phase, that when different to a normal maturation is not paralleled by the accumulation of flavour components that require more time to store. In particular, if veraison will occur earlier during the season there will be less time for the preveraison production of flavanols that can be assembled into tannins before and during ripening. A possible overcome to this problem is to wait the flavours to assemble and develop but doing so, grapes are harvested with very high sugar concentrations that will result in wines with high alcohol contents. Temperature is not the only factor that can influence phenological stages in grapevine, but also conditions of water deficit, both before and after veraison, can impact the formation of organic compounds in the berry and in particular in the skins (Ojeda et al., 2002). With such a scenario, different approaches may be taken in order to contrast the effect of climate change on grapevine phenology. The genetic approach, that is for example exploiting genetic diversity from the wild or from germplasm collections to integrate rare and natural alleles into existing cultivars, is a long process and it still needs to be addressed carefully in grapevine. The genetics underlying complex traits like phenology is still poorly known. Here, by integrating different genetic
approaches to identify the genetic determinants of grapevine veraison process, we aimed to produce relevant information towards the disclosure of the major loci contributing to the berry ripening process.

In the first chapter we created the first integrated consensus map of grapevine from different individual grapevine genetic maps with the aim to perform the first metaQTL analysis for grapevine. In detail, we collected genetic maps info from 42 different publications and integrated them with a statistical approach, in order to obtain the most significant and reliable grapevine consensus map that can be used to summarize and integrate different QTL results. We collected all the available QTL information that were produced for grapevine and then we focused on phenological traits. We identified 4 meta-QTLs for the veraison trait that represent the most reliable genomic spots where the most significant QTLs mapped for veraison so far have been identified. One of the main aims was to reduce the number of possible positional candidate genes comprised in the high number of veraison associated QTL present in literature. In performing such meta-analysis, we were able to reduce the number of candidate genes by almost 4 -fold. Indeed, one of the main limitations of the QTL approach is the downstream analysis represented by the search of the candidate genes; since the number of genes within the interval of a QTL can be really high, an approach that tries to reduce this number may be eventually useful, especially in situations where the trait studied is a complex trait, and the loci governing the trait are numerous. After performing the meta-analysis of the QTLs, we integrated data from different transcriptomic experiments, given the idea that the genes resulting differentially expressed across veraison time in grapevine berries may be also in common with positional candidate genes deriving from QTL studies. In particular, different previous transcriptomic experiments (Fasoli et al., 2012; Massonnet et al., 2017; Palumbo et al., 2014) have identified a number of candidate genes that are either up or down regulated across the veraison stage of grapevine berries and are considered as master regulators of the transition from the green to the mature stage of the grapevine. For example, some special sets of the genes identified, for example the "switch genes" (Palumbo et al., 2014) characterized with a network analysis as having many significant negative
correlations outside their own group in the network, represent a group of genes, mainly transcription factors, considered as master regulators of the transcriptome remodelling marking the developmental shift from immature to mature growth. We found some of these special master regulators to lie within the meta-QTL intervals, indicating that the integration of different approaches, from summarizing QTL data to integrating existent transcriptomic data, may be helpful to prioritize most significant results, with the final aim of identifying a restricted number of candidate genes to be explored individually.

In the second chapter we studied the genetic diversity of a grapevine germplasm collection, in order to identify the most genetically diverse and representative panel of individuals to be used for genetic association study. Studying the population structure of the collection, we firstly found out that the Conegliano collection is mostly composed by cultivars from Italian origin, and even so, the overall genetic diversity is comparable to the one of other germplasm collection, indicating that among all Vitis vinifera cultivars, the overall genetic diversity is easily reached with a small number of alleles. We were able to identify significant genetically distinct subgroups/substructure in the collection, and found that this subgroups resemble the geographical subdivision of the collection. We collected all these information and used them to construct diversity panels to be used for association analysis. Before, we made use of existing genotypic data, with the idea of integrating them with newly generated data. Through the use of the GrapeReseq Vitis Genotyping Chip we genotyped additional cultivars that from the genetic point of view summarize the entire collection, and tested the genotype-phenotype association with an incredible resource of grapevine phenology phenotypic data provided by our CREA-VIT Conegliano partners. Though some limitations due to sample size, genetic architecture of the trait, and not easily scorable phenotypic traits, we identified some significant genomic regions that would represent the most interesting regions linked to veraison time traits. For doing so, we did not only used the classical GWAS approach, but we had the idea of integrating different approaches represented by different software that apply different statistical procedures. Moreover, we integrated the results from the GWAS approach with the
results of the meta-analysis of the QTLs and found that one of the veraison time most significantly associated SNP is located precisely in a meta-QTL interval on chromosome 18.

We have shown that the integration of existing data, also of different sources, is a strategic procedure when the nature of the trait studied is complex and multifaceted. Also with limited resources, the summarization of existing results and the exploration of new results with the aid of reorganized and prioritized information, may disclose novel targets and shed light on the genetic control of complex traits.

## Acronyms

## CCC CREA-VIT Conegliano Collection

DAPC Discriminant Analysis of Principal Components

FB Flowering beginning

F-V Flowering-Veraison interval

GCM Grapevine Consensus Map

GWAS Genome Wide Association Study

LD Linkage Disequilibrium

PCA Principal Component Analysis

QTL Quantitative Trait Loci

SNP Single Nucleotide Polymorphism

SSR Simple Sequence Repeat

VB Veraison beginning

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## Appendix

Supplementary Table 1
External file (Available upon request
$\begin{array}{lllllllllllllllllll}1 & 2 & 3 & 4 & 5 & 6 & 7 & 8 & 9 & 10 & 11 & 12 & 13 & 14 & 15 & 16 & 17 & 18 & 19\end{array}$


Supplementary Figure 1. The grapevine consensus map

Supplementary Table 2. Contribution of each individual genetic map to the construction of the consensus map.

| Reference | I | II | III | IV | V | VI | VII | VIII | IX | X | XI | XII | XIII | XIV | XV | XVI | XVII | XVIII | XIX |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| Ban et al. 2014 | x | x | x | x | x | x | x | x | x | x | x | x | x | x | x | x | x | x | x |
| Bayo Canha, PhD thesis 2015 | x | x | x | x | x | x | x | x | x | X | x | x | x | x | x | x | x | x | x |
| Bellin et al. 2009 BI | x | x | x | x | x | x | x | x | x | x | x | x | x | x | x | x | x | x | x |
| Bellin et al. 2009 CH | x | x | x | x | x | x | x | x | x | x | x | x | x | x | x | x | x | x | x |
| Bert et al. 2013 | x | x | x | x | x | x | x | x | x | x | x | x | x | x | x | x | x | x | x |
| Blasi et al. 2011 | x | x | x | x | x | x | x | x | x | x | x | x | x | x | x | x | x | x | x |
| Cabezas et al. 2006 | x | x | x | x | x | x | x | x | x | x | x | x | x | x | x | x | x | x | x |
| Carreño Ruiz, PhD thesis 2012 | x | x | x | x | x | x | x | x | x | x | x | x | x | x | x | x | x | x | x |
| Correa et al. 2014 | x | x | x | x | x | x | x | x | x | x | x | x | x | x | x | x | x | x | x |
| Correa et al. 2015 | x | x | x | x | x | x | x | x | x | x | x | x | x | x | x | x | x | x | x |
| Correa et al. 2016 | x | x | x | x | x | x | x | x | x | x | x | x | x | x | x | x | x | x | x |
| Costantini et al. 2008 IT | x | x | x | x | x | x | x | x | x | x |  | x | x | x | x | x | x | x | x |
| Costantini et al. 2008 | x | x | x | x | x | x | x | x | x | x | x | x | x | x | x | x | x | x | x |
| Costantini et al. 2015 | x | x |  | x | x | x | x |  |  | x |  | x |  |  |  |  | x | x |  |
| Coupel-Ledru et al. 2014, 2016 | x | x | x | x | x | x | x | x | x | x |  | x | x | x | x | x | x | x | x |
| Doligez et al. 2006 | x | x | x | x | x | x | x | x | x | x | x | x | x | x | x | x | x | x | x |
| Doligez et al. 2010 | x | x | x | x | x | x | x | x | x | x | x | x | x | x | x | x | x | x | x |
| Duchêne et al. 2009 | x | x | x | x | x | x | x | x | x | x | x | x | x | x | x | x | x | x | x |
| Duchêne et al. 2012 | x | x | x | x | x | x | x | x | x | x | x | x | x | x | x | x | x | x | x |
| Fechter et al. 2014 | x | x | x | x | x | x | x | x | x | x | x | x | x | x | x | x | x | x | x |
| Garris et al. 2009 | x | x | x | x | x | x | x | x | x | x | x | x | x | x | x | x | x | x | x |
| Guo et al. 2015 | x | x | x | x | x | x | x | x | x | x |  | x | x | x | x | x | x | x | x |


| Huang et al. 2012 | x | x | x | x | x | x | x | x | x | x |  | x | x | x | x | x | x | x | x |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| Marguerit et al. 2009, 2011 CS | x | x | x | x | x | x | x | x | x | x |  | x | x | x | x | x | x | x | x |
| Marguerit et al. 2009, 2011 RGM | x | x | x | x | x | x | x | x | x | x |  | x | x | x | x | x | x | x | x |
| Mejía et al. 2007 | x | x | x | x | x | x | x | x | x | x |  | x | x | x | x | x | x | x | x |
| Mejía et al. 2011 | x | x | x | x | x | x | x | x | x | x |  | x | x | x | x | x | x | x | x |
| Moreira et al. 2011 RT | x | x | x | x | x | x | x | x | x | x | x | x | x | x | x | x | x | x | x |
| Moreira et al. 2011 AM | x | x | x | x | x | x | x | x | x | x | x | x | x | x | x | x | x | x | x |
| Ochssner et al. 2016 | x | x | x | x | x | x | x | x | x | x | x | x | x | x | x | x | x | x | x |
| Schwander et al. 2012 | x | x | x | x | x | x | x | x | x | x | x | x | x | x | x | x | x | x | x |
| Riaz et al. 2006 | x | x | x | x | x | x | x | x | x | x | x | x | x | x | x | x | x | x | x |
| van Heerden et al. 2014 | x | x | x | x | x | x | x | x | x | x | x | x | x | x | x | x | x | x | x |
| Venuti et al. 2013 | x | x | x | x | x | x | x | x | x | x | x | x | x | x | x | x | x | x | x |
| Viana et al. 2013 | x | x |  | x | x |  |  |  |  | x |  | x |  |  |  |  | x | x |  |
| Welter et al. 2007 | x | x | x | x | x | x | x | x |  | x |  | x | x | x | x |  | x | x | x |
| Zhang et al. 2009 | x | x | x | x | x | x | x | x |  | x |  | x | x | x | x |  | x | x | x |
| Zhao et al. 2015 | x | x | x | x | x | x | x | x | x | x |  | x | x | x | x | x | x | x | x |
| Zyprian et al. 2016 | x | x | x | x | x | x | x | x | x | x | x | x | x | x | x | x | x | x | x |
| Number Of Maps Used To Construct Linkage Group | 39 | 39 | 37 | 39 | 39 | 38 | 38 | 37 | 35 | 39 | 26 | 39 | 37 | 37 | 37 | 35 | 39 | 39 | 37 |



Supplementary Figure 2. Correlation plot displaying Pearson correlation values for each pair of input genetic maps.

## Supplementary Table 3

## External file (Available upon request)

## Supplementary Table 4

External file (Available upon request)

Supplementary Table 5

| LG | Meta-QTL |  | $\mathbf{R}^{\mathbf{2}}$ | $\begin{aligned} & \text { Start } \\ & \text { (cM) } \end{aligned}$ | End $(\mathbf{c M})$ | Start (bp) | End (bp) | Candidates | Original QTLs co-located | QTL Studies <br> (Populations) | Reference |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| II | anthocyanin_2_1 | 49.04 | 0.560143 | 48.72 | 49.36 | 8600542 | 9173540 | 25 | 14 | 5 | Carreño Ruiz 2012, Costantini et al. 2015, Azuma et al. 2015, Guo et al. 2015, Bayo Canha PhD thesis 2015 |
| II | anthocyanin_2_2 | 50.76 | 0.554231 | 50.315 | 51.205 | 10028560 | 10710740 | 22 | 13 | 3 | Costantini et al. 2015, Azuma et al. 2015, Bayo Canha PhD thesis 2015 |
| II | anthocyanin_2_3 | 52.4 | 0.567667 | 52.17 | 52.63 | 11130194 | 12095023 | 19 | 15 | 3 | Costantini et al. 2015, Azuma et al. 2015, Bayo Canha PhD thesis 2015 |
| II | anthocyanin_2_4 | 52.94 | 0.563625 | 52.76 | 53.12 | 12753898 | 14578477 | 57 | 16 | 3 | Costantini et al. 2015, Azuma et al. 2015, Bayo Canha PhD thesis 2015 |
| II | anthocyanin_2_5 | 54.03 | 0.561125 | 53.86 | 54.2 | 16667197 | 16683683 | 1 | 16 | 3 | Costantini et al. 2015, Azuma et al. 2015, Bayo Canha PhD thesis 2015 |
| II | anthocyanin_2_6 | 54.4 | 0.574389 | 54.4 | 54.4 | 16693380 | 16693380 | 0 | 18 | 3 | Costantini et al. 2015, Azuma et al. 2015, Bayo Canha PhD thesis 2015 |
| II | anthocyanin_2_7 | 56.58 | 0.588667 | 56.51 | 56.65 | 17799696 | 17822818 | 1 | 12 | 2 | Costantini et al. 2015, Bayo Canha PhD thesis 2015 |

Supplementary Table 6

| Meta-QTL | Gene ID | Chr | Start (bp) | End(bp) | Annotation_V1 |
| :---: | :---: | :---: | :---: | :---: | :---: |
| anthocyanin_2_1 | VIT_02s0012g01940 | 2 | 8639809 | 8644529 | Myb KAN2 (KANADI 2) |
|  | VIT_02s0012g01950 | 2 | 8684996 | 8685382 | Photosystem II protein D1 |
|  | VIT_02s0012g01960 | 2 | 8685632 | 8694337 | Transcription factor jumonji (jmj) |
|  | VIT_02s0012g01970 | 2 | 8725299 | 8725604 | No hit |
|  | VIT_02s0012g01980 | 2 | 8737610 | 8738292 | Valyl tRNA synthetase |
|  | VIT_02s0012g01990 | 2 | 8745023 | 8746458 | 4-hydroxyphenylpyruvate dioxygenase |
|  | VIT_02s0012g02000 | 2 | 8805267 | 8815050 | Binding |
|  | VIT_02s0012g02020 | 2 | 8846265 | 8846784 | GTP binding protein |
|  | VIT_02s0012g02030 | 2 | 8851816 | 8858007 | Homeobox-7 |
|  | VIT_02s0012g02050 | 2 | 8869228 | 8874707 | PAP/fibrillin family |
|  | VIT_02s0012g02060 | 2 | 8876177 | 8876668 | Calmodulin |
|  | VIT_02s0012g02070 | 2 | 8877753 | 8893038 | Dolichol phosphate-mannose biosynthesis regulatory protein |
|  | VIT_02s0012g02080 | 2 | 8903756 | 8904732 | No hit |
|  | VIT_02s0012g02090 | 2 | 8918570 | 8923074 | Unknown protein |
|  | VIT_02s0012g02110 | 2 | 8936032 | 8953681 | No hit |
|  | VIT_02s0012g02120 | 2 | 8961449 | 8968305 | Defender against cell death 1 (DAD1) |
|  | VIT_02s0012g02130 | 2 | 9009706 | 9010443 | Unknown |
|  | VIT_02s0012g02140 | 2 | 9020625 | 9024203 | No hit |
|  | VIT_02s0012g02150 | 2 | 9028389 | 9028757 | Reduced sugar response 4 RSR4 |
|  | VIT_02s0012g02160 | 2 | 9055038 | 9090180 | Glycerol-3-phosphate acyltransferase |
|  | VIT_02s0012g02170 | 2 | 9098719 | 9099667 | No hit |
|  | VIT_02s0012g02180 | 2 | 9103155 | 9107033 | Abhydrolase domain-containing protein |
|  | VIT_02s0012g02190 | 2 | 9124133 | 9129245 | Cellulose synthase CSLD2 |
|  | VIT_02s0012g02200 | 2 | 9133993 | 9144207 | No hit |
|  | VIT_02s0012g02210 | 2 | 9151187 | 9151438 | No hit |
| anthocyanin _2_2 | VIT_02s0012g02520 | 2 | 10095290 | 10107351 | Vacuolar protein sorting 25 |
|  | VIT_02s0012g02530 | 2 | 10144914 | 10157269 | Phosphomevalonate kinase |
|  | VIT_02s0012g02540 | 2 | 10157476 | 10159508 | Chlororespiratory reduction 4 (CRR4) |
|  | VIT_02s0012g02560 | 2 | 10211013 | 10211294 | No hit |
|  | VIT_02s0012g02570 | 2 | 10303823 | 10313431 | Guanine nucleotide-binding protein alpha-1 subunit |
|  | VIT_02s0012g02580 | 2 | 10333777 | 10333986 | No hit |
|  | VIT_02s0012g02590 | 2 | 10333987 | 10335087 | Galacturonosyltransferase 1 |
|  | VIT_02s0012g02600 | 2 | 10337139 | 10337655 | Octicosapeptide/Phox/Bemlp (PB1) domain-containing protein |
|  | VIT_02s0012g02610 | 2 | 10346830 | 10347282 | SAR1 (suppressor of auxin resistance1) ( Nup160) |
|  | VIT_02s0012g02620 | 2 | 10358627 | 10359350 | PFT1 (phytochrome and flowering time 1) MED25 |
|  | VIT_02s0012g02640 | 2 | 10371559 | 10372237 | Aspartic-type endopeptidase |
|  | VIT_02s0012g02650 | 2 | 10379257 | 10380646 | Purple acid phosphatase 2 PAP2 |
|  | VIT_02s0012g02660 | 2 | 10381950 | 10405928 | Proteasome 26S regulatory subunit (RPN9) |
|  | VIT_02s0012g02670 | 2 | 10482684 | 10485983 | Unknown protein |
|  | VIT_02s0012g02680 | 2 | 10538523 | 10542564 | No hit |
|  | VIT_02s0012g02690 | 2 | 10556046 | 10556298 | No hit |
|  | VIT_02s0012g02700 | 2 | 10603213 | 10607154 | Unknown |
|  | VIT_02s0012g02710 | 2 | 10617021 | 10618719 | Unknown |
|  | VIT_02s0012g02720 | 2 | 10620819 | 10623614 | RPM1 (resistance to p. syringae pv maculicola 1) |


|  | VIT_02s0012g02760 | 2 | 10685743 | 10699497 | Methionyl-tRNA synthetase |
| :---: | :---: | :---: | :---: | :---: | :---: |
|  | VIT_02s0012g02770 | 2 | 10703356 | 10703813 | ABC transporter g family pleiotropic drug resistance 12 PDR 12 |
|  | VIT_02s0012g02780 | 2 | 10706920 | 10720569 | Cytochrome b5 domain-containing protein |
| anthocyanin _2_3 | VIT_02s0012g02920 | 2 | 11133984 | 11136347 | Acyl-CoA oxidase ACX3 |
|  | VIT_02s0012g02970 | 2 | 11183822 | 11186832 | No hit |
|  | VIT_02s0012g03040 | 2 | 11666980 | 11687641 | AAA-type ATPase |
|  | VIT_02s0012g03050 | 2 | 11688053 | 11691156 | AAA-type ATPase |
|  | VIT_02s0012g03060 | 2 | 11710057 | 11710461 | 6-phosphogluconate dehydrogenase, decarboxylating |
|  | VIT_02s0012g03090 | 2 | 11751038 | 11752949 | AAA-type ATPase |
|  | VIT_02s0012g03100 | 2 | 11771975 | 11773559 | No hit |
|  | VIT_02s0012g03110 | 2 | 11774822 | 11776354 | CTV. 22 |
|  | VIT_02s0012g03130 | 2 | 11817589 | 11831312 | No hit |
|  | VIT_02s0012g03140 | 2 | 11831313 | 11832597 | Ribosomal protein S7 30S |
|  | VIT_02s0012g03160 | 2 | 11840324 | 11841710 | No hit |
|  | VIT_02s0012g03170 | 2 | 11873089 | 11874714 | No hit |
|  | VIT_02s0012g03180 | 2 | 11879297 | 11879706 | Unknown |
|  | VIT_02s0012g03190 | 2 | 11885508 | 11886222 | CTV. 22 |
|  | VIT_02s0012g03200 | 2 | 11893087 | 11894138 | Armadillo/beta-catenin repeat |
|  | VIT_02s0012g03210 | 2 | 11897304 | 11897873 | No hit |
|  | VIT_02s0012g03240 | 2 | 11947893 | 11959569 | AAA-type ATPase |
|  | VIT_02s0012g03250 | 2 | 11959578 | 11960203 | AAA-type ATPase |
|  | VIT_02s0012g03260 | 2 | 11987891 | 11988792 | No hit |
| anthocyanin _2_4 | VIT_02s0109g00230 | 2 | 12786746 | 12808921 | Early-responsive to dehydration protein / ERD protein |
|  | VIT_02s0109g00240 | 2 | 12814813 | 12815037 | No hit |
|  | VIT_02s0109g00250 | 2 | 12822109 | 12827876 | 4-coumarate-CoA ligase |
|  | VIT_02s0109g00260 | 2 | 12868384 | 12876281 | Retrotransposon gag protein |
|  | VIT_02s0109g00280 | 2 | 12912306 | 12929427 | No hit |
|  | VIT_02s0109g00290 | 2 | 12929477 | 12929908 | CYP76C6 |
|  | VIT_02s0109g00300 | 2 | 12929909 | 12931565 | CYP76B1 |
|  | VIT_02s0109g00310 | 2 | 12936238 | 12938714 | flavonoid 3-monooxygenase |
|  | VIT_02s0109g00320 | 2 | 12982398 | 12982920 | Translation initiation factor eIF-3 subunit 8 |
|  | VIT_02s0109g00350 | 2 | 13058347 | 13059533 | Carboxyl-terminal proteinase |
|  | VIT_02s0109g00360 | 2 | 13116317 | 13119876 | Translocase of chloroplast 34 |
|  | VIT_02s0109g00370 | 2 | 13163595 | 13167989 | RNA recognition motif (RRM)-containing protein |
|  | VIT_02s0109g00380 | 2 | 13170239 | 13193614 | Stress response suppressor 1 |
|  | VIT_02s0109g00390 | 2 | 13282201 | 13282765 | N-hydroxycinnamoyl/benzoyltransferase 5 |
|  | VIT_02s0109g00400 | 2 | 13282778 | 13283707 | Serine-type peptidase SLP3 |
|  | VIT_02s0109g00410 | 2 | 13308757 | 13312393 | No hit |
|  | VIT_02s0109g00420 | 2 | 13341559 | 13342323 | Disease resistance protein (CC-NBS-LRR class) |
|  | VIT_02s0109g00430 | 2 | 13347364 | 13349570 | Nitrilase 4 (NIT4) |
|  | VIT_02s0109g00440 | 2 | 13353462 | 13353985 | No hit |
|  | VIT_02s0033g00010 | 2 | 13457765 | 13460162 | Unknown protein |
|  | VIT_02s0033g00020 | 2 | 13460163 | 13461470 | Unknown protein |
|  | VIT_02s0033g00030 | 2 | 13506518 | 13511727 | No hit |
|  | VIT_02s0033g00040 | 2 | 13512257 | 13512649 | No hit |
|  | VIT_02s0033g00050 | 2 | 13518893 | 13519179 | Scarecrow transcription factor 3 (SCL3) |
|  | VIT_02s0033g00060 | 2 | 13531787 | 13538078 | DNAJ plastid division protein (ARC6) |


|  | VIT_02s0033g00070 | 2 | 13568564 | 13569348 | No hit |
| :---: | :---: | :---: | :---: | :---: | :---: |
|  | VIT_02s0033g00080 | 2 | 13590737 | 13637352 | Pleckstriny ( PH ) domain-containing protein |
|  | VIT_02s0033g00090 | 2 | 13639223 | 13640252 | DNA replication licensing factor MCM2 |
|  | VIT_02s0033g00120 | 2 | 13705067 | 13785560 | Pleckstriny ( PH ) domain-containing protein |
|  | VIT_02s0033g00130 | 2 | 13786111 | 13787768 | Cis-zeatin O-beta-D-glucosyltransferase |
|  | VIT_02s0033g00150 | 2 | 13832782 | 13833455 | No hit |
|  | VIT_02s0033g00160 | 2 | 13856081 | 13859536 | Pleckstriny (PH) domain-containing protein |
|  | VIT_02s0033g00180 | 2 | 13880089 | 13881941 | Phenylpropanoid:glucosyltransferase 2 |
|  | VIT_02s0033g00190 | 2 | 13884200 | 13933646 | DNA replication licensing factor MCM2 |
|  | VIT_02s0033g00230 | 2 | 13998724 | 14007202 | Unknown protein |
|  | VIT_02s0033g00240 | 2 | 14031671 | 14033252 | Glucosyltransferase twil |
|  | VIT_02s0033g00250 | 2 | 14051742 | 14052882 | No hit |
|  | VIT_02s0033g00260 | 2 | 14104114 | 14107827 | Pinoresinol-lariciresinol reductase |
|  | VIT_02s0033g00270 | 2 | 14123184 | 14126331 | Pinoresinol-lariciresinol reductase |
|  | VIT_02s0033g00290 | 2 | 14139881 | 14141542 | Pinoresinol-lariciresinol reductase |
|  | VIT_02s0033g00300 | 2 | 14144838 | 14148929 | myb family |
|  | VIT_02s0033g00310 | 2 | 14158366 | 14159002 | No hit |
|  | VIT_02s0033g00320 | 2 | 14159003 | 14183582 | PRLI-interacting factor L |
|  | VIT_02s0033g00350 | 2 | 14253249 | 14254161 | No hit |
|  | VIT_02s0033g00360 | 2 | 14265226 | 14266015 | Urease |
|  | VIT_02s0033g00370 | 2 | 14275372 | 14275752 | myb domain protein 113 |
|  | VIT_02s0033g00380 | 2 | 14291438 | 14291683 | VvMybA2 (C-term) |
|  | VIT_02s0033g00390 | 2 | 14291727 | 14292732 | VvMybA2 |
|  | VIT_02s0033g00400 | 2 | 14302187 | 14303282 | myb domain protein 113 |
|  | VIT_02s0033g00410 | 2 | 14351791 | 14352807 | VvMybA1 |
|  | VIT_02s0033g00420 | 2 | 14386896 | 14387181 | Ribosomal protein S28 (RPS28C) 40S |
|  | VIT_02s0033g00430 | 2 | 14392775 | 14393842 | myb domain protein 113 |
|  | VIT_02s0033g00440 | 2 | 14400037 | 14401092 | Myb VvMYBA1 |
|  | VIT_02s0033g00450 | 2 | 14420525 | 14421283 | VvMybA3 |
|  | VIT_02s0033g00460 | 2 | 14425416 | 14426478 | myb domain protein 113 |
|  | VIT_02s0033g00480 | 2 | 14498732 | 14499419 | No hit |
|  | VIT_02s0033g00490 | 2 | 14567216 | 14568104 | No hit |
| anthocyanin _2_5 | VIT_02s0033g01190 | 2 | 16677461 | 16690102 | Unknown protein |
| anthocyanin _2_6 |  |  |  |  |  |
| anthocyanin _2_7 | VIT_02s0087g00390 | 2 | 17799720 | 17825371 | Endonuclease |

Supplementary Table 7. Description of the phenology meta-QTLs

| LG | MetaQTL | Peak Position (cM) | $\mathbf{R}^{2}$ | $\begin{aligned} & \text { Start } \\ & \text { (cM) } \end{aligned}$ | $\begin{aligned} & \text { End } \\ & \text { (cM) } \end{aligned}$ | $\begin{aligned} & \text { Original QTLs } \\ & \text { co-located } \end{aligned}$ | QTL Studies <br> (Populations) | Traits | Reference |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| I | ver_1_1 | 31.29 | 0.11 | 29.15 | 33.43 | 2 | 2 | VT | Fechter et al 2014, Zyprian et al 2016 |
|  | pheno_1_1 | 30.30 | 0.15 | 28.08 | 32.53 | 9 | 5 | VT, FBL, FT, FS, RT | Carreño Ruiz 2012, Costantini et al 2008, Fechter et al 2014, Zyprian et al 2016 |
|  | pheno_1_2 | 37.94 | 0.15 | 36.68 | 39.21 | 9 | 3 | VT, FBL, RT, FS | Carreño Ruiz 2012, Fechter et al 2014, Zyprian et al 2016, |
| II | ver_2_1 | 31.34 | 0.17 | 28.79 | 33.89 | 7 | 2 | $\mathrm{VB}, \mathrm{Vr}$ | Bayo Canha 2015, Grzeskowiak et al 2013 |
|  | ver_2_2 | 41.55 | 0.13 | 40.00 | 43.30 | 5 | 3 | Vr, VB, VE, VT | Costantini et al 2008, Bayo Canha 2015, Grzeskowiak et al 2013 |
|  | ver_2_3 | 53.47 | 0.34 | 52.88 | 54.07 | 5 | 3 | Vr, VE, VP | Costantini et al 2008, Bayo Canha 2015, Grzeskowiak et al 2013 |
|  | pheno_2_1 | 31.22 | 0.18 | 30.03 | 32.42 | 15 | 5 | $\begin{gathered} \text { Ac, VE, Rp, FT, ssc, VB, VE, } \\ \text { V-R, Vr, Vr-Rp } \end{gathered}$ | Carreño Ruiz 2012, Costantini et al 2008, Ban et al 2016, Bayo Canha 2015, Grzeskowiak et al 2013 |
|  | pheno_2_2 | 41.58 | 0.18 | 40.23 | 42.94 | 9 | 4 | $\begin{gathered} \mathrm{Ac}, \mathrm{VB}, \mathrm{FT}, \mathrm{Rp}, \mathrm{VE}, \mathrm{Vr}, \mathrm{Vr-} \\ \mathrm{Rp}, \mathbf{V T} \end{gathered}$ | Carreño Ruiz 2012, Costantini et al 2008, Bayo Canha 2015, Grzeskowiak et al 2013 |

pheno_2_3
48.83
53.60
pheno 24 .

Supplementary Table 8

| Meta- <br> QTL | Gene ID | Chr | Start <br> (bp) | End(bp) |
| :---: | :---: | :---: | :---: | :---: | Annotation_V1 (


| VIT_01s0011g03040 | 1 | 2714906 | 2715758 |
| :--- | :--- | :--- | :--- |
| VIT_01s0011g03050 | 1 | 2717642 | 2719224 |
| VIT_01s0011g03060 | 1 | 2726727 | 2743646 |
| VIT_01s0011g03070 | 1 | 2751566 | 2753036 |
| VIT_01s0011g03080 | 1 | 2754305 | 2765766 |
| VIT_01s0011g03090 | 1 | 2766597 | 2767444 |
| VIT_01s0011g03100 | 1 | 2770673 | 2771827 |
| VIT_01s0011g03110 | 1 | 2781518 | 2783517 |
| VIT_01s0011g03120 | 1 | 2786238 | 2790806 |
| VIT_01s0011g03130 | 1 | 2791893 | 2796533 |
| VIT_01s0011g03140 | 1 | 2806814 | 2809091 |
| VIT_01s0011g03160 | 1 | 2819342 | 2877060 |
| VIT_01s0011g03170 | 1 | 2877767 | 2878493 |
| VIT_01s0011g03180 | 1 | 2879352 | 2882108 |
| VIT_01s0011g03190 | 1 | 2886723 | 2897786 |
| VIT_01s0011g03200 | 1 | 2898690 | 2906357 |
| VIT_01s0011g03210 | 1 | 2924829 | 2926924 |
| VIT_01s0011g03220 | 1 | 2928959 | 2935184 |
| VIT_01s0011g03230 | 1 | 2941266 | 2946275 |
| VIT_01s0011g03240 | 1 | 2948265 | 2949046 |
| VIT_01s0011g03250 | 1 | 2951539 | 2951847 |
| VIT_01s0011g03260 | 1 | 2954243 | 2955469 |
| VIT_01s0011g03270 | 1 | 2958052 | 2959131 |
| VIT_01s0011g03280 | 1 | 2967860 | 2979364 |
| VIT_01s0011g03290 | 1 | 2981226 | 2996541 |
| VIT_01s0011g03300 | 1 | 3007049 | 3007411 |
| VIT_01s0011g03310 | 1 | 3008153 | 3011404 |
| VIT_01s0011g03320 | 1 | 3011621 | 3013806 |
| VIT_01s0011g03330 | 1 | 3014379 | 3016292 |
| VIT_01s0011g03340 | 1 | 3028681 | 3030154 |
| VIT_01s0011g03350 | 1 | 3030155 | 3040612 |
| VIT_01s0011g03360 | 1 | 3044557 | 3045849 |
| VIT_01s0011g03370 | 1 | 3046873 | 3051454 |
| VIT_01s0011g03380 | 1 | 3061306 | 3065510 |
| VIT_01s0011g03390 | 1 | 3065990 | 3068987 |
| VIT_01s0011g03400 | 1 | 3076982 | 3080343 |
|  |  |  |  |
| VIs |  |  |  |

Unknown protein
Unknown protein
Symbiosis receptor-like kinase
ERF/AP2 Gene Family (VvRAV1)
Methionine aminopeptidase 1B, chloroplast precursor Allene oxide cyclase (jasmonates from fatty acids) Zinc finger (C2H2 type) protein (WIP6)

## myb family

Unknown protein
Unknown protein
PFT1 (phytochrome and flowering time 1) MED25 PFT1 (phytochrome and flowering time 1) MED25 PFT1 (phytochrome and flowering time 1) MED25 Lysine and histidine specific transporter ATP-dependent Cl p protease adaptor protein ClpS containing protein Transmembrane protein FT27/PFT27

Aspartic Protease (VvAP1)
Choline transporter
Basic Leucine Zipper Transcription Factor (VvbZIP01)
PGPS/D12
Fw2.2 ORFX
Fw2.2 ORFX
CXE carboxylesterase
Protein phosphatase 2 regulatory subunit A
Aluminum-activated malate transporter 9
Plastid-specific 30S ribosomal protein 3
Unknown protei
Arabidopsis histidine phosphotransfer AHP2
Unknown protein
No hi
PRP4 pre-mRNA processing factor 4 homolog B
Unknown
Agenet domain-containing protein Dihydrolipoamide S-acetyltransferase (LTA2)

Unknown protein
Proton-dependent oligopeptide transport (POT) family protein

|  | VIT_01s0011g03410 | 1 | 3084191 | 3092672 | DNA repair protein RAD23 |
| :---: | :---: | :---: | :---: | :---: | :---: |
|  | VIT_01s0011g03420 | 1 | 3100807 | 3101627 | Unknown protein |
|  | VIT_01s0011g03430 | 1 | 3112762 | 3113407 | No hit |
|  | VIT_01s0011g03440 | 1 | 3117073 | 3117578 | DNA mismatch repair protein |
|  | VIT_01s0011g03450 | 1 | 3127812 | 3132261 | Alpha-glucosidase |
|  | VIT_01s0011g03460 | 1 | 3141923 | 3142868 | No hit |
|  | VIT_01s0011g03470 | 1 | 3142869 | 3144296 | ERF/AP2 Gene Family (VvERF122) |
|  | VIT_01s0011g03480 | 1 | 3147476 | 3149151 | Cinnamoyl CoA reductase |
|  | VIT_01s0011g03490 | 1 | 3167509 | 3169015 | 3-ketoacyl-CoA synthase |
|  | VIT_01s0011g03500 | 1 | 3175430 | 3178998 | Homocysteine S-methyltransferase 1 |
|  | VIT_01s0011g03510 | 1 | 3180020 | 3185492 | Protein kinase PKN/PRK1 |
|  | VIT_01s0011g03520 | 1 | 3190826 | 3192777 | Constans-like 16 |
|  | VIT_01s0011g03530 | 1 | 3204646 | 3205573 | Lateral organ boundaries protein 41 |
|  | VIT_01s0011g03540 | 1 | 3210273 | 3211389 | Lateral organ boundaries protein 41 |
|  | VIT_01s0011g03550 | 1 | 3217781 | 3218222 | Unknown |
|  | VIT_01s0011g03560 | 1 | 3223281 | 3227712 | fiber protein Fb 34 |
|  | VIT_01s0011g03570 | 1 | 3232043 | 3232750 | Unknown |
|  | VIT_01s0011g03580 | 1 | 3235418 | 3237772 | Unknown protein |
|  | VIT_01s0011g03590 | 1 | 3246882 | 3249663 | Ribosomal protein L15, chloroplast (CL15) 50S |
|  | VIT_01s0011g03600 | 1 | 3252820 | 3255824 | JAGGED |
| ver_2_1 | VIT_02s0025g04660 | 2 | 4037933 | 4039464 | Senescence-inducible chloroplast stay-green protein 1 |
|  | VIT_02s0025g04670 | 2 | 4043652 | 4045210 | Senescence-inducible chloroplast stay-green protein 2 |
|  | VIT_02s0025g04680 | 2 | 4047267 | 4056403 | Cell division cycle 20-like protein 1 |
|  | VIT_02s0025g04690 | 2 | 4059360 | 4060286 | Unknown protein |
|  | VIT_02s0025g04700 | 2 | 4061719 | 4063223 | Expansin (VvEXPA2) |
|  | VIT_02s0025g04710 | 2 | 4064018 | 4073343 | Unknown protein |
|  | VIT_02s0025g04720 | 2 | 4094895 | 4096363 | Leucoanthocyanidin dioxygenase (VvLDOX) [Vitis vinifera] |
|  | VIT_02s0025g04730 | 2 | 4100066 | 4103095 | Glyoxylate reductase |
|  | VIT_02s0025g04750 | 2 | 4108905 | 4111169 | Glycerate dehydrogenase |
|  | VIT_02s0025g04760 | 2 | 4121700 | 4124366 | Splicing factor YT521-B |
|  | VIT_02s0025g04770 | 2 | 4135934 | 4139911 | Unknown protein |
|  | VIT_02s0025g04780 | 2 | 4141626 | 4144449 | Subtilisin stomatal density and distribution |
|  | VIT_02s0025g04790 | 2 | 4145024 | 4146976 | Hydrogenobyrinic acid a,c-diamide synthase |
|  | VIT_02s0025g04800 | 2 | 4149493 | 4151715 | Hydrogenobyrinic acid a,c-diamide synthase |
|  | VIT_02s0025g04810 | 2 | 4153569 | 4155803 | Subtilisin P69F protein |
|  | VIT_02s0025g04820 | 2 | 4166469 | 4168208 | Subtilisin P69E protein |


| VIT_02s0025g04830 | 2 | 4169732 | 4176688 |
| :--- | :--- | :--- | :--- |
| VIT_02s0025g04840 | 2 | 4177464 | 4181953 |
| VIT_02s0025g04850 | 2 | 4191899 | 4193146 |
| VIT_02s0025g04860 | 2 | 4193239 | 4194582 |
| VIT_02s0025g04870 | 2 | 4210862 | 4211722 |
| VIT_02s0025g04880 | 2 | 4212083 | 4213721 |
| VIT_02s0025g04890 | 2 | 4216365 | 4217374 |
| VIT_02s0025g04900 | 2 | 4226153 | 4229768 |
| VIT_02s0025g04910 | 2 | 4233942 | 4235225 |
| VIT_02s0025g04920 | 2 | 4236968 | 4238438 |
| VIT_02s0025g04930 | 2 | 4243946 | 4246738 |
| VIT_02s0025g04940 | 2 | 4246754 | 4248676 |
| VIT_02s0025g04950 | 2 | 4251382 | 4252445 |
| VIT_02s0025g04960 | 2 | 4257725 | 4263946 |
| VIT_02s0025g04970 | 2 | 4267189 | 4268564 |
| VIT_02s0025g04980 | 2 | 4274342 | 4280444 |
| VIT_02s0025g04990 | 2 | 4282395 | 4290690 |
| VIT_02s0025g05000 | 2 | 4300566 | 4305985 |
| VIT_02s0025g05010 | 2 | 4307128 | 4313693 |
| VIT_02s0025g05020 | 2 | 4313694 | 4314578 |
| VIT_02s0025g05030 | 2 | 4314983 | 4315368 |
| VIT_02s0025g05040 | 2 | 4315576 | 4316706 |
| VIT_02s0025g05050 | 2 | 4321904 | 4323519 |
| VIT_02s0025g05060 | 2 | 4324307 | 4346662 |
| VIT_02s0025g05070 | 2 | 4346663 | 4347651 |
| VIT_02s0025g05080 | 2 | 4348262 | 4351307 |
| VIT_02s0025g05090 | 2 | 4356581 | 4373218 |
| VIT_02s0025g05100 | 2 | 4376256 | 4377810 |
| VIT_02s0025g05110 | 2 | 4390487 | 4392478 |
| VIT_02s0025g05120 | 2 | 4410051 | 4413324 |
| VIT_02s0025g05130 | 2 | 4421391 | 4429393 |
| VIT_02s0025g05140 | 2 | 4437507 | 4440897 |
| VIT_02s0025g05150 | 2 | 4444146 | 4448453 |
| VIT_02s0025g05160 | 2 | 4450431 | 4454056 |
| VIT_02s0025g05170 | 2 | 4454528 | 4456260 |
| VIT_00s2563g00010 | 2 | 4456792 | 4458501 |
|  |  |  |  |

Copper chaperone for superoxide dismutase Nudix hydrolase 14

CYP76B1
CYP76B1
No hit
Geraniol 10-hydroxylase
CYP76B1
Phosphoglycerate mutase Growth-regulating factor 5
Phosphate translocator protein2, Plastidic
CDI3/OZS1/RCD3/SLAC1 (slow anion channel-associated 1)
Carboxyl-terminal peptidase
No hit
Unknown protein Subtilisin protease 1-aminocyclopropane-1-carboxylate synthase

Unknown protein
Bile acid:sodium symporter

Unknown protein
Nuclear pore complex component Zf A20 and AN1 domain-containing stress-associated protein 1 Microtubule motor PAK (phosphatidic acid kinase) KHC Microtubule motor PAK (phosphatidic acid kinase) KHC Zf A20 and AN1 domain-containing stress-associated protein 1 Microtubule motor PAK (phosphatidic acid kinase) KHC

AT-hook DNA-binding protein MATE efflux family protein ACT domain-containing protein (ACR8) High mobility group HMG-I and HMG-Y, DNA-binding ATP-dependent Clp protease proteolytic subunit (ClpP2) TFIIH basal transcription factor complex TTD-A subunit Cornichon family
Unknown protein
Glycogenin glucosyltransferase

| VIT_02s0241g00010 | 2 | 4481714 | 4483309 |
| :--- | :--- | :--- | :--- |
| VIT_02s0241g00020 | 2 | 4486591 | 4488109 |
| VIT_02s0241g00030 | 2 | 4497165 | 4497380 |
| VIT_02s0241g00040 | 2 | 4500666 | 4508486 |
| VIT_02s0241g00050 | 2 | 4516706 | 4519215 |
| VIT_02s0241g00060 | 2 | 4528876 | 4533648 |
| VIT_02s0241g00070 | 2 | 4545465 | 4546112 |
| VIT_02s0241g00080 | 2 | 4546113 | 4551868 |
| VIT_02s0241g00090 | 2 | 4553230 | 4560602 |
| VIT_02s0241g00100 | 2 | 4560763 | 4561723 |
| VIT_02s0241g00110 | 2 | 4563842 | 4567239 |
| VIT_02s0241g00120 | 2 | 4567672 | 4570586 |
| VIT_02s0241g00130 | 2 | 4572517 | 4573830 |
| VIT_02s0241g00140 | 2 | 4577280 | 4577983 |
| VIT_02s0241g00150 | 2 | 4581818 | 4583479 |
| VIT_02s0241g00160 | 2 | 4583480 | 4597122 |
| VIT_02s0241g00170 | 2 | 4599598 | 4602678 |
| VIT_02s0241g00180 | 2 | 4617301 | 4618383 |
| VIT_02s0241g00190 | 2 | 4626070 | 4626875 |
| VIT_00s0555g00010 | 2 | 4641190 | 4641853 |
| VIT_00s0555g00020 | 2 | 4648183 | 4650503 |
| VIT_00s0555g00050 | 2 | 4665292 | 4666049 |
| VIT_00s0323g00010 | 2 | 4668106 | 4670512 |
| VIT_00s0323g00020 | 2 | 4671193 | 4672013 |
| VIT_00s0323g00030 | 2 | 4678930 | 4682628 |
| VIT_00s0323g00040 | 2 | 4697703 | 4701979 |
| VIT_00s0323g00050 | 2 | 4703890 | 4704749 |
| VIT_00s0323g00060 | 2 | 4707844 | 4708734 |
| VIT_00s0323g00070 | 2 | 4716445 | 4717351 |
| VIT_00s0323g00080 | 2 | 4728265 | 4732585 |
| VIT_00s0323g00100 | 2 | 4758426 | 4761706 |
| VIT_02s0154g00010 | 2 | 4765224 | 4766070 |
| VIT_02s0154g00020 | 2 | 4779757 | 4782259 |
| VIT_02s0154g00030 | 2 | 4784152 | 4784510 |
| VIT_02s0154g00040 | 2 | 4788746 | 4794088 |
| VIT_02s0154g00050 | 2 | 4794446 | 4795025 |
|  |  |  |  |

Pentatricopeptide (PPR) repeat-containing protein
Unknown
Sterol 4-alpha-methyl-oxidase 1 (SMO1)
Cyclobutane pyrimidine dimer photolyase
Unknown protein
Unknown protein
Zinc finger (DHHC type)
Glycine-rich protein-like Translation release factor

Unknown protein Aminomethyltransferase Receptor protein kinase

No hit
Calmodulin (A)

## No hit

Esterase/lipase/thioesterase family Zinc finger ( C 2 H 2 type) family
UDP-D- glcucuronate 4-epimerase 5 GAE5
No hit
CYP706A12
flavonoid 3-monooxygenase
CHCH domain containing protein
No hit
HSP associated protein
HSP associated protein
Unknown protein
Invertase/pectin methylesterase inhibitor Invertase/pectin methylesterase inhibitor Pectin methylesterase inhibitor Elongation factor G, chloroplast precursor Vacuolar processing enzyme beta Auxin-responsive SAUR11
NAC domain-containing protein (VvNAC02)
No hit
Thylakoid lumenal protein Glycosyl hydrolase family 5

| VIT_02s0154g00060 | 2 | 4795026 | 4798608 |
| :--- | :--- | :--- | :--- |
| VIT_02s0154g00070 | 2 | 4804832 | 4807460 |
| VIT_02s0154g00080 | 2 | 4813347 | 4818031 |
| VIT_02s0154g00090 | 2 | 4824906 | 4827102 |
| VIT_02s0154g00100 | 2 | 4830177 | 4840110 |
| VIT_02s0154g00110 | 2 | 4840895 | 4843817 |
| VIT_02s0154g00120 | 2 | 4856955 | 4857771 |
| VIT_02s0154g00130 | 2 | 4864567 | 4866368 |
| VIT_02s0154g00140 | 2 | 4866564 | 4878108 |
| VIT_02s0154g00150 | 2 | 4891997 | 4893321 |
| VIT_02s0154g00160 | 2 | 4895022 | 4898791 |
| VIT_02s0154g00170 | 2 | 4900531 | 4903560 |
| VIT_02s0154g00180 | 2 | 4905399 | 4908515 |
| VIT_02s0154g00190 | 2 | 4917402 | 4920926 |
| VIT_02s0154g00200 | 2 | 4926234 | 4929441 |
| VIT_02s0154g00210 | 2 | 4932248 | 4932565 |
| VIT_02s0154g00220 | 2 | 4968010 | 4968965 |
| VIT_02s0154g00230 | 2 | 4979771 | 4980739 |
| VIT_02s0154g00240 | 2 | 5005578 | 5017004 |
| VIT_02s0154g00250 | 2 | 5021111 | 5041753 |
| VIT_02s0154g00260 | 2 | 5048952 | 5052974 |
| VIT_02s0154g00270 | 2 | 5059261 | 5059470 |
| VIT_02s0154g00280 | 2 | 5063840 | 5064500 |
| VIT_02s0154g00290 | 2 | 5068196 | 5068884 |
| VIT_02s0154g00300 | 2 | 5073755 | 5074467 |
| VIT_02s0154g00310 | 2 | 5077143 | 5078172 |
| VIT_02s0154g00320 | 2 | 5084359 | 5085114 |
| VIT_02s0154g00330 | 2 | 5090858 | 5100329 |
| VIT_02s0154g00340 | 2 | 5100931 | 5102577 |
| VIT_02s0154g00350 | 2 | 5107218 | 5109108 |
| VIT_02s0154g00360 | 2 | 5109216 | 5109392 |
| VIT_02s0154g00370 | 2 | 5110262 | 5115144 |
| VIT_02s0154g00380 | 2 | 5117118 | 5118512 |
| VIT_02s0154g00390 | 2 | 5127909 | 5131694 |
| VIT_02s0154g00400 | 2 | 5145761 | 5147651 |
| VIT_02s0154g00410 | 2 | 5164705 | 5167011 |
|  |  |  |  |

Thylakoid lumenal protein
Abnormal floral organs
Multi-copper oxidase (SKU5)
Vacuolar invertase 2, GIN2
Transaldolase totaL2
Trehalose-6-phosphate phosphatase (AtTPPA)
Unknown
Exostosin (Xyloglucan galactosyltransferase KATAMARI 1) 3-oxoacyl-[acyl-carrier-protein] synthase 3 A , chloroplast precursor

PLATZ transcription factor
flavin-containing monooxygenase family protein / FMO family protein flavin-containing monooxygenase 3 flavin-containing monooxygenase 3 flavin-containing monooxygenase 3

Unknown protein
WRKY DNA-binding protein 21
Zinc finger ( C 2 H 2 type) family
Phosphatidic acid phosphatase / PAP2
Oxysterol binding protein
Oxysterol binding protein
Nitrate transporter

## No hit

Arachidonic acid-induced DEA1
Extensin
Small nuclear ribonucleoprotein Sm D3 Protease inhibitor/seed storage/lipid transfer protein (LTP) Protease inhibitor/seed storage/lipid transfer protein (LTP)

L-lactate dehydrogenase A
No hit
$\mathrm{YbaK} /$ prolyl-tRNA synthetase associated region

Scarecrow transcription factor 6 (SCL6)

|  | VIT_02s0154g00420 | 2 | 5169940 | 5170164 | No hit |
| :---: | :---: | :---: | :---: | :---: | :---: |
|  | VIT_02s0154g00430 | 2 | 5170165 | 5170573 | Unknown |
|  | VIT_02s0154g00440 | 2 | 5170574 | 5171512 | Unknown protein |
|  | VIT_02s0154g00450 | 2 | 5174203 | 5183898 | Zinc knuckle |
|  | VIT_02s0154g00460 | 2 | 5184751 | 5188467 | Unknown protein |
|  | VIT_02s0154g00470 | 2 | 5188779 | 5190703 | No hit |
|  | VIT_02s0154g00480 | 2 | 5191329 | 5192828 | Heat shock protein MTSHP |
|  | VIT_02s0154g00490 | 2 | 5197928 | 5198941 | Heat shock 22 kDa protein mitochondrial |
|  | VIT_02s0154g00500 | 2 | 5201356 | 5203907 | VAP27-1 (VAMP/synaptobrevin-associated protein 27-1) |
|  | VIT_02s0154g00510 | 2 | 5205962 | 5214005 | Aspartic Protease (VvAP2) |
|  | VIT_02s0154g00520 | 2 | 5216152 | 5221223 | Aspartyl protease |
|  | VIT_02s0154g00530 | 2 | 5227339 | 5229999 | Histidine triad nucleotide binding protein 3 |
|  | VIT_02s0154g00540 | 2 | 5231330 | 5239367 | Protein arginine N-methyltransferase |
|  | VIT_02s0154g00550 | 2 | 5239749 | 5268651 | DnaJ homolog, subfamily C, member 11 |
|  | VIT_02s0154g00560 | 2 | 5280620 | 5283758 | No hit |
|  | VIT_02s0154g00580 | 2 | 5295292 | 5296235 | Unknown |
|  | VIT_02s0154g00590 | 2 | 5308401 | 5308985 | Unknown |
|  | VIT_02s0154g00600 | 2 | 5321389 | 5323536 | Pectinesterase family |
|  | VIT_02s0154g00610 | 2 | 5324724 | 5332701 | Pex19 protein |
|  | VIT_02s0154g00620 | 2 | 5333217 | 5334885 | Unknown |
|  | VIT_00s1338g00010 | 2 | 5339303 | 5340709 | DNA-directed RNA polymerase III subunit C34 |
|  | VIT_00s1338g00020 | 2 | 5343106 | 5344525 | Protein transport protein Sec61 subunit alpha |
|  | VIT_00s0229g00010 | 2 | 5346706 | 5349092 | No hit |
| ver_2_2 | VIT_02s0012g00010 | 2 | 5737022 | 5737770 | No hit |
|  | VIT_02s0012g00020 | 2 | 5764306 | 5772117 | Glycine-rich protein |
|  | VIT_02s0012g00030 | 2 | 5772754 | 5778969 | Unknown |
|  | VIT_02s0012g00040 | 2 | 5781659 | 5781796 | No hit |
|  | VIT_02s0012g00050 | 2 | 5786859 | 5796200 | Glycine-rich protein |
|  | VIT_02s0012g00060 | 2 | 5796703 | 5801313 | Unknown |
|  | VIT_02s0012g00070 | 2 | 5801456 | 5801882 | Unknown protein |
|  | VIT_02s0012g00080 | 2 | 5804657 | 5813125 | Casein kinase II subunit beta-4 |
|  | VIT_02s0012g00090 | 2 | 5813865 | 5826392 | Phosphatidylinositol-4-phosphate 5-kinase |
|  | VIT_02s0012g00100 | 2 | 5828133 | 5830613 | Ribosomal protein L37a (RPL37aB) 60S |
|  | VIT_02s0012g00110 | 2 | 5835376 | 5837313 | Chromatin remodeling 31 |
|  | VIT_02s0012g00140 | 2 | 5869675 | 5887801 | Novel plant snare 11 |
|  | VIT_02s0012g00150 | 2 | 5897718 | 5902809 | NAK-type protein kinase |


| VIT_02s0012g00160 | 2 | 5910037 | 5913981 | Unknown |
| :---: | :---: | :---: | :---: | :---: |
| VIT_02s0012g00170 | 2 | 5916093 | 5918986 | 1,4-alpha-D-glucan maltohydrolase |
| VIT_02s0012g00180 | 2 | 5923457 | 5932422 | Serine protease |
| VIT_02s0012g00190 | 2 | 5938773 | 5978552 | Unknown protein |
| VIT_02s0012g00240 | 2 | 5999054 | 6002113 | Unknown protein |
| VIT_02s0012g00250 | 2 | 6005214 | 6012306 | Vacuolar protein sorting 13C protein |
| VIT_02s0012g00270 | 2 | 6027694 | 6106280 | Pleckstriny ( PH ) domain-containing protein |
| VIT_02s0012g00280 | 2 | 6111583 | 6112872 | Senescence-associated protein |
| VIT_02s0012g00290 | 2 | 6133593 | 6135728 | S-ribonuclease binding protein SBP1 |
| VIT_02s0012g00300 | 2 | 6151132 | 6189859 | Unknown protein |
| VIT_02s0012g00310 | 2 | 6204735 | 6223899 | Lon protease |
| VIT_02s0012g00320 | 2 | 6225175 | 6226474 | Norcoclaurine synthase |
| VIT_02s0012g00350 | 2 | 6233046 | 6236366 | Norcoclaurine synthase |
| VIT_02s0012g00360 | 2 | 6238852 | 6242299 | 1-aminocyclopropane-1-carboxylate oxidase |
| VIT_02s0012g00370 | 2 | 6242887 | 6258576 | No hit |
| VIT_02s0012g00380 | 2 | 6273340 | 6285802 | Norcoclaurine synthase |
| VIT_02s0012g00390 | 2 | 6286306 | 6288341 | Norcoclaurine synthase |
| VIT_02s0012g00400 | 2 | 6296189 | 6299377 | 1-aminocyclopropane-1-carboxylate oxidase |
| VIT_02s0012g00410 | 2 | 6305682 | 6310741 | Naringenin, 2-oxoglutarate 3-dioxygenase |
| VIT_02s0012g00420 | 2 | 6311954 | 6315948 | No hit |
| VIT_02s0012g00430 | 2 | 6320484 | 6320851 | Oxidoreductase, 20G-Fe(II) oxygenase |
| VIT_02s0012g00440 | 2 | 6320999 | 6321467 | Norcoclaurine synthase |
| VIT_02s0012g00450 | 2 | 6323850 | 6325555 | 1-aminocyclopropane-1-carboxylate oxidase |
| VIT_02s0012g00460 | 2 | 6332076 | 6381980 | Glycine-rich protein |
| VIT_02s0012g00470 | 2 | 6397883 | 6398726 | Unknown |
| VIT_02s0012g00480 | 2 | 6399524 | 6430074 | Regulator of nonsense transcripts 1 |
| VIT_02s0012g00490 | 2 | 6446158 | 6446783 | Cytochrome c oxidase subunit 6b |
| VIT_02s0012g00500 | 2 | 6449814 | 6450360 | Invertase/pectin methylesterase inhibitor |
| VIT_02s0012g00510 | 2 | 6455739 | 6464806 | Golgi transport Got1 |
| VIT_02s0012g00530 | 2 | 6490169 | 6496086 | Ribose-phosphate pyrophosphokinase 1 |
| VIT_02s0012g00540 | 2 | 6516342 | 6518185 | Peroxidase |
| VIT_02s0012g00550 | 2 | 6518911 | 6526733 | Inositol polyphosphate 5-phosphatase II |
| VIT_02s0012g00560 | 2 | 6533749 | 6545389 | Tobamovirus multiplication 2A TOM2A |
| VIT_02s0012g00570 | 2 | 6554241 | 6560259 | Pseudo-response regulator 2 (APRR2) (TOC2) |
| VIT_02s0012g00580 | 2 | 6586031 | 6594054 | Ankyrin protein kinase |
| VIT_02s0012g00590 | 2 | 6600611 | 6610281 | Unknown protein |


| VIT_02s0012g00600 | 2 | 6611782 | 6615934 | Vacuolar protein sorting 55 |
| :---: | :---: | :---: | :---: | :---: |
| VIT_02s0012g00610 | 2 | 6616375 | 6616852 | Allyl alcohol dehydrogenase |
| VIT_02s0012g00620 | 2 | 6625135 | 6636514 | Phosphoacetylglucosamine mutase |
| VIT_02s0012g00630 | 2 | 6639102 | 6640237 | Myb family |
| VIT_02s0012g00640 | 2 | 6645562 | 6645999 | PBP1 (pinoid-binding protein 1) |
| VIT_02s0012g00650 | 2 | 6653658 | 6654223 | PBP1 (pinoid-binding protein 1) |
| VIT_02s0012g00660 | 2 | 6660229 | 6660825 | Calcium-binding EF-hand |
| VIT_02s0012g00670 | 2 | 6670534 | 6672390 | No hit |
| VIT_02s0012g00680 | 2 | 6682700 | 6682873 | No hit |
| VIT_02s0012g00690 | 2 | 6707170 | 6707803 | No hit |
| VIT_02s0012g00700 | 2 | 6709030 | 6709203 | No hit |
| VIT_02s0012g00710 | 2 | 6733243 | 6741163 | RAB GTPase RAB6 |
| VIT_02s0012g00720 | 2 | 6746761 | 6748648 | Unknown |
| VIT_02s0012g00730 | 2 | 6749467 | 6750865 | Purine permease 10 PUP10 |
| VIT_02s0012g00740 | 2 | 6758931 | 6769661 | Dynamin-like protein 2 b |
| VIT_02s0012g00750 | 2 | 6769877 | 6773816 | Haloacid dehalogenase hydrolase |
| VIT_02s0012g00760 | 2 | 6778127 | 6782078 | Haloacid dehalogenase hydrolase |
| VIT_02s0012g00770 | 2 | 6787173 | 6789662 | No hit |
| VIT_02s0012g00780 | 2 | 6791455 | 6791610 | No hit |
| VIT_02s0012g00790 | 2 | 6792634 | 6793661 | Ras-related protein RAB6A |
| VIT_02s0012g00800 | 2 | 6795685 | 6795849 | No hit |
| VIT_02s0012g00810 | 2 | 6799938 | 6800021 | No hit |
| VIT_02s0012g00820 | 2 | 6804399 | 6806143 | Unknown |
| VIT_02s0012g00830 | 2 | 6817359 | 6819815 | Expansin (VvEXLB1) |
| VIT_02s0012g00840 | 2 | 6820451 | 6823470 | Pentatricopeptide (PPR) repeat-containing protein |
| VIT_02s0012g00850 | 2 | 6823971 | 6851054 | Splicing factor PWI domain-containing protein |
| VIT_02s0012g00860 | 2 | 6855160 | 6859171 | No hit |
| VIT_02s0012g00870 | 2 | 6859219 | 6873440 | RNA-binding protein 10 |
| VIT_02s0012g00880 | 2 | 6897090 | 6898500 | AT-hook DNA-binding protein |
| VIT_02s0012g00890 | 2 | 6914782 | 6915482 | Unknown protein |
| VIT_02s0012g00900 | 2 | 6917525 | 6928933 | AT-hook protein 1 (AHP1) |
| VIT_02s0012g00910 | 2 | 6930227 | 6946661 | Adaptor-related protein complex 2, mu 2 subunit |
| VIT_02s0012g00920 | 2 | 6947431 | 6967357 | Conserved oligomeric complex COG6 |
| VIT_02s0012g00930 | 2 | 6972956 | 6979797 | Unknown protein |
| VIT_02s0012g00940 | 2 | 6980980 | 6981681 | Late embryogenesis abundant group 1 |
| VIT_02s0012g00950 | 2 | 6991276 | 6991983 | H(+)-ATPase 9 AHA9 |


|  | VIT_02s0012g00960 | 2 | 6993519 | 6994847 | 1-deoxy-D-xylulose-5-phosphate synthase |
| :---: | :---: | :---: | :---: | :---: | :---: |
|  | VIT_02s0012g00970 | 2 | 7008648 | 7012385 | Cu2+-exporting ATPase HMA5 (heavy metal ATPase 5) |
|  | VIT_02s0012g00980 | 2 | 7027209 | 7037654 | ferredoxin-6, chloroplast precursor |
|  | VIT_02s0012g00990 | 2 | 7043508 | 7046965 | LOL1 (LSD ONE like 1) |
|  | VIT_02s0012g01000 | 2 | 7050270 | 7060977 | Protein tyrosine phosphatase |
|  | VIT_02s0012g01010 | 2 | 7087110 | 7089452 | Leucine-rich repeat |
|  | VIT_02s0012g01020 | 2 | 7092594 | 7099500 | Zinc finger (C3HC4-type ring finger) |
|  | VIT_02s0012g01030 | 2 | 7100088 | 7118988 | AL-activated malate transporter 1 |
|  | VIT_02s0012g01040 | 2 | 7120118 | 7122681 | NAC domain-containing protein (VvNAC13) |
|  | VIT_02s0012g01050 | 2 | 7131585 | 7133584 | ABC protein 6 non-intrinsic |
|  | VIT_02s0012g01060 | 2 | 7154618 | 7155584 | ABC Transporter (VvNAP3 - VvABCI3) |
| ver_2_3 | VIT_02s0109g00420 | 2 | 13341559 | 13342323 | Disease resistance protein (CC-NBS-LRR class) |
|  | VIT_02s0109g00430 | 2 | 13347364 | 13349570 | Nitrilase 4 (NIT4) |
|  | VIT_02s0109g00440 | 2 | 13353462 | 13353985 | No hit |
|  | VIT_02s0033g00010 | 2 | 13457765 | 13460162 | Unknown protein |
|  | VIT_02s0033g00020 | 2 | 13460163 | 13461470 | Unknown protein |
|  | VIT_02s0033g00030 | 2 | 13506518 | 13511727 | No hit |
|  | VIT_02s0033g00040 | 2 | 13512257 | 13512649 | No hit |
|  | VIT_02s0033g00050 | 2 | 13518893 | 13519179 | Scarecrow transcription factor 3 (SCL3) |
|  | VIT_02s0033g00060 | 2 | 13531787 | 13538078 | DNAJ plastid division protein (ARC6) |
|  | VIT_02s0033g00070 | 2 | 13568564 | 13569348 | No hit |
|  | VIT_02s0033g00080 | 2 | 13590737 | 13637352 | Pleckstriny ( PH ) domain-containing protein |
|  | VIT_02s0033g00090 | 2 | 13639223 | 13640252 | DNA replication licensing factor MCM2 |
|  | VIT_02s0033g00120 | 2 | 13705067 | 13785560 | Pleckstriny ( PH ) domain-containing protein |
|  | VIT_02s0033g00130 | 2 | 13786111 | 13787768 | Cis-zeatin O-beta-D-glucosyltransferase |
|  | VIT_02s0033g00150 | 2 | 13832782 | 13833455 | No hit |
|  | VIT_02s0033g00160 | 2 | 13856081 | 13859536 | Pleckstriny (PH) domain-containing protein |
|  | VIT_02s0033g00180 | 2 | 13880089 | 13881941 | Phenylpropanoid:glucosyltransferase 2 |
|  | VIT_02s0033g00190 | 2 | 13884200 | 13933646 | DNA replication licensing factor MCM2 |
|  | VIT_02s0033g00230 | 2 | 13998724 | 14007202 | Unknown protein |
|  | VIT_02s0033g00240 | 2 | 14031671 | 14033252 | Glucosyltransferase twil |
|  | VIT_02s0033g00250 | 2 | 14051742 | 14052882 | No hit |
|  | VIT_02s0033g00260 | 2 | 14104114 | 14107827 | Pinoresinol-lariciresinol reductase |
|  | VIT_02s0033g00270 | 2 | 14123184 | 14126331 | Pinoresinol-lariciresinol reductase |
|  | VIT_02s0033g00290 | 2 | 14139881 | 14141542 | Pinoresinol-lariciresinol reductase |
|  | VIT_02s0033g00300 | 2 | 14144838 | 14148929 | myb family |


| VIT_02s0033g00310 | 2 | 14158366 | 14159002 |
| :--- | :--- | :--- | :--- |
| VIT_02s0033g00320 | 2 | 14159003 | 14183582 |
| VIT_02s0033g00350 | 2 | 14253249 | 14254161 |
| VIT_02s0033g00360 | 2 | 14265226 | 14266015 |
| VIT_02s0033g00370 | 2 | 14275372 | 14275752 |
| VIT_02s0033g00380 | 2 | 14291438 | 14291683 |
| NIT_02s0033g00390 hit |  |  |  |
| VIT_02s0033g00400 | 2 | 14291727 | 14292732 |
| VIT_02s0033g00410 | 2 | 14351791 | 14352807 |
| VIT_02s0033g00420 | 2 | 14386896 | 14387181 |
| VIT_02s0033g00430 | 2 | 14392775 | 14393842 |$\quad$ No hit


| VIT_02s0033g00750 | 2 | 15496243 | 15499175 | Kinesin motor protein |
| :---: | :---: | :---: | :---: | :---: |
| VIT_02s0033g00770 | 2 | 15560608 | 15562469 | Nitrilase 4B |
| VIT_02s0033g00780 | 2 | 15573189 | 15573552 | 14-3-3 protein GF14 omega (GRF2) |
| VIT_02s0033g00790 | 2 | 15591008 | 15592898 | Nitrilase 4B |
| VIT_02s0033g00800 | 2 | 15606321 | 15608743 | Nitrilase 4 (NIT4) |
| VIT_02s0033g00810 | 2 | 15612721 | 15613040 | No hit |
| VIT_02s0033g00830 | 2 | 15651191 | 15651721 | RAB GTPase RABA4A |
| VIT_02s0033g00840 | 2 | 15659957 | 15669174 | Nitrilase 4B |
| VIT_02s0033g00850 | 2 | 15669871 | 15671771 | Nitrilase |
| VIT_02s0033g00870 | 2 | 15700024 | 15702438 | Nitrilase 4B |
| VIT_02s0033g00880 | 2 | 15714340 | 15714678 | Ribosomal protein L8 (RPL8C) 60S |
| VIT_02s0033g00900 | 2 | 15829983 | 15834399 | Unc51-like kinase |
| VIT_02s0033g00910 | 2 | 15835533 | 15836148 | Kinesin motor protein |
| VIT_02s0033g00920 | 2 | 15921498 | 15922178 | No hit |
| VIT_02s0033g00930 | 2 | 15941055 | 15952365 | Single-strand DNA binding protein |
| VIT_02s0033g00960 | 2 | 15974263 | 15978140 | Zinc-binding protein |
| VIT_02s0033g00970 | 2 | 16053537 | 16062151 | No hit |
| VIT_02s0033g00980 | 2 | 16081574 | 16084857 | NADH-plastoquinone oxidoreductase subunit 2 |
| VIT_02s0033g00990 | 2 | 16084858 | 16085540 | Ribosomal protein S7 30S |
| VIT_02s0033g01000 | 2 | 16093114 | 16095441 | Anthraniloyal-CoA: methanol anthraniloyal transferase |
| VIT_02s0033g01010 | 2 | 16152695 | 16154133 | Ribosomal protein S7 30S |
| VIT_02s0033g01020 | 2 | 16190948 | 16193334 | Anthraniloyal-CoA: methanol anthraniloyal transferase |
| VIT_02s0033g01030 | 2 | 16252684 | 16254941 | Anthraniloyal-CoA: methanol anthraniloyal transferase |
| VIT_02s0033g01050 | 2 | 16299329 | 16301080 | Anthraniloyal-CoA: methanol anthraniloyal transferase |
| VIT_02s0033g01060 | 2 | 16349045 | 16351278 | Anthraniloyal-CoA: methanol anthraniloyal transferase |
| VIT_02s0033g01070 | 2 | 16356696 | 16359431 | Anthraniloyal-CoA: methanol anthraniloyal transferase |
| VIT_02s0033g01100 | 2 | 16435669 | 16439357 | No hit |
| VIT_02s0033g01110 | 2 | 16439358 | 16447346 | Male germ cell-associated kinase |
| VIT_02s0033g01120 | 2 | 16515095 | 16520470 | Dehydration-responsive protein |
| VIT_02s0033g01130 | 2 | 16555004 | 16556115 | SAR1 (secretion-associated ras) |
| VIT_02s0033g01150 | 2 | 16602266 | 16611445 | Unknown protein |
| VIT_02s0033g01160 | 2 | 16649637 | 16654775 | Unknown protein |
| VIT 02s0033g01170 | 2 | 16657559 | 16664675 | Replication protein RPA 70kDa subunit |


| Meta-QTL | Gene ID | Chr | $\begin{aligned} & \hline \begin{array}{l} \text { Start } \\ \text { (bp) } \end{array} \\ & \hline \end{aligned}$ | End(bp) | Annotation_V1 |
| :---: | :---: | :---: | :---: | :---: | :---: |
| pheno_3_1 | VIT_03s0038g00620 | 3 | 547345 | 562437 | Zinc knuckle |
|  | VIT_03s0038g00630 | 3 | 565597 | 568081 | Endo-1,4-beta-glucanase |
|  | VIT_03s0038g00640 | 3 | 574493 | 578920 | Unknown |
|  | VIT_03s0038g00650 | 3 | 579366 | 583281 | Coenzyme Q10 homolog B |
|  | VIT_03s0038g00660 | 3 | 584427 | 587912 | Unknown protein |
|  | VIT_03s0038g00670 | 3 | 592539 | 594503 | fructose-bisphosphate aldolase, chloroplast precursor |
|  | VIT_03s0038g00690 | 3 | 594846 | 598517 | Zinc finger (B-box type) |
|  | VIT_03s0038g00700 | 3 | 606114 | 613813 | Kinesin motor protein |
|  | VIT_03s0038g00710 | 3 | 614468 | 614930 | Unknown |
|  | VIT_03s0038g00720 | 3 | 615777 | 618174 | NADH dehydrogenase (ubiquinone) 1 beta subcomplex 9 |
|  | VIT_03s0038g00730 | 3 | 623867 | 624348 | No hit |
|  | VIT_03s0038g00740 | 3 | 624648 | 632238 | GC1 (GIANT chloroplast 1) |
|  | VIT_03s0038g00750 | 3 | 632713 | 639536 | Ubiquitin fusion degradation protein UFD1 |
|  | VIT_03s0038g00760 | 3 | 644491 | 647420 | Arginine decarboxylase (Fragment) |
|  | VIT_03s0038g00790 | 3 | 662174 | 664796 | V-type H+-transporting ATPase 16kDa proteolipid subunit |
|  | VIT_03s0038g00800 | 3 | 666647 | 670774 | NADP-dependent D-sorbitol-6-phosphate dehydrogenase |
|  | VIT_03s0038g00820 | 3 | 670775 | 676616 | 3-oxo-5-alpha-steroid 4-dehydrogenase, C-terminal |
|  | VIT_03s0038g00840 | 3 | 679207 | 684375 | BPC4/BBR/BPC4/BPC4 |
|  | VIT_03s0038g00860 | 3 | 689247 | 693308 | Basic Leucine Zipper Transcription Factor (VvbZIP05) |
|  | VIT_03s0038g00870 | 3 | 694566 | 700828 | Dihydrouridine synthase 3 |
|  | VIT_03s0038g00880 | 3 | 702486 | 704460 | Amidophosphoribosyltransferase 2 |
|  | VIT_03s0038g00890 | 3 | 709401 | 710989 | No hit |
|  | VIT_03s0038g00910 | 3 | 713857 | 721463 | Ribosome-binding factor A , chloroplast precursor |
|  | VIT_03s0038g00920 | 3 | 722303 | 732914 | Carbohydrate kinase, PfkB |
|  | VIT_03s0038g00930 | 3 | 736435 | 737319 | Auxin responsive SAUR protein |
|  | VIT_03s0038g00940 | 3 | 738127 | 738887 | Auxin-responsive |
|  | VIT_03s0038g00950 | 3 | 749716 | 750398 | Auxin-responsive SAUR9 |
|  | VIT_03s0038g00960 | 3 | 755323 | 760266 | Transcription factor TFIIF |
|  | VIT_03s0038g00980 | 3 | 783897 | 784664 | Unknown |
|  | VIT_03s0038g01010 | 3 | 799108 | 799188 | No hit |


| VIT_03s0038g01020 | 3 | 815278 | 815673 |
| :--- | :--- | :--- | :--- |
| VIT_03s0038g01060 | 3 | 854295 | 855373 |
| VIT_03s0038g01070 | 3 | 856872 | 858394 |
| VIT_03s0038g01080 | 3 | 858886 | 859170 |
| VIT_03s0038g01090 | 3 | 862995 | 863398 |
| VIT_03s0038g01100 | 3 | 864608 | 864876 |
| VIT_03s0038g01110 | 3 | 866357 | 866897 |
| VIT_03s0038g01120 | 3 | 867890 | 868156 |
| VIT_03s0038g01130 | 3 | 869586 | 872380 |
| VIT_03s0038g01140 | 3 | 876348 | 878337 |
| VIT_03s0038g01150 | 3 | 880653 | 881315 |
| VIT_03s0038g01160 | 3 | 883689 | 883976 |
| VIT_03s0038g01170 | 3 | 885397 | 885684 |
| VIT_03s0038g01180 | 3 | 888635 | 888955 |
| VIT_03s0038g01190 | 3 | 892009 | 892382 |
| VIT_03s0038g01210 | 3 | 894854 | 895141 |
| VIT_03s0038g01220 | 3 | 897891 | 898180 |
| VIT_03s0038g01230 | 3 | 900918 | 901205 |
| VIT_03s0038g01250 | 3 | 903282 | 903569 |
| VIT_03s0038g01260 | 3 | 905892 | 906181 |
| VIT_03s0038g01270 | 3 | 908817 | 909399 |
| VIT_03s0038g01280 | 3 | 910867 | 911175 |
| VIT_03s0038g01290 | 3 | 915230 | 915499 |
| VIT_03s0038g01300 | 3 | 917424 | 917748 |
| VIT_03s0038g01310 | 3 | 921733 | 927965 |
| VIT_03s0038g01320 | 3 | 930985 | 948815 |
| VIT_03s0038g01330 | 3 | 955035 | 956506 |
| VIT_03s0038g01360 | 3 | 964891 | 966169 |
| VIT_03s0038g01370 | 3 | 967748 | 969830 |
| VIT_03s0038g01380 | 3 | 971266 | 975571 |
| VIT_03s0038g01390 | 3 | 976131 | 977121 |
| VIT_03s0038g01400 | 3 | 978958 | 982622 |
| VIT_03s0038g01410 | 3 | 984045 | 985466 |
| VIT_03s0038g01420 | 3 | 991431 | 993320 |
| VIT_03s0038g01430 | 3 | 993344 | 994100 |
| VIT_03s0038g01440 | 3 | 996232 | 1004637 |
|  |  |  |  |

Metalloprotease
Metalloprotease
Pentatricopeptide (PPR) repeat-containing protein Auxin responsive SAUR protein Auxin responsive SAUR protein Auxin responsive SAUR protein

Auxin-responsive SAUR31
Auxin responsive SAUR protein
Auxin-responsive
Metalloendoproteinase 1 precursor
Auxin-responsive
Auxin responsive SAUR protein Auxin responsive SAUR protein Auxin responsive SAUR protein

Auxin-induced SAUR
Auxin responsive SAUR protein
Auxin-induced protein 15A
Auxin responsive SAUR protein Auxin responsive SAUR protein Auxin responsive SAUR protein Auxin responsive SAUR protein Auxin responsive SAUR protein Auxin responsive SAUR protein

Auxin-induced SAUR
Auxin responsive SAUR protein
Crp1 protein
Anthranilate N -benzoyltransferase protein 1
Unknown
ABA-responsive protein (HVA22)HVA22H
Calcium-binding EF hand Aquaporin TIP3;1
R protein PRF disease resistance protein
Aquaporin PIP PIP1A
Phytochelatin synthetase
Phytochelatin synthetase
Adenosylhomocysteine nucleosidase.

| VIT_03s0038g01450 | 3 | 1004638 | 1005093 |
| :--- | :--- | :--- | :--- |
| VIT_03s0038g01460 | 3 | 1006320 | 1009717 |
| VIT_03s0038g01470 | 3 | 1009718 | 1014601 |
| VIT_03s0038g01480 | 3 | 1015472 | 1018780 |
| VIT_03s0038g01490 | 3 | 1025473 | 1040019 |
| VIT_03s0038g01510 | 3 | 1041388 | 1041967 |
| VIT_03s0038g01520 | 3 | 1044307 | 1047456 |
| VIT_03s0038g01530 | 3 | 1050448 | 1058022 |
| VIT_03s0038g01540 | 3 | 1062491 | 1063732 |
| VIT_03s0038g01550 | 3 | 1066258 | 1068897 |
| VIT_03s0038g01580 | 3 | 1077896 | 1078534 |
| VIT_03s0038g01590 | 3 | 1082216 | 1083954 |
| VIT_03s0038g01610 | 3 | 1084197 | 1085815 |
| VIT_03s0038g01620 | 3 | 1088445 | 1096163 |
| VIT_03s0038g01630 | 3 | 1111137 | 1113800 |
| VIT_03s0038g01670 | 3 | 1168819 | 1177208 |
| VIT_03s0038g01740 | 3 | 1215576 | 1222056 |
| VIT_03s0038g01750 | 3 | 1224505 | 1226936 |
| VIT_03s0038g01760 | 3 | 1231040 | 1235642 |
| VIT_03s0038g01770 | 3 | 1235713 | 1237448 |
| VIT_03s0038g01780 | 3 | 1238202 | 1248527 |
| VIT_03s0038g01790 | 3 | 1252286 | 1255340 |
| VIT_03s0038g01810 | 3 | 1263090 | 1274110 |
| VIT_03s0038g01820 | 3 | 1275024 | 1279206 |
| VIT_03s0038g01830 | 3 | 1283253 | 1285733 |
| VIT_03s0038g01840 | 3 | 1286455 | 1288037 |
| VIT_03s0038g01850 | 3 | 1298173 | 1299515 |
| VIT_03s0038g01860 | 3 | 1301501 | 1302939 |
| VIT_03s0038g01870 | 3 | 1303910 | 1304387 |
| VIT_03s0038g01880 | 3 | 1313644 | 1323526 |
| VIT_03s0038g01920 | 3 | 1332326 | 1347193 |
| VIT_03s0038g01930 | 3 | 1347905 | 1348611 |
| VIT_03s0038g01940 | 3 | 1349912 | 1356622 |
| VIT_03s0038g01950 | 3 | 1357739 | 1360559 |
| VIT_03s0038g01960 | 3 | 1362167 | 1363750 |
| VIT_03s0038g01970 | 3 | 1367294 | 1369340 |
|  |  |  |  |


Translocon-associated protein alpha TRAP complex
Proline-rich protein 4
Proline-rich protein 4
Proline-rich protein 4
Proline-rich protein 4
Nuclear pore complex protein Nup188 Nuclear pore complex protein Nup 188 Nuclear pore complex protein Nup 188
Peptidyl-prolyl cis-trans isomerase ROC5 (rotamase CYP 5) Permease nonimprinted in Prader-Willi/Angelman

Glycine-rich protein
Proline-rich family protein
Amidase
$\left.\begin{array}{llll}\text { VIT_03s0038g01990 } & 3 & 1374043 & 1381233 \\ \text { VIT_03s0038g02000 } & 3 & 1383516 & 1393182 \\ \text { VIT_03s0038g02010 } & 3 & 1393183 & 1395922 \\ \text { VIT_03s0038g02020 } & 3 & 1395923 & 1400744 \\ \text { VIT_03s0038g02030 } & 3 & 1402181 & 1405517 \\ \text { VIT_03s0038g02040 } & 3 & 1406571 & 1408542 \\ \text { VIT_03s0038g02050 } & 3 & 1412118 & 1412819 \\ \text { VIT_03s0038g02060 } & 3 & 1415776 & 1416475 \\ \text { VIT_03s0038g02070 } & 3 & 1424949 & 1429038 \\ \text { VIT_03s0038g02080 } & 3 & 1436852 & 1438855 \\ \text { AIT_03s0038g02090 } & 3 & 1441262 & 1442535 \\ \text { Amidase } \\ \text { VIT_03s0038g02100 } & 3 & 1443063 & 1447359 \\ \text { VIT_03s0038g02110 } & 3 & 1449330 & 1451521\end{array}\right]$ Amidase
 VIT 03s0180g00050 $3 \quad 5981525 \quad 5989159$ VIT_03s0180g00060 $3 \quad 5989504 \quad 599394$ VIT 03s0180g00070 $3 \quad 5995174 \quad 6000876$ VIT 03s0180g00080 $3 \quad 6009317 \quad 6020720$ VIT_-03s0180g00090 $\quad 3 \quad 6023040 \quad 6024531$ VIT_03s0180g00100 $\quad 3 \quad 6030251 \quad 6032840$ VIT_03s0180g00110 360337236036694 VIT 03s0180g00120 $366037870 \quad 6059709$ VIT_03s0180g00130 $3 \quad 6060227 \quad 6064528$ VIT_03s0180g00140 $3 \quad 6071530 \quad 6072486$ VIT 03s0180g00150 $3 \quad 6078670 \quad 6081676$ VIT_-03s0180g00160 $\quad 3 \quad 6090668 \quad 6100225$ VIT_03s0180g00170 $\quad 3 \quad 6101310 \quad 6105927$ VIT_03s0180g00180 3661066026118797 VIT 03s0180g00190 $3 \quad 6119889 \quad 612437$ VIT_03s0180g00200 $\quad 3 \quad 6124465 \quad 6126202$ VIT_03s0180g00210 $\quad 3 \quad 6147352 \quad 6148726$ VIT_03s0180g00230 $3 \quad 6166608 \quad 6177981$ VIT ${ }^{-} 03 \mathrm{~s} 0180 \mathrm{~g} 00240 \quad 3 \quad 6179625 \quad 6179759$ VIT_03s0180g00250 $3 \quad 6179858 \quad 6181360$ VIT_03s0180g00260 3662123296214039 VIT 03s $0180 \mathrm{~g} 00270 \quad 3 \quad 6217552 \quad 6226884$ VIT_03s0180g00280 $\quad 3 \quad 6255489 \quad 6257208$ VIT_03s0180g00290 $366263737 \quad 6264635$ VIT_03s0180g00300 $366267395 \quad 6267971$ VIT_03s0180g00310 $3 \quad 6269156 \quad 6269856$ VIT_03s0180g00320 $\quad 3 \quad 6281986 \quad 6283705$ VIT_03s0091g00010 $366310807 \quad 6311602$ $\begin{array}{lllll}\text { VIT 03s0091g00020 } & 3 & 6314436 & 6314606\end{array}$ VIT_03s0091g00030 $\quad 3 \quad 6314607 \quad 6315780$ VIT_03s0091g00040 $\quad 3 \quad 6338109 \quad 6339850$ VIT_03s0091g00050 $36357496 \quad 6364352$

Cysteine proteinase
Phosphoric monoester hydrolase
Calcium-binding protein
Cyclin D3_2
DNA-directed RNA polymerase II subunit C
Late embryonic abundant protein EMB7
Cyclase
Cyclase
Receptor kinase
Unknown protein
Stress enhanced protein 1 (SEP1)
Sodium hydrogen antiporter NHD1
D-3-phosphoglycerate dehydrogenase, chloroplast precursor
Acetyl xylan esterase AxeA
Receptor protein kinase
ATTIC21/CIA5/TIC21 (chloroplast import apparatus 5) Reticulon-like protein B11 RTNLB11

Unknown protein
No hit
Limonoid UDP-glucosyltransferase (VvGT2)
Myb domain protein R1
UNC-50
No hit
Cinnamyl alcohol dehydrogenase
Cinnamyl alcohol dehydrogenase
Pseudouridine synthase
Indole-3-acetate beta-glucosyltransferase Gibberellin-regulated protein 4 (GASA4)

ABC transporter F member 2 ATP-dependent Clp protease ClpB protein indole-3-acetate beta-glucosyltransferase (VvGT3)

No hit
No hit
No hit
Limonoid UDP-glucosyltransferase (VvGT1) Triacylglycerol/steryl ester hydrolase

| VIT_03s0091g00060 | 3 | 6374562 | 6374820 |
| :--- | :--- | :--- | :--- |
| VIT_03s0091g00070 | 3 | 6374821 | 6376254 |
| VIT_03s0091g00080 | 3 | 6384738 | 6388178 |
| VIT_03s0091g00090 | 3 | 6388280 | 6390379 |
| VIT_03s0091g00100 | 3 | 6391617 | 6395252 |
| VIT_03s0091g00110 | 3 | 6395575 | 6397456 |
| VIT_03s0091g00120 | 3 | 6398105 | 6398826 |
| VIT_03s0091g00130 | 3 | 6399870 | 6400567 |
| VIT_03s0091g00140 | 3 | 6404905 | 6405345 |
| VIT_03s0091g00150 | 3 | 6405346 | 6405711 |
| VIT_03s0091g00160 | 3 | 6406167 | 6406991 |
| VIT_03s0091g00180 | 3 | 6419877 | 6420461 |
| VIT_03s0091g00190 | 3 | 6447349 | 6451553 |
| VIT_03s0091g00200 | 3 | 6479048 | 6503929 |
| VIT_03s0091g00210 | 3 | 6507392 | 6509263 |
| VIT_03s0091g00220 | 3 | 6511172 | 6511647 |
| VIT_03s0091g00230 | 3 | 6512832 | 6514492 |
| VIT_03s0091g00240 | 3 | 6521904 | 6537905 |
| VIT_03s0091g00250 | 3 | 6538966 | 6539935 |
| VIT_03s0091g00260 | 3 | 6548677 | 6549577 |
| VIT_03s0091g00270 | 3 | 6551840 | 6559848 |
| VIT_03s0091g00280 | 3 | 6566136 | 6569256 |
| VIT_03s0091g00290 | 3 | 6571900 | 6589424 |
| VIT_03s0091g00300 | 3 | 6623124 | 6623416 |
| VIT_03s0091g00310 | 3 | 6633708 | 6635996 |
| VIT_03s0091g00320 | 3 | 6642216 | 6651075 |
| VIT_03s0091g00350 | 3 | 6664651 | 6666079 |
| VIT_03s0091g00360 | 3 | 6668161 | 6688642 |
| VIT_03s0091g00370 | 3 | 6691248 | 6693137 |
| VIT_03s0091g00380 | 3 | 6699445 | 6705656 |
| VIT_03s0091g00390 | 3 | 6715491 | 6716068 |
| VIT_03s0091g00400 | 3 | 6717421 | 6752391 |
| VIT_03s0091g00410 | 3 | 6773316 | 6776359 |
| VIT_03s0091g00420 | 3 | 6785458 | 6787059 |
| VIT_03s0091g00430 | 3 | 6798217 | 6802152 |
| VIT_03s0091g00440 | 3 | 6804406 | 6807382 |
|  |  |  |  |

ACT domain-containing protein (ACR6)
ACT domain-containing protein (ACR6)
Methylthioribose kinase
No hit
Methylthioribose kinase
Plant basic secretory protein (BSP) family Prp27-like protein

Prp27
Plant basic secretory protein (BSP) family NtPRp27 secretory protein NtPRp27 secretory protein

Unknown
WD40
DNA repair protein REV1
Ethylene-responsive protein Unknown protein Unknown protein
Haloacid dehalogenase hydrolase
Unknown
Zinc finger protein 4 TIP41
Cyclin A3;4
Pathogenesis-related homeodomain protein (PRHA) B-cell receptor-associated protein 31
Indole-3-acetic acid-amido synthetase GH3.8
Ribosomal rna assembly protein mis3
No hit
Seed maturation protein PM23
Pentatricopeptide (PPR) repeat-containing protein
Thioredoxin domain-containing protein 9
Snakin-1
Transducin family protein / WD-40 repeat
Unknown protein
No hit
SWIB complex BAF60b domain-containing protein
Pentatricopeptide (PPR) repeat-containing protein

| VIT_03s0091g00450 | 3 | 6811798 | 6813280 |
| :--- | :--- | :--- | :--- |
| VIT_03s0091g00460 | 3 | 6817116 | 6818768 |
| VIT_03s0091g00470 | 3 | 6846055 | 6849796 |
| VIT_03s0091g00480 | 3 | 6851584 | 6855075 |
| VIT_03s0091g00490 | 3 | 6859815 | 6860409 |
| VIT_03s0091g00500 | 3 | 6871858 | 6872341 |
| VIT_03s0091g00510 | 3 | 6894959 | 6895595 |
| VIT_03s0091g00520 | 3 | 6905764 | 6914781 |
| VIT_03s0091g00530 | 3 | 6917800 | 6921320 |
| VIT_03s0091g00540 | 3 | 6976055 | 6984113 |
| VIT_03s0091g00550 | 3 | 6987237 | 6990594 |
| VIT_03s0091g00560 | 3 | 6995098 | 6998522 |
| VIT_03s0091g00570 | 3 | 6998808 | 6999512 |
| VIT_03s0091g00580 | 3 | 7035451 | 7039010 |
| VIT_03s0091g00590 | 3 | 7040488 | 7042974 |
| VIT_03s0091g00600 | 3 | 7044909 | 7047565 |
| VIT_03s0091g00610 | 3 | 7050934 | 7055804 |
| VIT_03s0091g00620 | 3 | 7055805 | 7059068 |
| VIT_03s0091g00630 | 3 | 7065044 | 7065631 |
| VIT_03s0091g00640 | 3 | 7072474 | 7073358 |
| VIT_03s0091g00650 | 3 | 7076200 | 7076677 |
| VIT_03s0091g00660 | 3 | 7082071 | 7098207 |
| VIT_03s0091g00670 | 3 | 7099015 | 7099742 |
| VIT_03s0091g00680 | 3 | 7103281 | 7113666 |
| VIT_03s0091g00690 | 3 | 7125371 | 7126295 |
| VIT_03s0091g00700 | 3 | 7137444 | 7143607 |
| VIT_03s0091g00710 | 3 | 7149908 | 7160638 |
| VIT_03s0091g00720 | 3 | 7168741 | 7177092 |
| VIT_03s0091g00730 | 3 | 7192504 | 7193911 |
| VIT_03s0091g00740 | 3 | 7195947 | 7206249 |
| VIT_03s0091g00750 | 3 | 7219508 | 7223633 |
| VIT_03s0091g00760 | 3 | 7223730 | 7230484 |
| VIT_03s0091g00770 | 3 | 7244888 | 7245217 |
| VIT_03s0091g00810 | 3 | 7287360 | 7289100 |
| VIT_03s0091g00820 | 3 | 7291711 | 7302903 |
| VIT_03s0091g00830 | 3 | 7303408 | 7311191 |
|  |  |  |  |


| VIT_03s0091g00840 | 3 | 7321433 | 7322062 |
| :--- | :--- | :--- | :--- |
| VIT_03s0091g00850 | 3 | 7323004 | 7324112 |
| VIT_03s0091g00860 | 3 | 7324113 | 7339946 |
| VIT_03s0091g00870 | 3 | 7342165 | 7357074 |
| VIT_03s0091g00880 | 3 | 7374686 | 7382483 |
| VIT_03s0091g00890 | 3 | 7385745 | 7389061 |
| VIT_03s0091g00900 | 3 | 7389633 | 7402240 |
| VIT_03s0091g00920 | 3 | 7415296 | 7415826 |
| VIT_03s0091g00930 | 3 | 7417454 | 7418376 |
| VIT_03s0091g00940 | 3 | 7428322 | 7430945 |
| VIT_03s0091g00950 | 3 | 7441440 | 7444729 |
| VIT_03s0091g00960 | 3 | 7446465 | 7449614 |
| VIT_03s0091g00970 | 3 | 7469828 | 7484328 |
| VIT_03s0091g00990 | 3 | 7502149 | 7502253 |
| VIT_03s0091g01010 | 3 | 7535063 | 7536574 |
| VIT_03s0091g01020 | 3 | 7537306 | 7559608 |
| VIT_03s0091g01030 | 3 | 7563890 | 7582120 |
| VIT_03s0091g01040 | 3 | 7603310 | 7604326 |
| VIT_03s0091g01050 | 3 | 7652632 | 7661288 |
| VIT_03s0091g01060 | 3 | 7673917 | 7675754 |
| VIT_03s0091g01080 | 3 | 7693399 | 7695038 |
| VIT_03s0091g01090 | 3 | 7704746 | 7705934 |
| VIT_03s0091g01100 | 3 | 7710736 | 7711700 |
| VIT_03s0091g01110 | 3 | 7716202 | 7726782 |
| VIT_03s0091g01120 | 3 | 7739421 | 7740245 |
| VIT_03s0091g01130 | 3 | 7741682 | 7743818 |
| VIT_03s0091g01140 | 3 | 7747402 | 7749276 |
| VIT_03s0091g01150 | 3 | 7759653 | 7762665 |
| VIT_03s0091g01160 | 3 | 7771297 | 7771879 |
| VIT_03s0091g01170 | 3 | 7813667 | 7815342 |
| VIT_03s0091g01180 | 3 | 7826669 | 7826884 |
| VIT_03s0091g01190 | 3 | 7831919 | 7853120 |
| VIT_03s0091g01200 | 3 | 7855565 | 7861870 |
| VIT_03s0091g01220 | 3 | 7888986 | 7890085 |
| VIT_03s0091g01230 | 3 | 7890272 | 7892417 |
| VIT_03s0091g01240 | 3 | 7903958 | 7907570 |
|  |  |  |  |
| VI |  |  |  |

DNA polymerase kappa subunit Adenylylsulfate kinase 1 (AKN1)

Endoxylanase
Endoxylanase
Endoxylanase
Zfwd2 protein (ZFWD2)
GRAM domain-containing protein / ABA-responsive
Zfwd2 protein (ZFWD2)
Endoxylanase
No hit
ADP-ribosylation factor-like A1D
Heat shock protein 70
LIM domain protein WLIM1
Unknown protein
Telomere repeat binding factor 1

## No hit

Nucleobase-ascorbate transporter 4 (NAT4)
Cyclin delta-2
-aminocyclopropane-1-carboxylate oxidase
Unknown
Unknown protein
Meprin and TRAF homology domain-containing protein Meprin and TRAF homology domain-containing protein Meprin and TRAF homology domain-containing protein Meprin and TRAF homology domain-containing protein

Cleavage and polyadenylation specificity factor
No hit
Meprin and TRAF homology domain-containing protein
No hit
Magnesium transporter CorA
Sinapoylglucose:malate sinapoyltransferase (SNG1) Enhancer of mRNA-decapping protein 4
Enhancer of mRNA-decapping protein 4
Serine carboxypeptidase S10 / Anthocyanin Acyl-transferase

| VIT_03s0091g01260 | 3 | 7918875 | 7934080 |
| :--- | :--- | :--- | :--- |
| VIT_03s0091g01270 | 3 | 7935398 | 7938767 |
| VIT_03s0091g01280 | 3 | 7942831 | 7944961 |
| VIT_03s0091g01290 | 3 | 7951175 | 7955690 |
| VIT_03s0088g00010 | 3 | 7987351 | 7989378 |
| VIT_03s0088g00030 | 3 | 8005949 | 8006486 |
| VIT_03s0088g00040 | 3 | 8018981 | 8020373 |
| VIT_03s0088g00050 | 3 | 8035215 | 8039037 |
| VIT_03s0088g00060 | 3 | 8069862 | 8072808 |
| VIT_03s0088g00070 | 3 | 8084352 | 8086688 |
| VIT_03s0088g00100 | 3 | 8122234 | 8124112 |
| VIT_03s0088g00110 | 3 | 8141417 | 8145242 |
| VIT_03s0088g00140 | 3 | 8186019 | 8188472 |
| VIT_03s0088g00150 | 3 | 8193838 | 8194272 |
| VIT_03s0088g00160 | 3 | 8195668 | 8197771 |
| VIT_03s0088g00170 | 3 | 8209417 | 8210614 |
| VIT_03s0088g00180 | 3 | 8212188 | 8213029 |
| VIT_03s0088g00190 | 3 | 8220525 | 8221777 |
| VIT_03s0088g00200 | 3 | 8228928 | 8232449 |
| VIT_03s0088g00210 | 3 | 8232628 | 8239401 |
| VIT_03s0088g00220 | 3 | 8240353 | 8241018 |
| VIT_03s0088g00230 | 3 | 8241019 | 8242075 |
| VIT_03s0088g00240 | 3 | 8242076 | 8244718 |
| VIT_03s0088g00250 | 3 | 8244719 | 8251952 |
| VIT_03s0088g00260 | 3 | 8252549 | 8257129 |
| VIT_03s0088g00270 | 3 | 8267214 | 8267760 |
| VIT_03s0088g00280 | 3 | 8281268 | 8302843 |
| VIT_03s0088g00290 | 3 | 8315170 | 8315924 |
| VIT_03s0088g00300 | 3 | 8320984 | 8322831 |
| VIT_03s0088g00310 | 3 | 8329921 | 8335645 |
| VIT_03s0088g00320 | 3 | 8343365 | 8347681 |
| VIT_03s0088g00330 | 3 | 8350297 | 8352177 |
| VIT_03s0088g00340 | 3 | 8371381 | 8372242 |
| VIT_03s0088g00350 | 3 | 8374871 | 8376359 |
| VIT_03s0088g00370 | 3 | 8403058 | 8408371 |
| VIT_03s0088g00380 | 3 | 8412010 | 8414515 |
|  |  |  |  |

sinapoylglucose-choline O-sinapoyltransferase
No hit
Serine carboxypeptidase S10
Concanavalin A lectin
Serine carboxypeptidase-like 6 Concanavalin A lectin
Serine carboxypeptitase 1 Isoflavone reductase

No hit
Concanavalin A lectin
Serine carboxypeptidase SCPL17
Isoflavone reductase protein 2 Serine carboxypeptidase SCPL17 Serine carboxypeptidase SCPL17 Brassinosteroid insensitive 1-associated receptor kinase 1 No hit
Brassinosteroid insensitive 1-associated receptor kinase 1
Unknown
STT3B (staurosporin and temperature sensitive 3-like B)
Replication factor A 1, rfa1
Replication protein A 70 kDa DNA-binding subunit
Replication protein A 70b
Isoflavone reductase
Serine carboxypeptidase S10
No hit
Ribosomal RNA small subunit methyltransferase E
Phytosulfokines PSK2
Unknown
No hit
Peptidase M50 F-box protein 7

No hit
Unknown
Unknown protein
Tubulin alpha

| VIT_03s0088g00390 | 3 | 8414516 | 8423780 |
| :--- | :--- | :--- | :--- |
| VIT_03s0088g00400 | 3 | 8427000 | 8430780 |
| VIT_03s0088g00410 | 3 | 8432142 | 8436432 |
| VIT_03s0088g00420 | 3 | 8443429 | 8445079 |
| VIT_03s0088g00450 | 3 | 8486720 | 8618518 |
| VIT_03s0088g00460 | 3 | 8630875 | 8631169 |
| VIT_03s0088g00470 | 3 | 8631326 | 8631646 |
| VIT_03s0088g00490 | 3 | 8653859 | 8658460 |
| VIT_03s0088g00500 | 3 | 8660759 | 8679489 |
| VIT_03s0088g00510 | 3 | 8696026 | 8697570 |
| VIT_03s0088g00540 | 3 | 8736929 | 8743307 |
| VIT_03s0088g00550 | 3 | 8744237 | 8744878 |
| VIT_03s0088g00560 | 3 | 8763759 | 8765911 |
| VIT_03s0088g00570 | 3 | 8766129 | 8766881 |
| VIT_03s0088g00590 | 3 | 8776002 | 8776646 |
| VIT_03s0088g00600 | 3 | 8793246 | 8793890 |
| VIT_03s0088g00610 | 3 | 8820035 | 8820694 |
| VIT_03s0088g00620 | 3 | 8825504 | 8827218 |
| VIT_03s0088g00630 | 3 | 8828264 | 8830122 |
| VIT_03s0088g00650 | 3 | 8842942 | 8844322 |
| VIT_03s0088g00680 | 3 | 8883856 | 8884465 |
| VIT_03s0088g00690 | 3 | 8895244 | 8909155 |
| VIT_03s0088g00700 | 3 | 8911524 | 8912111 |
| VIT_03s0088g00710 | 3 | 8916223 | 8916893 |
| VIT_03s0088g00720 | 3 | 8923838 | 8924209 |
| VIT_03s0088g00730 | 3 | 8927274 | 8927983 |
| VIT_03s0088g00750 | 3 | 8949588 | 8950233 |
| VIT_03s0088g00760 | 3 | 8955534 | 8962261 |
| VIT_03s0088g00780 | 3 | 8986184 | 8986955 |
| VIT_03s0088g00810 | 3 | 9044529 | 9045211 |
| VIT_03s0088g00820 | 3 | 9050245 | 9055257 |
| VIT_03s0088g00840 | 3 | 9072903 | 9073766 |
| VIT_03s0088g00860 | 3 | 9080159 | 9081145 |
| VIT_03s0088g00880 | 3 | 9083004 | 9083780 |
| VIT_03s0088g00890 | 3 | 9101007 | 9101495 |
| VIT_03s0088g00900 | 3 | 9106967 | 9110222 |
|  |  |  |  |

DnaJ homolog, subfamily C, member 17
Tubulin alpha-6 chain
Pyruvate kinase isozyme A, chloroplast precursor Gag-pol polyprotein FKBP12-rapamycin complex-associated protein

Angustifolia
Auxin responsive SAUR protein
Unknown
TRNA synthetase class II (D, K and N)
hypothetical MADS-box type 1 alpha 1b (VviMADS1A1b)
No hit
putative MADS-box type 1 alpha 1a (VviMADS1A1a) Citrate synthase
Glutamine synthetase B1 GLB1
hypothetical MADS-box type 1 alpha 1 g (VviMADS1A1g) putative MADS-box type 1 alpha 1c (VviMADS1A1c) hypothetical MADS-box type 1 alpha 1 f (VviMADS1A1f)

Stress-induced
Unknown protein
Xyloglucan:xyloglucosyl transferase Pathogenesis-related protein 1 precursor (PRP 1) Pathogenesis-related protein 1 precursor (PRP 1) Pathogenesis related protein 1 precursor (pr1 gene) Pathogenesis-related protein 1 precursor (PRP 1) Pathogenesis-related protein 1 precursor (PRP 1)

Methionine sulfoxide reductase Pathogenesis related protein 1 precursor [Vitis vinifera]

No hit
Pathogenesis-related protein 1 precursor (PRP 1) Pathogenesis-related protein 1 precursor (PRP 1)

Unknown
No hit
Lectin-like receptor kinase Kinase
Pathogenesis related protein 1 precursor [Vitis vinifera]
Pathogenesis-related protein 1B

| VIT_03s0088g00910 | 3 | 9119915 | 9130668 |
| :--- | :--- | :--- | :--- |
| VIT_03s0088g00920 | 3 | 9145287 | 9145858 |
| VIT_03s0088g00930 | 3 | 9146189 | 9147403 |
| VIT_03s0088g00940 | 3 | 9150089 | 9150575 |
| VIT_03s0088g00950 | 3 | 9153293 | 9162432 |
| VIT_03s0088g00960 | 3 | 9168413 | 9182543 |
| VIT_03s0088g00970 | 3 | 9190121 | 9193027 |
| VIT_03s0088g00980 | 3 | 9205881 | 9213548 |
| VIT_03s0088g00990 | 3 | 9216064 | 9223316 |
| VIT_03s0088g01000 | 3 | 9223670 | 9231652 |
| VIT_03s0088g01030 | 3 | 9249737 | 9251605 |
| VIT_03s0088g01040 | 3 | 9270928 | 9285459 |
| VIT_03s0088g01050 | 3 | 9286043 | 9295905 |
| VIT_03s0088g01060 | 3 | 9296849 | 9309519 |
| VIT_03s0088g01070 | 3 | 9310070 | 9311966 |
| VIT_03s0088g01080 | 3 | 9311967 | 9314129 |
| VIT_03s0088g01090 | 3 | 9340156 | 9341562 |
| VIT_03s0088g01100 | 3 | 9342100 | 9347312 |
| VIT_03s0088g01130 | 3 | 9373663 | 9376242 |
| VIT_03s0088g01140 | 3 | 9385888 | 9388788 |
| VIT_03s0088g01150 | 3 | 9394421 | 9397946 |
| VIT_03s0088g01160 | 3 | 9408764 | 9412394 |
| VIT_03s0088g01170 | 3 | 9414656 | 9433536 |
| VIT_03s0088g01180 | 3 | 9438885 | 9442060 |
| VIT_03s0088g01190 | 3 | 9452241 | 9454982 |
| VIT_03s0088g01200 | 3 | 9455172 | 9457915 |
| VIT_03s0088g01220 | 3 | 9506440 | 9508439 |
| VIT_03s0088g01240 | 3 | 9525109 | 9527109 |
| VIT_03s0088g01250 | 3 | 9538993 | 9552353 |
| VIT_03s0088g01260 | 3 | 9553280 | 9614184 |
| VIT_03s0088g01270 | 3 | 9672111 | 9672900 |
| VIT_03s0088g01280 | 3 | 9675335 | 9678621 |
| VIT_03s0088g01290 | 3 | 9698489 | 9706945 |
| VIT_03s0088g01300 | 3 | 9707224 | 9708876 |
| VIT_00s0282g00060 | 3 | 9723604 | 9724492 |
| VIT_00s0282g00050 | 3 | 9731330 | 9732112 |
|  |  |  |  |

Pathogenesis related protein 1 precursor [Vitis vinifera]
Unknown
Zinc finger (C3HC4-type ring finger) Pathogenesis-related protein 1 precursor (PRP 1)

Unknown protein
Unknown protein
No hit
Unknown protein
Metal transporter CNNM4 (Cyclin-M4) Unknown protein

No hit Aminotransferase AGD2
Transposon protein, Mutator sub-class Aminotransferase AGD2 Mutator-like transposase

No hit
Zinc finger (C3HC4-type ring finger) Leaf senescence related protein-like EMB2758 (embryo defective 2758)

Squalene monooxygenase 2
Squalene monooxygenase
Unknown protein
Proline iminopeptidase Proline iminopeptidase
Malate dehydrogenase, glyoxysomal precursor
Unknown protein
Unknown protein
basic helix-loop-helix (bHLH) family
D-threo-aldose 1-dehydrogenase TIP growth defective 1

No hit
No hit
Inorganic pyrophosphatase
No hit
No hit
Separase

| VIT_00s0282g00040 | 3 | 9752559 | 9770979 | Major facilitator superfamily protein (MFS) Spinster |
| :---: | :---: | :---: | :---: | :---: |
| VIT_00s0282g00030 | 3 | 9781570 | 9797118 | Leaf senescence protein |
| VIT_00s0282g00020 | 3 | 9805549 | 9820005 | K+ efflux antiporter (KEA3) |
| VIT_00s0282g00010 | 3 | 9821313 | 9827537 | Methionine sulfoxide reductase |
| VIT_03s0097g00010 | 3 | 9843656 | 9843781 | No hit |
| VIT_03s0097g00030 | 3 | 9871902 | 9873992 | Concanavalin A lectin |
| VIT_03s0097g00040 | 3 | 9876165 | 9877081 | No hit |
| VIT_03s0097g00060 | 3 | 9887464 | 9888181 | Arginine-tRNA-protein transferase 1 |
| VIT_03s0097g00070 | 3 | 9889442 | 9890273 | Lectin protein kinase |
| VIT_03s0097g00080 | 3 | 9894840 | 9900035 | Lectin protein kinase |
| VIT_03s0097g00090 | 3 | 9909044 | 9909574 | Lectin protein kinase |
| VIT_03s0097g00110 | 3 | 9958876 | 9960414 | Concanavalin A lectin |
| VIT_03s0097g00120 | 3 | 9991550 | 9993565 | Concanavalin A lectin |
| VIT_03s0097g00130 | 3 | 10022997 | 10024832 | Concanavalin A lectin |
| VIT_03s0097g00140 | 3 | 10025188 | 10025953 | No hit |
| VIT_03s0097g00160 | 3 | 10121969 | 10123630 | putative MADS-box Agamous-like 17b (VviAGL17b) |
| VIT_03s0097g00210 | 3 | 10224330 | 10224440 | No hit |
| VIT_03s0097g00220 | 3 | 10252090 | 10253094 | 1,4-beta-mannan endohydrolase |
| VIT_03s0097g00230 | 3 | 10258196 | 10258417 | No hit |
| VIT_03s0097g00240 | 3 | 10269562 | 10272060 | No hit |
| VIT_03s0097g00250 | 3 | 10282948 | 10283841 | No hit |
| VIT_03s0097g00280 | 3 | 10349623 | 10350180 | Unknown protein |
| VIT_03s0097g00290 | 3 | 10361093 | 10362571 | Unknown |
| VIT_03s0097g00300 | 3 | 10363405 | 10365027 | Myosin-like protein XIK |
| VIT_03s0097g00330 | 3 | 10451345 | 10452028 | No hit |
| VIT_03s0097g00340 | 3 | 10470864 | 10472031 | Monofunctional aspartokinase |
| VIT_03s0097g00350 | 3 | 10472994 | 10475771 | Myosin-like protein XIK |
| VIT_03s0097g00360 | 3 | 10534330 | 10535472 | No hit |
| VIT_03s0097g00370 | 3 | 10558884 | 10559186 | No hit |
| VIT_03s0097g00380 | 3 | 10665097 | 10666177 | Monofunctional aspartokinase |
| VIT_03s0097g00390 | 3 | 10666726 | 10676862 | Unknown protein |
| VIT_03s0097g00400 | 3 | 10714440 | 10714920 | No hit |
| VIT_03s0097g00410 | 3 | 10717626 | 10717748 | No hit |
| VIT_03s0097g00450 | 3 | 10822230 | 10823579 | No hit |
| VIT_03s0097g00460 | 3 | 10833705 | 10835724 | Geraniol 10-hydroxylase |

VIT_05s0029g00990 5 1707179217090213 VIT_05s0029g01000 $\quad 5 \quad 17106964 \quad 17111001$ $\begin{array}{lllll}\text { VIT_05s0029g01030 } & 5 & 17241274 & 17241571\end{array}$ VIT_05s0029g01040 $5 \quad 1724158217243823$ $\begin{array}{lllll}\text { VIT_05s0029g01050 } & 5 & 17254865 & 17255236\end{array}$ $\begin{array}{lllll}\text { VIT_05s0029g01060 } & 5 & 17287266 & 17288837\end{array}$ VIT_05s0029g01070 $5 \quad 17290076 \quad 17305071$ VIT_05s0029g01080 $5 \quad 17305146 \quad 17321891$ VIT_05s0029g01090 5 5 $17330068 \quad 17349574$ $\begin{array}{lllll}\text { VIT_05s } 0029 \mathrm{~g} 01100 & 5 & 17352483 & 17364242\end{array}$ VIT_05s0029g01110 $5 \quad 1738065317382173$ VIT_05s0029g01120 $5 \quad 17387012 \quad 17388607$ $\begin{array}{lllll}\text { VIT_05s } 0029 \mathrm{~g} 01130 & 5 & 17397255 & 17402916\end{array}$ $\begin{array}{lllll}\text { VIT_05s0029g01140 } & 5 & 17405622 & 17411752\end{array}$ VIT_05s0029g01150 $5 \quad 17421093 \quad 17421226$ VIT_05s0029g01160 501742122717421628 VIT_05s0029g01180 5 VIT_05s0029g01200 $5 \quad 1748177317510650$ VIT_05s0029g01210 $5 \quad 17512328 \quad 17526686$ $\begin{array}{lllll}\text { VIT_-05s } 0029 \mathrm{~g} 01220 & 5 & 17527161 & 17548530\end{array}$ $\begin{array}{lllll}\text { VIT_05s0029g01230 } & 5 & 17566105 & 17570227\end{array}$ $\begin{array}{lllll}\text { VIT_05s0029g01240 } & 5 & 17581401 & 17589938\end{array}$ $\begin{array}{lllll}\text { VIT_05s0029g01260 } & 5 & 17601845 & 17608169\end{array}$ VIT_05s0029g01270 5 VIT_05s0029g01280 5 VIT_05s0029g01290 $5 \quad 17661362 \quad 17667411$ VIT_-05s0029g01300 5 $\begin{array}{lllll}\text { VIT_05s0029g01310 } & 5 & 17672690 & 17691790\end{array}$ VIT_05s0029g01340 $5 \quad 17729528 \quad 17779782$ VIT_05s0029g01350 501778068317808865 $\begin{array}{lllll}\text { VIT_05s } 0029 \mathrm{~g} 01360 & 5 & 17816849 & 17817058\end{array}$ VIT_05s0029g01370 $5 \quad 1781823717852241$ VIT_05s0029g01380 $5 \quad 1786545017890042$

Tripeptidyl-peptidase 2
No hit
Tripeptidyl-peptidase 2 HcrVf3 protein Aspartate aminotransferase Aspartate aminotransferase

No hit
ATELC/ELC
Ribosome biogenesis protein Bms1 Ribosome biogenesis protein Bms1

Alpha 1,3-glucosidase
RNA recognition motif (RRM)-containing protein
Unknown protein
WD-repeat protein
Pentatricopeptide (PPR) repeat-containing protein
Sucrose-phosphate synthase
Elongation factor 1-alpha 1
Elongation factor 1-alpha 1
Unknown
Methionyl-tRNA synthetase
Alkaline phytoceramidase Unknown protein
Unknown protein
Unknown protein
Unknown protein
Unknown protein
Unknown protein
Aspartic protease
MAGE (melanoma antigen-encoding gene) Actin related protein $2 / 3$ complex, subunit 4 Auxin-independent growth promoter (axi 1) RabGAP/TBC domain-containing protein

No hit
Ubiquitin-associated (UBA)
UV radiation resistance associated

| VIT_05s0029g01390 | 5 | 17892783 | 17894587 |
| :--- | :--- | :--- | :--- |
| VIT_05s0029g01410 | 5 | 17910253 | 17910654 |
| VIT_05s0029g01420 | 5 | 17958264 | 17962588 |
| VIT_05s0029g01430 | 5 | 17962589 | 17980463 |
| VIT_05s0029g01440 | 5 | 17982084 | 17983215 |
| VIT_05s0029g01450 | 5 | 17991431 | 17992219 |
| VIT_05s0029g01460 | 5 | 17997058 | 18001928 |
| VIT_05s0029g01470 | 5 | 18038062 | 18038862 |
| VIT_05s0029g01480 | 5 | 18039108 | 18041182 |
| VIT_05s0029g01490 | 5 | 18051810 | 18062080 |
| VIT_05s0029g01500 | 5 | 18083541 | 18084599 |
| VIT_05s0029g01510 | 5 | 18145463 | 18145669 |
| VIT_05s0029g01520 | 5 | 18164134 | 18166380 |
| VIT_05s0029g01530 | 5 | 18176485 | 18176941 |
| VIT_05s0029g01540 | 5 | 18204070 | 18207297 |
| VIT_05s0029g01570 | 5 | 18305556 | 18306917 |
| VIT_05s0029g01580 | 5 | 18309911 | 18310471 |
| VIT_05s0062g00010 | 5 | 18385412 | 18406790 |
| VIT_05s0062g00020 | 5 | 18414218 | 18419262 |
| VIT_05s0062g00050 | 5 | 18586887 | 18587566 |
| VIT_05s0062g00060 | 5 | 18595309 | 18596094 |
| VIT_05s0062g00110 | 5 | 18634743 | 18638371 |
| VIT_05s0062g00120 | 5 | 18674775 | 18679369 |
| VIT_05s0062g00130 | 5 | 18682670 | 18707735 |
| VIT_05s0062g00140 | 5 | 18711528 | 18719113 |
| VIT_05s0062g00150 | 5 | 18721220 | 18732785 |
| VIT_05s0062g00160 | 5 | 18735084 | 18743648 |
| VIT_05s0062g00200 | 5 | 18783093 | 18784137 |
| VIT_05s0062g00210 | 5 | 18784791 | 18791811 |
| VIT_05s0062g00220 | 5 | 18792041 | 18801353 |
| VIT_05s0062g00240 | 5 | 18818659 | 18820264 |
| VIT_05s0062g00250 | 5 | 18831568 | 18833016 |
| VIT_05s0062g00260 | 5 | 18834498 | 18839309 |
| VIT_05s0062g00270 | 5 | 18843664 | 18845204 |
| VIT_05s0062g00300 | 5 | 18872492 | 18879621 |
| VIT_05s0062g00310 | 5 | 18881653 | 18883335 |
|  |  |  |  |

[^0]|  |  |  |  |
| :--- | :--- | :--- | :--- |
| VIT_05s0062g00320 | 5 | 18886880 | 18887014 |
| VIT_05s0062g00340 | 5 | 18889800 | 18892378 |
| VIT_05s0062g00350 | 5 | 18895153 | 18896550 |
| VIT_05s0062g00360 | 5 | 18899918 | 18901613 |
| VIT_05s0062g00370 | 5 | 18920244 | 18920915 |
| VIT_05s0062g00400 | 5 | 18966256 | 18968160 |
| VIT_05s0062g00410 | 5 | 18968252 | 18970326 |
| VIT_05s0062g00430 | 5 | 18991958 | 18993513 |
| VIT_05s0062g00450 | 5 | 19023771 | 19025352 |
| VIT_05s0062g00460 | 5 | 19027377 | 19028774 |
| VIT_05s0062g00470 | 5 | 19035904 | 19037295 |
| VIT_05s0062g00480 | 5 | 19056125 | 19057777 |
| VIT_05s0062g00490 | 5 | 19060743 | 19062243 |
| VIT_05s 0062 g 00500 | 5 | 19065380 | 19066637 |
| VIT_05s0062g00510 | 5 | 19074901 | 19075119 |
| VIT_05s 0062 g 00520 | 5 | 19088785 | 19090792 |
| VIT_05s0062g00560 | 5 | 19149824 | 19158516 |
| VIT_05s0062g00570 | 5 | 19171107 | 19175569 |
| VIT_05s0062g00590 | 5 | 19182683 | 19184074 |
| VIT_05s0062g00610 | 5 | 19193950 | 19195592 |
| VIT_05s 0062 g 00620 | 5 | 19200908 | 19216029 |
| VIT_05s0062g00630 | 5 | 19226293 | 19228218 |
| VIT_05s0062g00640 | 5 | 19275169 | 19276859 |
| VIT_05s0062g00650 | 5 | 19303220 | 19312340 |
| VIT_05s0062g00660 | 5 | 19312821 | 19313569 |
| VIT_05s0062g00670 | 5 | 19328962 | 19334904 |
| VIT_05s0062g00680 | 5 | 19337594 | 19340715 |
| VIT_05s0062g00690 | 5 | 19347255 | 19348250 |
| VIT_05s0062g00700 | 5 | 19355732 | 19357242 |
| VIT_05s0062g00710 | 5 | 19377833 | 19379259 |
| VIT_05s0062g00720 | 5 | 19413520 | 19414986 |
| VIT_05s0062g00730 | 5 | 19415117 | 19418358 |
| VIT_05s0062g00740 | 5 | 19421537 | 19423163 |
| VIT_05s0062g00760 | 5 | 19469712 | 19473848 |
| VIT_05s 0062 g 00770 | 5 | 19490732 | 19492576 |
| VIT_05s0062g00780 | 5 | 19498492 | 19500009 |
|  |  |  |  |

No hit
UDP-glucose:flavonoid 7-O-glucosyltransferase
UDP-glucose:flavonoid 7-O-glucosyltransferase
UDP-glucose:flavonoid 7-O-glucosyltransferase
No hit
Indole-3-acetate beta-glucosyltransferase 1
No hit
UDP-glucose:flavonoid 7-O-glucosyltransferase
No hit
UDP-glucose:flavonoid 7-O-glucosyltransferase
UDP-glucose:flavonoid 7-O-glucosyltransferase
Xyloglucan endo-transglycosylase, C-terminal
No hit
No hit
Unknown
UDP-glucose:flavonoid 7-O-glucosyltransferase
No hit
UDP-glucose:flavonoid 7-O-glucosyltransferase
UDP-glucose:flavonoid 7-O-glucosyltransferase
Xyloglucan endotransglucosylase/hydrolase 23
Unknown
UDP-glucose transferase (UGT75B2)
UDP-glucose:flavonoid 7-O-glucosyltransferase
No hit
UDP-glucose:flavonoid 7-O-glucosyltransferase
No hit
Maturase
Heat shock protein 81-2 (HSP81-2)
UDP-glucose:flavonoid 7-O-glucosyltransferase
UDP-glucose:flavonoid 7-O-glucosyltransferase
UDP-glucoronosyl/UDP-glucosyl transferase UGT75C1
No hit
UDP-glucoronosyl/UDP-glucosyl transferase UGT75C1 Receptor kinase RHG4

Beta-1,3-glucanase
Rho GDP-dissociation inhibitor 2

|  | VIT_05s0062g00790 | 5 | 19501484 | 19509384 | NSL1 (necrotic spotted lesions 1) |
| :---: | :---: | :---: | :---: | :---: | :---: |
|  | VIT_05s0062g00800 | 5 | 19530733 | 19532850 | No hit |
|  | VIT 05s0062g00810 | 5 | 19532960 | 19534168 | Lipoprotein |
| pheno_7_1 | VIT_07s0151g00020 | 7 | 1111699 | 1112960 | Ankyrin |
|  | VIT_07s0151g00030 | 7 | 1116653 | 1117940 | Ankyrin |
|  | VIT_07s0151g00070 | 7 | 1121621 | 1122629 | Ankyrin |
|  | VIT_07s0151g00080 | 7 | 1124567 | 1125468 | Ankyrin |
|  | VIT_07s0151g00100 | 7 | 1127267 | 1139316 | Ankyrin |
|  | VIT_07s0151g00110 | 7 | 1143815 | 1145064 | Chlorophyllase 1 |
|  | VIT_07s0151g00130 | 7 | 1150474 | 1152362 | Chlorophyllase 1 |
|  | VIT_07s0151g00150 | 7 | 1156509 | 1163510 | Ankyrin |
|  | VIT_07s0151g00170 | 7 | 1164369 | 1165503 | Ankyrin |
|  | VIT_07s0151g00180 | 7 | 1168055 | 1169359 | Ankyrin |
|  | VIT_07s0151g00190 | 7 | 1171172 | 1172409 | Chlorophyllase 1 |
|  | VIT_07s0151g00210 | 7 | 1176483 | 1178421 | Chlorophyllase 1 |
|  | VIT_07s0151g00220 | 7 | 1180875 | 1191846 | Ankyrin |
|  | VIT_07s0151g00240 | 7 | 1194385 | 1195514 | Ankyrin |
|  | VIT_07s0151g00250 | 7 | 1197507 | 1198777 | Chlorophyllase |
|  | VIT_07s0151g00270 | 7 | 1206559 | 1206948 | Chlorophyllase 1 |
|  | VIT_07s0151g00280 | 7 | 1223073 | 1225810 | Bax inhibitor |
|  | VIT_07s0151g00290 | 7 | 1226423 | 1230024 | Unknown protein |
|  | VIT_07s0151g00300 | 7 | 1231259 | 1234402 | Unknown protein |
|  | VIT_07s0151g00310 | 7 | 1234739 | 1241465 | NIMA protein kinase NEK1 |
|  | VIT_07s0151g00340 | 7 | 1264579 | 1270254 | Sulfate transporter 3.1 (AST12) (AtST1) |
|  | VIT_07s0151g00360 | 7 | 1272684 | 1275182 | Unknown protein |
|  | VIT_07s0151g00370 | 7 | 1276735 | 1277605 | Kinase interacting |
|  | VIT_07s0151g00390 | 7 | 1284716 | 1284960 | Ribosomal protein S14 30S |
|  | VIT_07s0151g00410 | 7 | 1288652 | 1294516 | Sulfate transporter 3.1 (AST12) (AtST1) |
|  | VIT_07s0151g00430 | 7 | 1295689 | 1305019 | Kinase interacting family protein |
|  | VIT_07s0151g00460 | 7 | 1313821 | 1325350 | 3-methylcrotonyl-CoA carboxylase |
|  | VIT_07s0151g00500 | 7 | 1326958 | 1332156 | MLK/Raf-related protein kinase 1 |
|  | VIT_07s0151g00520 | 7 | 1334807 | 1343225 | Homocysteine S-methyltransferase 3 |
|  | VIT_07s0151g00530 | 7 | 1344129 | 1346891 | Unknown protein |
|  | VIT_07s0151g00550 | 7 | 1350567 | 1353627 | BUD32 family protein kinase |
|  | VIT_07s0151g00570 | 7 | 1356706 | 1357645 | OB-fold nucleic acid binding domain containing protein |
|  | VIT_07s0151g00590 | 7 | 1363622 | 1368249 | Inositol-1,4,5-trisphosphate 5-phosphatase CVP2, type I |


|  | VIT_07s0151g00610 | 7 | 1386497 | 1394126 |
| :---: | :---: | :---: | :---: | :---: |
|  | VIT_07s0151g00620 | 7 | 1402324 | 1404679 |
|  | VIT_07s0151g00630 | 7 | 1405392 | 1407468 |
|  | VIT_07s0151g00640 | 7 | 1408097 | 1410570 |
|  | VIT_07s0151g00660 | 7 | 1411745 | 1415073 |
|  | VIT_07s0151g00670 | 7 | 1415670 | 1416297 |
|  | VIT_07s0151g00690 | 7 | 1417935 | 1421821 |
|  | VIT_07s0151g00700 | 7 | 1422111 | 1423298 |
|  | VIT_07s0151g00720 | 7 | 1424054 | 1424497 |
|  | VIT_07s0151g00730 | 7 | 1428488 | 1429004 |
|  | VIT_07s0151g00740 | 7 | 1430006 | 1434168 |
|  | VIT_07s0151g00760 | 7 | 1435365 | 1436689 |
|  | VIT_07s0151g00770 | 7 | 1437817 | 1442530 |
|  | VIT_07s0151g00790 | 7 | 1443772 | 1448869 |
|  | VIT_07s0151g00800 | 7 | 1454149 | 1456559 |
|  | VIT_07s0151g00810 | 7 | 1459489 | 1462135 |
|  | VIT_07s0151g00830 | 7 | 1467714 | 1468455 |
|  | VIT_07s0151g00840 | 7 | 1469525 | 1471546 |
|  | VIT_07s0151g00850 | 7 | 1475572 | 1476564 |
|  | VIT_07s0151g00870 | 7 | 1478086 | 1496830 |
|  | VIT_07s0151g00900 | 7 | 1499105 | 1501968 |
|  | VIT_07s0151g00910 | 7 | 1507287 | 1509238 |
|  | VIT_07s0151g00930 | 7 | 1535022 | 1535998 |
|  | VIT_07s0151g00950 | 7 | 1540668 | 1543574 |
|  | VIT_07s0151g00960 | 7 | 1545126 | 1547795 |
|  | VIT_07s0151g00970 | 7 | 1548436 | 1560633 |
| pheno_11_1 | VIT_11s0016g03590 | 11 | 2932300 | 2940485 |
|  | VIT_11s0016g03600 | 11 | 2943205 | 2945209 |
|  | VIT_11s0016g03610 | 11 | 2948392 | 2950502 |
|  | VIT_11s0016g03630 | 11 | 2964436 | 2968009 |
|  | VIT_11s0016g03640 | 11 | 2972017 | 2974625 |
|  | VIT_11s0016g03650 | 11 | 2976690 | 2979682 |
|  | VIT_11s0016g03660 | 11 | 2982086 | 2991775 |
|  | VIT_11s0016g03670 | 11 | 2994483 | 3005396 |
|  | VIT_11s0016g03680 | 11 | 3008220 | 3017923 |
|  | VIT 11s0016g03690 | 11 | 3021953 | 3031545 |

Pentatricopeptide (PPR) repeat-containing protein
Globulin-1 S allele precursor
Globulin-1 S allele precursor
Ribosomal protein L24 (At5g23535) 50S
No hit
Proteasome 20S beta subunit D (PBD1) (PRGB)
EDA4 (embryo sac development arrest 4)
Lipid transfer protein
Unknown
Non-specific lipid-transfer protein
Lipid transfer protein
Kelch repeat-containing F-box protein Transducin family protein / WD-40 repeat

Unknown
Unknown protein HSP20 chaperone
Late embryogenesis abundant protein D-34 (LEA D-34)
Unknown
PHD finger transcription factor
Ribosomal protein S24 (RPS24A) 40S
Unknown
Unknown protein
Unknown protein
WD repeat domain 5
F-box protein
Proton-dependent oligopeptide transport (POT) family protein Proton-dependent oligopeptide transport (POT) family protein Peroxiredoxin-5
Rac-like GTP-binding protein ARAC7 (GTPase protein ROP9)
CDKF; 1 (CDK-activating kinase 1A Glucose transporter 2 plastidic Zinc finger (Ran-binding)
Tesmin/TSO1-like CXC domain-containing
Peroxisomal fatty acid beta-oxidation multifunctional protein (aim1)

| VIT_11s0016g03700 | 11 | 3037438 | 3043744 |
| :--- | :--- | :--- | :--- |
| VIT_11s0016g03710 | 11 | 3044618 | 3049139 |
| VIT_11s0016g03720 | 11 | 3049799 | 3055105 |
| VIT_11s0016g03730 | 11 | 3057394 | 3057525 |
| VIT_11s0016g03740 | 11 | 3057526 | 3060476 |
| VIT_11s0016g03750 | 11 | 3061340 | 3072065 |
| VIT_11s0016g03760 | 11 | 3079092 | 3081830 |
| VIT_11s0016g03770 | 11 | 3082194 | 3082576 |
| VIT_11s0016g03780 | 11 | 3088819 | 3105779 |
| VIT_11s0016g03790 | 11 | 3107842 | 3108617 |
| VIT_11s0016g03800 | 11 | 3108853 | 3113767 |
| VIT_11s0016g03810 | 11 | 3121899 | 3123934 |
| VIT_11s0016g03820 | 11 | 3124371 | 3128849 |
| VIT_11s0016g03830 | 11 | 3130738 | 3133954 |
| VIT_11s0016g03840 | 11 | 3134022 | 3136115 |
| VIT_11s0016g03850 | 11 | 3138660 | 3143394 |
| VIT_11s0016g03860 | 11 | 3143481 | 3153202 |
| VIT_11s0016g03870 | 11 | 3153879 | 3162165 |
| VIT_11s0016g03880 | 11 | 3163900 | 3169609 |
| VIT_11s0016g03890 | 11 | 3174051 | 3179257 |
| VIT_11s0016g03900 | 11 | 3182349 | 3186809 |
| VIT_11s0016g03910 | 11 | 3186960 | 3190875 |
| VIT_11s0016g03920 | 11 | 3191384 | 3193086 |
| VIT_11s0016g03930 | 11 | 3213014 | 3215223 |
| VIT_11s0016g03940 | 11 | 3224068 | 3225265 |
| VIT_11s0016g03950 | 11 | 3234837 | 3236255 |
| VIT_11s0016g03960 | 11 | 3236395 | 3243211 |
| VIT_11s0016g03970 | 11 | 3245946 | 3247926 |
| VIT_11s0016g03980 | 11 | 3249496 | 3250787 |
| VIT_11s0016g03990 | 11 | 3251962 | 3254889 |
| VIT_11s0016g04000 | 11 | 3255776 | 3256586 |
| VIT_11s0016g04010 | 11 | 3263191 | 3268784 |
| VIT_11s0016g04020 | 11 | 3271684 | 3276160 |
| VIT_11s0016g04030 | 11 | 3276161 | 3276314 |
| VIT_11s0016g04040 | 11 | 3289562 | 3294574 |
| VIT_11s0016g04050 | 11 | 3298356 | 3299653 |
|  |  |  |  |

Pentatricopeptide (PPR) repeat-containing protein
Abl interactor protein 1 (ABIL1)
Aspartate aminotransferase, cytoplasmic (Transaminase A)

## No hit

Rho guanyl-nucleotide exchange factor ROPGEF8 Myb-related protein 3R-1 (Plant c-MYB-like protein 1) MYB3R1 Unknown protein
Dynein light chain LC6, flagellar outer arm Continuous vascular ring (COV1)

Unknown protein
Signal peptidase I
Protein kinase

## No hit

Peroxisomal membrane 22 kDa
Raspberry 3 Subtilisin
Receptor protein kinase PERK1
Short-chain dehydrogenase/reductase (SDR)
AAA-type ATPase
Unknown protein
Polyol transporter 5
Dimethylaniline monooxygenase, N -oxide-forming
Heat shock transcription factor C1
Dehydration-responsive protein (RD22)
Kinesin family member 2/24
Ubiquitin-conjugating enzyme E2 D/E
Unknown protein
R protein PRF disease resistance protein
Unknown protein
Unknown protein
Ras-related protein Rab-5A
No hit
Unknown protein
No hit

|  | VIT_11s0016g04060 | 11 | 3306255 | 3323305 | MUT9-related serine/threonine protein kinase |
| :---: | :---: | :---: | :---: | :---: | :---: |
|  | VIT_11s0016g04070 | 11 | 3329915 | 3336836 | Hydroxyproline-rich glycoprotein |
|  | VIT_11s0016g04080 | 11 | 3340651 | 3341251 | Multiprotein-bridging factor 1c MBF1C |
|  | VIT_11s0016g04090 | 11 | 3341776 | 3358113 | DNA repair protein MutS |
| pheno_12_1 | VIT_12s0035g01900 | 12 | 23792777 | 23795385 | Pectinesterase family |
|  | VIT_12s0035g01910 | 12 | 23795440 | 23796622 | Heat shock protein 18.2 kDa class II |
|  | VIT_12s0035g01920 | 12 | 23807633 | 23823774 | Methionine S-methyltransferase |
|  | VIT_12s0035g01930 | 12 | 23823775 | 23826921 | No hit |
|  | VIT_12s0035g01940 | 12 | 23827787 | 23828051 | Unknown protein |
|  | VIT_12s0035g01950 | 12 | 23830296 | 23836655 | 4-diphosphocytidyl-2-C-methyl-D-erythritol synthase |
|  | VIT_12s0035g01960 | 12 | 23837408 | 23847527 | Stichel |
|  | VIT_12s0035g01970 | 12 | 23852741 | 23854836 | Pentatricopeptide (PPR) repeat-containing |
|  | VIT_12s0035g01980 | 12 | 23855859 | 23862754 | PHD finger protein alfin |
|  | VIT_12s0035g01990 | 12 | 23866171 | 23877161 | Intron maturase, type II |
|  | VIT_12s0035g02000 | 12 | 23879356 | 23879778 | Arachidonic acid-induced protein DEA1 |
|  | VIT_12s0035g02010 | 12 | 23888449 | 23889619 | Ribosomal RNA 23S |
|  | VIT_12s0035g02020 | 12 | 23896510 | 23899220 | NAC domain-containing protein (VvNAC40) |
|  | VIT_12s0035g02030 | 12 | 23915422 | 23924545 | Unknown protein |
|  | VIT_12s0035g02040 | 12 | 23925751 | 23927332 | No hit |
|  | VIT_12s0035g02060 | 12 | 23949395 | 23955283 | Methionine S-methyltransferase |
|  | VIT_12s0035g02070 | 12 | 23964667 | 23970073 | Cinnamoyl-CoA reductase |
|  | VIT_12s0035g02080 | 12 | 23970815 | 23972655 | Unknown protein |
|  | VIT_12s0035g02090 | 12 | 23983677 | 23999372 | Leucine-rich repeat family protein |
|  | VIT_12s0035g02100 | 12 | 24024790 | 24027813 | Glutathione S-transferase Z1 GSTZ1 |
|  | VIT_12s0035g02110 | 12 | 24028288 | 24038974 | Glutathione S-transferase Z1 GSTZ1 |
|  | VIT_12s0035g02120 | 12 | 24046092 | 24050103 | Unknown |
|  | VIT_12s0035g02150 | 12 | 24097742 | 24106206 | ferric reduction oxidase 7 FRO7 |
|  | VIT_12s0035g02160 | 12 | 24122892 | 24125541 | No hit |
|  | VIT_12s0035g02170 | 12 | 24125542 | 24127807 | No hit |
|  | VIT_12s0035g02180 | 12 | 24145254 | 24150267 | Endo-1,4-beta-glucanase |
|  | VIT 12s0035g02190 | 12 | 24151051 | 24158793 | MLO-like protein 13 |
| pheno_14_1 | VIT_14s0083g00390 | 14 | 22431591 | 22456772 | Calcium exchanger |
|  | VIT_14s0083g00400 | 14 | 22465090 | 22468501 | Leaf senescence protein |
|  | VIT_14s0083g00410 | 14 | 22474094 | 22475171 | Unknown |
|  | VIT_14s0083g00420 | 14 | 22498682 | 22502947 | TRN2 (TORNADO 2) |


| VIT_14s0083g00430 | 14 | 22507642 | 22508026 |
| :--- | :--- | :--- | :--- |
| VIT_14s0083g00440 | 14 | 22511577 | 22526125 |
| VIT_14s0083g00450 | 14 | 22528546 | 22535692 |
| VIT_14s0083g00460 | 14 | 22547153 | 22550143 |
| VIT_14s0083g00470 | 14 | 22556235 | 22557761 |
| VIT_14s0083g00480 | 14 | 22557762 | 22560665 |
| VIT_14s0083g00490 | 14 | 22560666 | 22560982 |
| VIT_14s0083g00500 | 14 | 22560983 | 22561672 |
| VIT_14s0083g00510 | 14 | 22562747 | 22563358 |
| VIT_14s0083g00520 | 14 | 22568746 | 22570772 |
| VIT_14s0083g00530 | 14 | 22581005 | 22585214 |
| VIT_14s0083g00540 | 14 | 22586239 | 22598780 |
| VIT_14s0083g00550 | 14 | 22600241 | 22604340 |
| VIT_14s0083g00570 | 14 | 22609625 | 22614966 |
| VIT_14s0083g00580 | 14 | 22620849 | 22623743 |
| VIT_14s0083g00590 | 14 | 22625232 | 22637079 |
| VIT_14s0083g00600 | 14 | 22639538 | 22640532 |
| VIT_14s0083g00610 | 14 | 22649341 | 22651396 |
| VIT_14s0083g00620 | 14 | 22672469 | 22675655 |
| VIT_14s0083g00630 | 14 | 22677598 | 22687861 |
| VIT_14s0083g00640 | 14 | 22696160 | 22698346 |
| VIT_14s0083g00650 | 14 | 22700284 | 22702204 |
| VIT_14s0083g00660 | 14 | 22712276 | 22717744 |
| VIT_14s0083g00670 | 14 | 22722348 | 22722787 |
| VIT_14s0083g00680 | 14 | 22768564 | 22768923 |
| VIT_14s0083g00690 | 14 | 22797698 | 22802306 |
| VIT_14s0083g00700 | 14 | 22839619 | 22840360 |
| VIT_14s0083g00710 | 14 | 22855118 | 22856402 |
| VIT_14s0083g00720 | 14 | 22862598 | 22865888 |
| VIT_14s0083g00730 | 14 | 22866088 | 22869527 |
| VIT_14s0083g00740 | 14 | 22886669 | 22892408 |
| VIT_14s0083g00750 | 14 | 22893111 | 22952397 |
| VIT_14s0083g00760 | 14 | 22952398 | 22953336 |
| VIT_14s0083g00770 | 14 | 22957850 | 22959392 |
| VIT_14s0083g00780 | 14 | 22961013 | 22976952 |
| VIT 14s0083g00790 | 14 | 22977589 | 22994438 |
|  |  |  |  |

PHD finger transcription factor
Protein kinase PKN/PRK1, effector
Tryptophan synthase beta chain 2
2,3-biphosphoglycerate-independent phosphoglycerate mutase
2,3-biphosphoglycerate-independent phosphoglycerate mutase
Phosphoglycerate mutase
No hit
No hit
Proline oxidase
Cell division protein FtsH Plant adhesion molecule 1 (PAM1) Pentatricopeptide (PPR) repeat-containing Plant adhesion molecule 1 (PAM1)
S-adenosyl-L-methionine decarboxylase Ribosomal RNA-processing protein 7

Unknown
Pentatricopeptide (PPR) repeat-containing protein
NIK1 (NSP- interacting kinase 1)
F-box domain containing protein
Constans 2 (COL2)
No hit
Aspartic-type endopeptidase
No hit
No hit
Glycosyl transferase family 8 protein Basic Leucine Zipper Transcription Factor (VvbZIP39) Zinc finger (C3HC4-type ring finger) Pentatricopeptide (PPR) repeat-containing

Unknown protein
Amino acid permease
Unknown protein
No hit
Phytoene desaturase
Anion exchanger adaptor protein Kanadaptin
DNA-directed RNA polymerase

| VIT_14s0083g00800 | 14 | 23001150 | 23005055 |
| :--- | :--- | :--- | :--- |
| VIT_14s0083g00810 | 14 | 23017927 | 23019683 |
| VIT_14s0083g00820 | 14 | 23022233 | 23026183 |
| VIT_14s0083g00830 | 14 | 23043321 | 23046274 |
| VIT_14s0083g00840 | 14 | 23047679 | 23050334 |
| VIT_14s0083g00850 | 14 | 23056787 | 23058566 |
| VIT_14s0083g00870 | 14 | 23073964 | 23082418 |
| VIT_14s0083g00880 | 14 | 23082779 | 23088930 |
| VIT_14s0083g00890 | 14 | 23094965 | 23095719 |
| VIT_14s0083g00900 | 14 | 23095736 | 23096308 |
| VIT_14s0083g00910 | 14 | 23106621 | 23115040 |
| VIT_14s0083g00920 | 14 | 23121142 | 23123350 |
| VIT_14s0083g00930 | 14 | 23125718 | 23136127 |
| VIT_14s0083g00940 | 14 | 23137130 | 23143930 |
| VIT_14s0083g00950 | 14 | 23153810 | 23160185 |
| VIT_14s0083g00960 | 14 | 23160685 | 23194222 |
| VIT_14s0083g00970 | 14 | 23215781 | 23219760 |
| VIT_14s0083g00980 | 14 | 23255198 | 23256315 |
| VIT_14s0083g00990 | 14 | 23257443 | 23277391 |
| VIT_14s0083g01000 | 14 | 23286167 | 23293023 |
| VIT_14s0083g01010 | 14 | 23295051 | 23298839 |
| VIT_14s0083g01020 | 14 | 23306141 | 23319918 |
| VIT_14s0083g01030 | 14 | 23320331 | 23341036 |
| VIT_14s0083g01050 | 14 | 23363261 | 23379498 |
| VIT_14s0083g01060 | 14 | 23384136 | 23385107 |
| VIT_14s0083g01070 | 14 | 23391103 | 23396928 |
| VIT_14s0083g01090 | 14 | 23416450 | 23425111 |
| VIT_14s0083g01100 | 14 | 23425741 | 23428785 |
| VIT_14s0083g01110 | 14 | 23435436 | 23438457 |
| VIT_14s0083g01120 | 14 | 23456752 | 23469960 |
| VIT_14s0083g01130 | 14 | 23470715 | 23473749 |
| VIT_14s0083g01140 | 14 | 23478460 | 23480345 |
| VIT_14s0083g01150 | 14 | 23524839 | 23527545 |
| VIT_14s0083g01160 | 14 | 23527926 | 23532692 |
| VIT_14s0083g01170 | 14 | 23552658 | 23569757 |
| VIT_14s0083g01180 | 14 | 23573219 | 23576042 |
|  | 14 |  |  |

Esterase/lipase/thioesterase
F-box family protein
Lipase GDSL 7
Lipase GDSL 7
Lipase GDSL 7
Lipase GDSL 7
Nodulation receptor kinase
Phosphatidylinositol 4-kinase type-II Transport inhibitor response 1 protein

Aspartate aminotransferase
Auxin-independent growth promoter
No hit
BIM2 (BES1-interacting Myc-like protein 2) Auxin-independent growth promoter
U3 small nucleolar RNA-associated protein IMP3
Auxin transport protein (BIG)
Ribosomal protein L7A (RPL7aB) 60S
Pollen Ole e 1 allergen and extensin
IMP dehydrogenase
Zinc finger (C3HC4-type ring finger)
Protein disulfide oxidoreductase DSBA oxidoreductase
Regulator of chromosome condensation (RCC1) putative MADS-box Fruitfull 2 (VviFUL2) putative MADS-box sepallata 1 (VviSEP1)
myb domain protein 121
No hit
N2,N2-dimethylguanosine tRNA methyltransferase
Alpha-1,4-glucan-protein synthase 1 Brassinosteroid-6-oxidase MAP3K-like protein kinase, putative, expressed MAP3K-like protein kinase, putative, expressed

B12D
COBRA protein COBRA protein
Mitochondrial substrate carrier family protein
Unknown protein

| 14s0083g01190 | 14 | 23585947 | 23586226 |
| :---: | :---: | :---: | :---: |
| VIT_14s0083g01200 | 14 | 23603520 | 23606026 |
| VIT_14s0083g01210 | 14 | 23631468 | 23634185 |
| VIT_14s0083g01220 | 14 | 23647671 | 23648618 |
| VIT_14s0083g01230 | 14 | 23687371 | 23687925 |
| VIT_14 | 14 | 23691896 | 23694505 |
| VIT_14s0068g00020 | 14 | 23702632 | 23703602 |
| VIT_14s0068g00030 | 14 | 23710282 | 23713253 |
| VIT_14s0068g00040 | 14 | 23730955 | 23731566 |
| VIT_14s0068g00050 | 14 | 23741203 | 23741804 |
| VIT_14s0068g00060 | 14 | 23761143 | 23763198 |
| VIT_14s0068g00070 | 14 | 23763889 | 23765688 |
| VIT_14s0068g00080 | 14 | 23790042 | 23793197 |
| VIT_14s0068g00090 | 14 | 23793482 | 23796203 |
| VIT_14s0068g00100 | 14 | 23804737 | 23805618 |
| VIT_14s0068g00110 | 14 | 23806944 | 23824202 |
| VIT_14s0068g00120 | 14 | 2382420 | 23824397 |
| VIT_14s0068g00130 | 14 | 23824529 | 23825866 |
| VIT_14s0068g00140 | 14 | 23848301 | 23848892 |
| VIT_14s0068g00150 | 14 | 23858719 | 23859396 |
| VIT_14s0068g00160 | 14 | 23866963 | 23867506 |
| VIT_14s0068g00170 | 14 | 23880217 | 23880588 |
| VIT_14s0068g00190 | 14 | 23901133 | 23901667 |
| VIT_14s0068g00200 | 14 | 23903015 | 23903642 |
| VIT_14s0068g00210 | 14 | 23920911 | 23938861 |
| VIT_14s0068g00220 | 14 | 23939286 | 23945038 |
| VIT_14s0068g00230 | 14 | 23946758 | 23949751 |
| VIT_14s0068g00240 | 14 | 23956348 | 23964987 |
| VIT_14s0068g00250 | 14 | 23967557 | 23970244 |
| VIT_14s0068g00260 | 14 | 23972158 | 23975509 |
| VIT_14s0068g00270 | 14 | 23981569 | 23982426 |
| VIT_14s0068g00280 | 14 | 23982459 | 23984970 |
| VIT_14s0068g00290 | 14 | 23993049 | 23996736 |
| VIT_14s0068g00300 | 14 | 23997514 | 24000870 |
| VIT_14s0068g00310 | 14 | 24001531 | 24018716 |
| VIT_14s0068g00320 | 14 | 24031909 | 2403194 |

S-adenosylmethionine synthetase 1 (SAM1)
Nitrilase
feronia receptor-like kinase feronia receptor-like kinase Heat shock protein 81-4 (HSP81-4) feronia receptor-like kinase

No hit
feronia receptor-like kinase
No hit
No hit
No hit
feronia receptor-like kinase
Non-symbiotic hemoglobin class 1 feronia receptor-like kinase feronia receptor-like kinase feronia receptor-like kinase feronia receptor-like kinase feronia receptor-like kinase

ABC Transporter (VvMDR16-VvABCB16)
Proteasome
Spiral 1 like 2
Rough sheath2-interacting KH-domain protein Exocyst subunit EXO70 E1 Serine carboxypeptidase S10
Hydroxyproline-rich glycoprotein
Unknown protein
RPK1 (receptor-like protein kinase 1)
ABRC5
Nuclear transport factor 2 (NTF2)
No hit

| VIT_14s0068g00330 | 14 | 24046880 | 24048369 |
| :--- | :--- | :--- | :--- |
| VIT_14s0068g00340 | 14 | 24058921 | 24060831 |
| VIT_14s0068g00360 | 14 | 24075902 | 24080124 |
| VIT_14s0068g00370 | 14 | 24080150 | 24081477 |
| VIT_14s0068g00380 | 14 | 24088765 | 24091151 |
| VIT_14s0068g00390 | 14 | 24101322 | 24107332 |
| VIT_14s0068g00400 | 14 | 24127197 | 24145861 |
| VIT_14s0068g00410 | 14 | 24150294 | 24158106 |
| VIT_14s0068g00420 | 14 | 24160652 | 24204411 |
| VIT_14s0068g00430 | 14 | 24208142 | 24215337 |
| VIT_14s0068g00440 | 14 | 24220054 | 24221587 |
| VIT_14s0068g00450 | 14 | 24222577 | 24224432 |
| VIT_14s0068g00460 | 14 | 24225718 | 24227484 |
| VIT_14s0068g00470 | 14 | 24232271 | 24232869 |
| VIT_14s0068g00480 | 14 | 24235845 | 24236264 |
| VIT_14s0068g00490 | 14 | 24236265 | 24240475 |
| VIT_14s0068g00500 | 14 | 24240476 | 24246281 |
| VIT_14s0068g00510 | 14 | 24246986 | 24247718 |
| VIT_14s0068g00520 | 14 | 24253217 | 24259740 |
| VIT_14s0068g00600 | 14 | 24335233 | 24336088 |
| VIT_14s0068g00630 | 14 | 24364739 | 24437849 |
| VIT_14s0068g00640 | 14 | 24438706 | 24450994 |
| VIT_14s0068g00650 | 14 | 24461539 | 24463112 |
| VIT_14s0068g00660 | 14 | 24473597 | 24477950 |
| VIT_14s0068g00670 | 14 | 24478682 | 24479739 |
| VIT_14s0068g00680 | 14 | 24482406 | 24484689 |
| VIT_14s0068g00690 | 14 | 24485059 | 24492325 |
| VIT_14s0068g00700 | 14 | 24492483 | 24497875 |
| VIT_14s0068g00710 | 14 | 24498868 | 24504221 |
| VIT_14s0068g00720 | 14 | 24509679 | 24524403 |
| VIT_14s0068g00730 | 14 | 24525207 | 24525867 |
| VIT_14s0068g00740 | 14 | 24532101 | 24533135 |
| VIT_14s0068g00750 | 14 | 24534760 | 24535908 |
| VIT_14s0068g00760 | 14 | 24536260 | 24538272 |
| VIT_14s0068g00770 | 14 | 24540054 | 24544838 |
| VIT_14s0068g00780 | 14 | 24545158 | 24555142 |
|  |  |  |  |

PTF1 (plastid transcription factor 1) TCP13
Transcription initiation factor TFIID subunit 8
LNG1 (LONGIFOLIA1)
No hit
NLI interacting factor (NIF) family protein
Esterase PIR7B
Hydrolase, alpha/beta fold family
Transducin family protein / WD-40 repeat Alpha-amylase isozyme C2 precursor Dentin sialophosphoprotein
UDP-glucosyl transferase
UDP-glucoronosyl/UDP-glucosyl transferase
UDP-glucosyl transferase
UDP-glucosyl transferase
UDP-glucosyl transferase
UDP-glucuronosyl/UDP-glucosyltransferase
Indole-3-acetate beta-glucosyltransferase
Unknown
F-box domain containing protein Expansin (VvEXPA15)

Importin
Acetyl-CoA synthetase
Unknown
ATAN11 (ANTHOCYANIN11) (VvWDR2)
Heavy-metal-associated domain-containing protein
Glyceraldehyde-3-phosphate dehydrogenase A, chloroplast precursor OTU cysteine protease

EMB1895
DEAD/DEAH box helicase
Tudor domain protein 4 SNc
Unknown protein
Avr9/Cf-9 rapidly elicited protein 146
C 2 domain-containing protein
Pentatricopeptide (PPR) repeat-containing
FAR1-related sequence 5
Nematode chemoreceptor

|  | VIT_14s0068g00790 | 14 | 24569502 | 24570830 | Unknown protein |
| :---: | :---: | :---: | :---: | :---: | :---: |
|  | VIT_14s0068g00800 | 14 | 24574332 | 24590300 | Syntaxin 7 |
|  | VIT_14s0068g00810 | 14 | 24592155 | 24604605 | Helicase |
|  | VIT_14s0068g00820 | 14 | 24605414 | 24606138 | Lipid transfer protein |
|  | VIT_14s0068g00830 | 14 | 24606690 | 24609786 | 2,3,4,5-tetrahydropyridine-2,6-dicarboxylate N -succinyltransferase |
|  | VIT_14s0068g00840 | 14 | 24612133 | 24616683 | RBR1 (retinoblastoma-related 1) |
|  | VIT_14s0068g00850 | 14 | 24616889 | 24623539 | Potassium transporter (KUP3) |
|  | VIT_14s0068g00860 | 14 | 24626255 | 24627037 | ERF/AP2 Gene Family (VvERF070) |
|  | VIT 14s0068g00870 | 14 | 24631161 | 24634996 | Nudix hydrolase 13 |
| pheno_16_1 | VIT_16s0022g00940 | 16 | 14019798 | 14021005 | Pectinesterase PME3 |
|  | VIT_16s0022g00950 | 16 | 14025273 | 14027095 | No hit |
|  | VIT_16s0022g00960 | 16 | 14028716 | 14029381 | Invertase/pectin methylesterase inhibitor |
|  | VIT_16s0022g00970 | 16 | 14044085 | 14045832 | Pentatricopeptide (PPR) repeat-containing protein |
|  | VIT_16s0022g00980 | 16 | 14047757 | 14059196 | Adenylate kinase |
|  | VIT_16s0022g00990 | 16 | 14066536 | 14067062 | Acetolactate synthase SuRB |
|  | VIT_16s0022g01000 | 16 | 14102897 | 14103279 | Chalcone synthase [Vitis vinifera] |
|  | VIT_16s0022g01010 | 16 | 14114761 | 14116548 | Acetohydroxyacid synthase |
|  | VIT_16s0022g01020 | 16 | 14148552 | 14149551 | Chalcone synthase [Vitis vinifera] |
|  | VIT_16s0022g01030 | 16 | 14158599 | 14160569 | Acetolactate synthase SuRB |
|  | VIT_16s0022g01040 | 16 | 14229326 | 14229815 | Acetolactate synthase SuRB |
|  | VIT_16s0022g01050 | 16 | 14229816 | 14231180 | Acetolactate synthase 1, chloroplast precursor |
|  | VIT_16s0022g01070 | 16 | 14274118 | 14275623 | Chalcone synthase [Vitis vinifera] |
|  | VIT_16s0022g01080 | 16 | 14288985 | 14290858 | Acetolactate synthase 1, chloroplast precursor |
|  | VIT_16s0022g01100 | 16 | 14353302 | 14354444 | Acetohydroxyacid synthase 1 |
|  | VIT_16s0022g01110 | 16 | 14354445 | 14355089 | Acetohydroxyacid synthase 1 |
|  | VIT_16s0022g01120 | 16 | 14397803 | 14399181 | Acyl-CoA oxidase ACX3 |
|  | VIT_16s0022g01130 | 16 | 14433197 | 14459122 | No hit |
|  | VIT_16s0022g01140 | 16 | 14459402 | 14460328 | Chalcone synthase [Vitis vinifera] |
|  | VIT_16s0022g01150 | 16 | 14470887 | 14471577 | Acyl-CoA oxidase ACX3 |
|  | VIT_16s0022g01160 | 16 | 14471578 | 14472628 | Acyl-CoA oxidase ACX3 |
|  | VIT_16s0022g01190 | 16 | 14557320 | 14558246 | Chalcone synthase [Vitis vinifera] |
|  | VIT_16s0022g01210 | 16 | 14618249 | 14620000 | myb domain protein 85 |
|  | VIT_16s0022g01240 | 16 | 14679983 | 14700893 | IRE (incomplete root hair elongation) |
|  | VIT_16s0022g01250 | 16 | 14701443 | 14723650 | Unknown protein |
|  | VIT_16s0022g01260 | 16 | 14727207 | 14745148 | Metalloendopeptidase |
|  | VIT_16s0022g01280 | 16 | 14750817 | 14755409 | Metalloendopeptidase |

VIT_16s0022g01290 $16 \quad 14759440 \quad 14760595$ VIT_16s0022g01310 16 VIT_16s0022g01330 $16 \quad 14814938 \quad 14816839$ VIT 16s0022g01340 $16 \quad 1482150214838464$ VIT 16s $0022 \mathrm{~g} 01350 \quad 16 \quad 14853500 \quad 14855350$ VIT_16s0022g01360 16 VIT_16s0022g01370 $16 \quad 14864696 \quad 14869194$ VIT 16s0022g01380 $16 \quad 14886712 \quad 14887520$ VIT_16s0022g01390 $16 \quad 14889269 \quad 14893469$ VIT_16s0022g01400 $16 \quad 14900577 \quad 14901340$ VIT_16s0022g01410 $16 \quad 14908101 \quad 14910093$ VIT 16s0022g01420 $16 \quad 1491133414916210$ VIT_16s0022g01430 16 VIT_16s0022g01440 16 VIT 16s0022g01450 $16 \quad 14979177 \quad 14980434$ VIT_16s0022g01460 16 VIT_16s0022g01480 $16 \quad 16008412 \quad 15027993$ VIT_16s0022g01490 $16 \quad 1503059515031594$ VIT 16s0022g01500 $16 \quad 15042831 \quad 15045000$ VIT_16s0022g01510 16 VIT_16s0022g01520 $16 \quad 16117078 \quad 1511783$ $\begin{array}{lllll}\text { VIT } 16 \mathrm{~s} 0022 \mathrm{~g} 01540 & 16 & 15129838 & 1513108\end{array}$ VIT ${ }^{-} 16 \mathrm{~s} 0022 \mathrm{~g} 01560 \quad 16 \quad 15152968 \quad 15157791$ $\begin{array}{lllll}\text { VIT_16s0022g01570 } & 16 & 15160352 & 15162044\end{array}$ VIT_16s0022g01580 $16 \quad 15171729 \quad 15173776$ VIT 16s0022g01590 $16 \quad 15178350 \quad 15181780$ VIT_16s0022g01610 16 VIT_16s0022g01620 $16 \quad 15219297 \quad 15223055$ VIT 16s0022g01630 $16 \quad 15226817 \quad 15227624$ VIT 16s0022g01640 $16 \begin{array}{llll}15237111 & 15238586\end{array}$ VIT_16s0022g01650 $16 \quad 15243820 \quad 15246842$ VIT_16s0022g01660 $16 \quad 15252302 \quad 15254570$ VIT 16s0022g01670 $161515264752 \quad 15268846$ VIT_16s0022g01680 16 VIT_16s0022g01690 $16 \quad 1529013415296405$ VIT 16s0022g01700 $161615327451 \quad 1532766$

SDG29 (SET Domain group 29)
Hydroxymethylglutaryl coenzyme A synthase R protein PRF disease resistance protein Deoxyribodipyrimidine photolyase

No hit
Unknown
No hit
No hit
flavonoid 3-monooxygenase Cellulose synthase CSLD5

No hit
Unknown protein
Phosphate-induced protein 1
Transketolase, chloroplast precursor
UDP-D- glcucuronate 4-epimerase 2 GAE2 Ribosomal protein S20 (RPS20A) 40S

No hit
4,5-DOPA dioxygenase extradiol flavonoid $3^{\prime}, 5^{\prime}$-hydroxylase -like protein flavonoid $3^{\prime}, 5$-hydroxylase -like protein

4,5-DOPA dioxygenase extradiol
Flavonoid 3',5'-hydroxylase Nucleotide-sensitive chloride conductance regulator (ICln)

3-ketoacyl-CoA synthase
3-ketoacyl-CoA synthase
Ribosomal protein S13 (RPS13A) 40S
No hit
Receptor protein kinase
Unknown protein
Receptor serine/threonine kinase Receptor serine/threonine kinase PR5K

Receptor kinase homolog LRK10
Receptor kinase LRK10
Zinc finger (AN1-like)
Band 7 family
No hit
$\begin{array}{lllll}\text { VIT } & 16 \mathrm{~s} 0022 \mathrm{~g} 01710 & 16 & 15328128 & 15330621\end{array}$ VIT_16s0022g01720 $\begin{array}{lllll}16 & 15331961 & 15333786\end{array}$ VIT_16s0022g01730 $16 \quad 1534733815347814$ VIT 16s0022g01740 $1615361931 \quad 15364513$ VIT_16s0022g01760 16 VIT_16s0022g01770 $16 \begin{array}{llll}16467138 & 15473529\end{array}$ VIT_16s0022g01780 161548928115497566 $\begin{array}{lllll}\text { VIT } & 16 \mathrm{~s} 0022 \mathrm{~g} 01790 & 16 & 15503664 & 15504109\end{array}$ VIT_16s0022g01800 $\quad 16 \quad 15512356 \quad 15514461$ VIT_16s0022g01810 $16 \quad 1551899415527030$ VIT_16s0022g01820 $16 \quad 1552827815533919$ VIT_16s0022g01830 $16 \quad 15540620 \quad 15546227$ VIT_16s0022g01840 $16 \quad 15565361 \quad 15572463$ VIT_16s0022g01850 $16 \quad 15594331 \quad 15596264$ $\begin{array}{llllll}\text { VIT } & 16 s 0022 \mathrm{~g} 01860 & 16 & 15611019 & 15730450\end{array}$ VIT_16s0022g01870 $\quad 16 \quad 15736355 \quad 15738229$ VIT_16s0022g01880 $16 \quad 16 \quad 15741012 \quad 15741260$ VIT_16s0022g01890 $16 \quad 1574283215752020$ VIT 16s $0022 \mathrm{~g} 01900 \quad 16 \quad 15757973 \quad 15758305$ VIT_16s0022g01910 $16 \begin{array}{llll}15758306 & 15759314\end{array}$ VIT_16s0022g01920 $16 \quad 15760945 \quad 15763090$ VIT_16s0022g01960 $16 \quad 15771659 \quad 15773575$ $\begin{array}{lllll}\text { VIT_16s0022g01970 } & 16 & 15780583 & 15782142\end{array}$ VIT_16s0022g01980 $16 \begin{array}{lllll}15810041 & 15810499\end{array}$ VIT_16s0022g01990 $16 \quad 15812472 \quad 15812931$ VIT 16s0022g02000 $16 \quad 1581304715813946$ VIT_16s0022g02010 $16 \begin{array}{llll}16 & 15813947 & 15818102\end{array}$ VIT_16s0022g02020 $16 \begin{array}{llll}16847600 & 15848975\end{array}$ VIT_16s0022g02030 16161585559415861902 VIT_16s0022g02040 $16 \quad 1588856315895646$ VIT_16s0022g02050 $\quad 16 \quad 15931002 \quad 15931871$ VIT_16s0022g02060 $16 \quad 15955898 \quad 15988486$ VIT_16s0022g02070 $16 \quad 1603250716033281$ VIT_16s0022g02080 $\quad 16 \quad 16041486$ VIT_16s0022g02090 $16 \quad 16055649 \quad 16056600$ VIT_16s0022g02100 161606013616061359

Ser/Thr receptor-like kinase1
Receptor kinase homolog LRK10
6-phosphogluconate dehydrogenase
Unknown protein
FK506-binding protein genes family (VvFKBP43-1)
Phosphopyruvate hydratase.
Pentatricopeptide (PPR) repeat-containing protein
Histone deacetylase complex, SIN3 component -like 2 SNL2
RRNA-processing protein EBP2
AT-hook protein 1
Microtubule associated protein (MAP65/ASE1) pleiade
YLS7 (yellow-leaf-specific gene 7)
Clp amino terminal domain-containing protein
No hit
Cleavage and polyadenylation specificity factor subunit 1 Unknown protein

No hit
GTP-binding protein engA Pentatricopeptide (PPR) repeat Pentatricopeptide (PPR) repeat
Pentatricopeptide (PPR) repeat-containing protein
ATP-dependent Clp protease ATP-binding subunit ClpX1 (CLPX)
Anthocyanidin 3-O-glucosyltransferase
Zf A20 and AN1 domain-containing stress-associated protein 1
Unknown
Unknown
1-aminocyclopropane-1-carboxylate synthase
No hit
ERL1 (ERECTA 1)
PBS2 (PPHB susceptible 2)
Lateral organ boundaries domain gene 36
K+ efflux antiporter (KEA5)
Unknown protein
Exostosin family protein
Embryo-specific 3
Embryo-specific 3

|  | VIT_16s0022g02110 | 16 | 16067495 | 16075144 | DEAD box RNA helicase (PRH75) |
| :---: | :---: | :---: | :---: | :---: | :---: |
|  | VIT_16s0022g02120 | 16 | 16078156 | 16082489 | Metalloendopeptidase OMA1 |
|  | VIT_16s0022g02130 | 16 | 16084547 | 16085104 | No hit |
|  | VIT_16s0022g02140 | 16 | 16087685 | 16102990 | CYP704A2 |
|  | VIT_16s0022g02150 | 16 | 16103565 | 16105639 | Lectin |
|  | VIT_16s0022g02170 | 16 | 16130501 | 16143461 | Alpha-L-fucosidase |
|  | VIT_16s0022g02190 | 16 | 16182278 | 16183814 | Chalcone and stilbene synthases |
|  | VIT_16s0022g02200 | 16 | 16203664 | 16206181 | Subtilase |
|  | VIT_16s0022g02210 | 16 | 16215979 | 16220680 | Protein phosphatase 2CA AHG1 PP2CA (VvPP2C-8) |
|  | VIT_16s0022g02220 | 16 | 16230501 | 16231445 | Receptor Like Protein 48 RLP48 |
|  | VIT_16s0022g02230 | 16 | 16240572 | 16248680 | Leucine-rich repeat receptor protein kinase EXS |
|  | VIT_16s0022g02240 | 16 | 16255908 | 16257035 | basic helix-loop-helix (bHLH) family |
|  | VIT_16s0022g02250 | 16 | 16266462 | 16267491 | basic helix-loop-helix (bHLH) family |
|  | VIT_16s0022g02270 | 16 | 16287811 | 16288987 | basic helix-loop-helix (bHLH) family |
|  | VIT_16s0022g02280 | 16 | 16301165 | 16302698 | Receptor protein kinase PERK1 |
|  | VIT_16s0022g02290 | 16 | 16334395 | 16337736 | Unknown |
|  | VIT_16s0022g02300 | 16 | 16350758 | 16351750 | CXE carboxylesterase CXE20 |
|  | VIT_16s0022g02310 | 16 | 16372484 | 16374087 | Gibberellin 3-beta-dioxygenase. |
|  | VIT_16s0022g02320 | 16 | 16388979 | 16407441 | Phosphoglucomutase chloroplast precursor |
|  | VIT_16s0022g02330 | 16 | 16451140 | 16466299 | putative MADS-box Agamous-like 6b (VviAGL6b) |
|  | VIT_16s0022g02340 | 16 | 16470141 | 16475595 | fructokinase-2 |
|  | VIT_16s0022g02350 | 16 | 16481981 | 16491201 | ACI13 |
|  | VIT_16s0022g02370 | 16 | 16500433 | 16503019 | Unknown protein |
| pheno_16_2 | VIT_16s0022g02380 | 16 | 16506566 | 16519567 | putative MADS-box Supressor of Constans overexpression 1b (VviSOC1b) |
|  | VIT_16s0022g02400 | 16 | 16529521 | 16544563 | MADS-box agamous-like 42 |
|  | VIT_16s0022g02410 | 16 | 16558074 | 16566697 | Oxidoreductase |
|  | VIT_16s0022g02420 | 16 | 16568466 | 16571206 | Subtilisin protease C1 |
|  | VIT_16s0022g02430 | 16 | 16590597 | 16591356 | Carboxyesterase 20 CXE20 |
|  | VIT_16s0022g02440 | 16 | 16658913 | 16659634 | Carboxyesterase 20 CXE20 |
|  | VIT_16s0022g02450 | 16 | 16672729 | 16677077 | Subtilisin protease C1 |
|  | VIT_16s0022g02460 | 16 | 16707603 | 16710429 | Subtilisin protease C1 |
|  | VIT_16s0022g02470 | 16 | 16715289 | 16716489 | Cationic peroxidase |
|  | VIT_16s0022g02480 | 16 | 16721578 | 16723006 | TCP family transcription factor TCP9 |
|  | VIT_16s0022g02490 | 16 | 16725022 | 16732924 | Hydroxymethylglutaryl coenzyme A synthase |
|  | VIT_16s0022g02500 | 16 | 16753448 | 16753788 | No hit |
|  | VIT_16s0022g02510 | 16 | 16758120 | 16758664 | Protein phosphatase 2C |

$\begin{array}{lllll}\text { VIT } & 16 s 0022 \mathrm{~g} 02520 & 16 & 16783470 & 16784544\end{array}$ VIT_16s0022g02530 16 $\begin{array}{lllll}\text { VIT_16s0022g02540 } & 16 & 16787851 & 16788601\end{array}$ VIT_16s0100g00010 $16 \quad 16802160 \quad 16803228$ VIT_16s0100g00030 $\quad 16 \quad 1681886416821466$ VIT_16s0100g00040 $16 \begin{array}{llll}16 & 16331329 & 16832283\end{array}$ VIT_16s0100g00050 $161616833888 \quad 16834887$ $\begin{array}{lllll}\text { VIT 16s } 0100 \mathrm{~g} 00060 & 16 & 16867556 & 16872590\end{array}$ VIT_16s0100g00070 $\quad 16 \quad 16882780 \quad 16883663$ VIT_16s0100g00080 $16 \quad 168889920 \quad 16890662$ VIT_16s0100g00090 161689624616897439 VIT 16s0100g00110 $16 \quad 16928141 \quad 16929702$ VIT_16s0100g00120 $\quad 16 \quad 16951370 \quad 16951710$ VIT_16s0100g00130 $16 \quad 16955981 \quad 16956564$ VIT_16s0100g00140 $16 \quad 16974820 \quad 16975365$ VIT_16s0100g00150 $16 \quad 1697710316977433$ VIT_16s0100g00160 $16 \quad 1698760216993787$ VIT_16s0100g00170 $16 \quad 17000645 \quad 17002770$ VIT 16s0100g00180 $16 \quad 17071017 \quad 17071584$ VIT_16s0100g00190 $\quad 16 \quad 17115669 \quad 17121681$ VIT_16s0100g00200 $16 \begin{array}{lllll}17122769 & 17124779\end{array}$ VIT_16s0100g00210 $16 \quad 17124780 \quad 17125575$ $\begin{array}{lllll}\text { VIT }{ }^{-} 16 s 0100 \mathrm{~g} 00220 & 16 & 17132026 & 17132339\end{array}$ VIT_16s0100g00230 16171713255317134891 VIT_16s0100g00240 $16 \quad 17134892 \quad 17139634$ VIT 16s0100g00250 $16 \quad 17139635 \quad 17141094$ VIT_16s0100g00260 $16 \begin{array}{llll}16 & 17149698 & 17150478\end{array}$ VIT_16s0100g00270 $16 \begin{array}{lllll}17150479 & 17150928\end{array}$ VIT_16s0100g00280 $161717157127 \quad 17161415$ $\begin{array}{lllll}\text { VIT_16s0100g00290 } & 16 & 17162424 & 17180217\end{array}$ VIT_16s0100g00300 $16 \quad 17186416 \quad 17190028$ VIT_16s0100g00310 $16 \quad 17196699 \quad 17198841$ VIT_16s0100g00320 $16 \quad 17225835 \quad 17233936$ VIT_16s0100g00330 $\quad 16 \quad 17237416 \quad 17238326$ VIT_16s0100g00340 $16 \quad 17239840 \quad 17244491$ VIT_16s0100g00350 $16 \quad 17248816 \quad 17261155$

ZFWD1 (zinc finger WD40 repeat protein 1)
ZFWD1 (zinc finger WD40 repeat protein 1)
No hit
Carboxyesterase 20 CXE20
Cucumisin precursor
No hit
Protein kinase
Subtilisin protease C1
Prohibitin 1
Subtilisin serine protease
Cationic peroxidase
Hydroxymethylglutaryl coenzyme A synthase
No hit
Protein phosphatase 2C / PP2C
Protein phosphatase 2C / PP2C
COP9 signalosome complex subunit 5
HMG-CoA synthase 2
Xyloglucan endotransglucosylase/hydrolaseXTR2
ATP-dependent Clp protease ClpB protein Pentatricopeptide (PPR) repeat-containing protein

Unknown protein

## No hit

No hit
Glucose-methanol-choline (GMC) oxidoreductase family protein
Mandelonitrile lyase-like protein
Mandelonitrile lyase-like protein
No hit
Peptidoglycan-binding LysM domain-containing protein
EMB2758 (embryo defective 2758)
L-idonate dehydrogenase
Sorbitol dehydrogenase
Unknown protein
Zfwd2 protein (ZFWD2)
Unknown protein

ABC Transporter (VvTAP3 - VvABCB23)
$\begin{array}{lllll}\text { VIT } 16 \mathrm{~s} 0100 \mathrm{~g} 00360 & 16 & 17265204 & 17272005\end{array}$

|  | VIT_16s0100g00370 | 16 | $17274084$ | 17324348 | Valyl-tRNA synthetase |
| :---: | :---: | :---: | :---: | :---: | :---: |
| pheno_17_1 | VIT_17s0000g04630 | 17 | 4965012 | 5015746 | Phenylalanyl-tRNA synthetase beta chain |
|  | VIT_17s0000g04640 | 17 | 5017391 | 5020555 | H/ACA ribonucleoprotein complex subunit 2 |
|  | VIT_17s0000g04650 | 17 | 5021431 | 5035538 | Phosphoribulokinase/uridine kinase |
|  | VIT_17s0000g04660 | 17 | 5042400 | 5052215 | Agmatine deiminase |
|  | VIT_17s0000g04670 | 17 | 5052921 | 5063485 | Serine/threonine protein kinase ATPK3 |
|  | VIT_17s0000g04680 | 17 | 5090954 | 5095558 | Signal peptide peptidase SPPA |
|  | VIT_17s0000g04690 | 17 | 5096080 | 5107168 | Protease SppA |
|  | VIT_17s0000g04700 | 17 | 5145018 | 5145988 | Protease SppA |
|  | VIT_17s0000g04710 | 17 | 5236137 | 5239225 | Pentatricopeptide (PPR) repeat-containing protein |
|  | VIT_17s0000g04720 | 17 | 5241247 | 5243311 | No hit |
|  | VIT_17s0000g04730 | 17 | 5257791 | 5259482 | Zinc finger (C3HC4-type ring finger) |
|  | VIT_17s0000g04740 | 17 | 5261337 | 5267944 | Transducin family protein / WD-40 repeat |
|  | VIT_17s0000g04750 | 17 | 5279676 | 5281368 | UDP-glycosyltransferase 89B2 |
|  | VIT_17s0000g04760 | 17 | 5281484 | 5283173 | UDP-glycosyltransferase 89B2 |
|  | VIT_17s0000g04770 | 17 | 5286031 | 5291504 | PHD finger transcription factor |
|  | VIT_17s0000g04780 | 17 | 5292045 | 5296110 | Phosphatidylinositol 4-kinase type-II |
|  | VIT_17s0000g04790 | 17 | 5309123 | 5318428 | BIM1 (BES1-interacting Myc-like protein 1) |
|  | VIT_17s0000g04800 | 17 | 5318431 | 5318559 | No hit |
|  | VIT_17s0000g04810 | 17 | 5321964 | 5325762 | Auxin-independent growth promoter |
|  | VIT_17s0000g04820 | 17 | 5331026 | 5337299 | BSD domain-containing protein |
|  | VIT_17s0000g04830 | 17 | 5338567 | 5345150 | DNA Polymerase epsilon, subunit B |
|  | VIT_17s0000g04840 | 17 | 5357937 | 5362286 | Sterile alpha motif (SAM) domain-containing |
|  | VIT_17s0000g04850 | 17 | 5372091 | 5373566 | OBF binding protein 4 |
|  | VIT_17s0000g04860 | 17 | 5375984 | 5380301 | Unknown |
|  | VIT_17s0000g04870 | 17 | 5390557 | 5401864 | Microtubule-associated protein MBP2C |
|  | VIT_17s0000g04880 | 17 | 5408698 | 5409427 | Dirigent protein |
|  | VIT_17s0000g04890 | 17 | 5409996 | 5423899 | D-aminoacyl-tRNA deacylase GEKO1 |
|  | VIT_17s0000g04900 | 17 | 5426843 | 5441775 | ATP-dependent RNA helicase |
|  | VIT_17s0000g04910 | 17 | 5446682 | 5449753 | L-asparaginase |
|  | VIT_17s0000g04920 | 17 | 5486192 | 5489071 | Histidine kinase (AHK3) |
|  | VIT_17s0000g04930 | 17 | 5507759 | 5509033 | Desulfo-glucosinolate sulfotransferase 18 |
|  | VIT_17s0000g04940 | 17 | 5525332 | 5525670 | Brassinosteroid sulfotransferase ST4A |
|  | VIT_17s0000g04950 | 17 | 5528234 | 5529544 | Steroid sulfotransferase |
|  | VIT_17s0000g04960 | 17 | 5531301 | 5546181 | Polygalacturonate 4-alpha-galacturonosyltransferase GAUT11 |


| VIT_17s0000g04970 | 17 | 5551329 | 5561922 |
| :--- | :--- | :--- | :--- |
| VIT_17s0000g04980 | 17 | 5562484 | 5563956 |
| VIT_17s0000g04990 | 17 | 5565646 | 5584353 |
| VIT_17s0000g05000 | 17 | 5589362 | 5596096 |
| VIT_17s0000g05010 | 17 | 5614348 | 5626304 |
| VIT_17s0000g05020 | 17 | 5637669 | 5644801 |
| VIT_17s0000g05030 | 17 | 5645732 | 5648492 |
| VIT_17s0000g05040 | 17 | 5654593 | 5656105 |
| VIT_17s0000g05050 | 17 | 5659282 | 5660704 |
| VIT_17s0000g05060 | 17 | 5661333 | 5676012 |
| VIT_17s0000g05070 | 17 | 5676169 | 5679862 |
| VIT_17s0000g05080 | 17 | 5690884 | 5710034 |
| VIT_17s0000g05090 | 17 | 5710035 | 5711791 |
| VIT_17s0000g05100 | 17 | 5723289 | 5730063 |
| VIT_17s0000g05110 | 17 | 5731853 | 5734268 |
| VIT_17s0000g05120 | 17 | 5751736 | 5758856 |
| VIT_17s0000g05130 | 17 | 5759551 | 5761024 |
| VIT_17s0000g05140 | 17 | 5763617 | 5769816 |
| VIT_17s0000g05150 | 17 | 5773018 | 5780793 |
| VIT_17s0000g05160 | 17 | 5783842 | 5789162 |
| VIT_17s0000g05170 | 17 | 5793575 | 5794544 |
| VIT_17s0000g05180 | 17 | 5794802 | 5795725 |
| VIT_17s0000g05190 | 17 | 5801569 | 5810790 |
| VIT_17s0000g05200 | 17 | 5819731 | 5826246 |
| VIT_17s0000g05210 | 17 | 5832003 | 5838142 |
| VIT_17s0000g05220 | 17 | 5845847 | 5850744 |
| VIT_17s0000g05230 | 17 | 5851190 | 5861913 |
| VIT_17s0000g05240 | 17 | 5869290 | 5885095 |
| VIT_17s0000g05250 | 17 | 5885327 | 5886300 |
| VIT_17s0000g05260 | 17 | 5886472 | 5890773 |
| VIT_17s0000g05270 | 17 | 5892658 | 5896385 |
| VIT_17s0000g05280 | 17 | 5898984 | 5902122 |
| VIT_17s0000g05290 | 17 | 5924194 | 5926786 |
| VIT_17s0000g05300 | 17 | 5926787 | 5928753 |
| VIT_17s0000g05310 | 17 | 5933739 | 5939818 |
| VIT_17s0000g05320 | 17 | 5940805 | 5944765 |
|  |  |  |  |

ADP, ATP carrier protein Universal stress protein (USP) family protein putative MADS-box Fruitfull 1 (VviFUL1) putative MADS-box sepallata 2 (VviSEP2)

Unknown
Squamosa promoter-binding protein 6 (SPL6)
Alpha-1,4-glucan-protein synthase 1
No hit
COBRA-like protein 4 Phytochelatin synthetase Phytochelatin synthetase Cleavage stimulation factor subunit 1 Pentatricopeptide (PPR) repeat-containing protein

No hit
CYP78A4
ABC Transporter (VvTAP2 - VvABCB22)
Transcription termination factor mitochondrial mTERF
Rhomboid
Enoyl-CoA hydratase SP1L2 (SPIRAL2)

No hit
Unknown
Unknown protein
NLI interacting factor (NIF) family protein LNG1 (LONGIFOLIA1) TCP family transcription factor TCP5

Unknown protein
Nuclear transport factor 2 (NTF2)
Unknown protein
Unknown protein
Unknown protein
UPF0737 protein AFP3
Pre-mRNA cleavage complex II protein Clp1 Pre-mRNA cleavage complex II protein Clp1 Pre-mRNA cleavage complex II protein Clp 1

IFA binding protein

| VIT_17s0000g05330 | 17 | 5946645 | 5949934 |
| :--- | :--- | :--- | :--- |
| VIT_17s0000g05350 | 17 | 5958587 | 5963699 |
| VIT_17s0000g05360 | 17 | 5965138 | 5966181 |
| VIT_17s0000g05370 | 17 | 5966462 | 5970848 |
| VIT_17s0000g05380 | 17 | 5975939 | 5976998 |
| VIT_17s0000g05390 | 17 | 5978634 | 5988830 |
| VIT_17s0000g05400 | 17 | 5992385 | 5993924 |
| VIT_17s0000g05410 | 17 | 5997792 | 6001685 |
| VIT_17s0000g05420 | 17 | 6003755 | 6005138 |
| VIT_17s0000g05430 | 17 | 6005483 | 6026193 |
| VIT_17s0000g05440 | 17 | 6061730 | 6069222 |
| VIT_17s0000g05450 | 17 | 6070433 | 6082026 |
| VIT_17s0000g05460 | 17 | 6088077 | 6090457 |
| VIT_17s0000g05470 | 17 | 6090728 | 6093181 |
| VIT_17s0000g05480 | 17 | 6100712 | 6112781 |
| VIT_17s0000g05490 | 17 | 6117662 | 6118846 |
| VIT_17s0000g05500 | 17 | 6120295 | 6131114 |
| VIT_17s0000g05510 | 17 | 6132145 | 6137273 |
| VIT_17s0000g05520 | 17 | 6144571 | 6153251 |
| VIT_17s0000g055530 | 17 | 6153633 | 6161033 |
| VIT_17s0000g05540 | 17 | 6174485 | 6182145 |
| VIT_17s0000g05550 | 17 | 6183508 | 6195130 |
| VIT_17s0000g05560 | 17 | 6199502 | 6202250 |
| VIT_17s0000g05570 | 17 | 6207677 | 6212987 |
| VIT_17s0000g05580 | 17 | 6213229 | 6221132 |
| VIT_17s0000g05600 | 17 | 6255629 | 6256773 |
| VIT_17s0000g05610 | 17 | 6257443 | 6258438 |
| VIT_17s0000g05620 | 17 | 6259762 | 6262348 |
| VIT_17s0000g05630 | 17 | 6265801 | 6268050 |
| VIT_17s0000g05640 | 17 | 6276619 | 6279624 |
| VIT_17s0000g05650 | 17 | 6279745 | 6280799 |
| VIT_17s0000g05660 | 17 | 6282923 | 6285609 |
| VIT_17s0000g05670 | 17 | 6286277 | 6288346 |
| VIT_17s0000g05680 | 17 | 6293644 | 6298086 |
| VIT_17s0000g05690 | 17 | 6299282 | 6301763 |
| VIT_17s0000g05700 | 17 | 6304590 | 6308202 |
| VIT |  |  |  |

Serine/threonine protein phosphatase PP2A-5 catalytic subunit (PP2A5)
5-formyltetrahydrofolate cycloligase
Germin
MYC ZCW32 (bigpetal, bigpetalub) Dynein light chain LC8-type

Unknown protein
myb domain protein 35
Galactosyltransferase
AAA-type ATPase family Aminoacyl-tRNA synthetase Unknown protein ELF5 (early flowering 5) Nodulin
Nodulin
EMB2454 (embryo defective 2454) Lateral organ boundaries domain family protein (LBD27)

Cysteine endopeptidase
Pentatricopeptide (PPR) repeat-containing protein
Calcium Dependent Protein Kinase (VvCPK14)
DnaJ homolog, subfamily B, member 4
$\mathrm{H}(+)$-ATPase 11 AHA11
Proton-dependent oligopeptide transport (POT) family protein Proton-dependent oligopeptide transport (POT) family protein

Receptor protein kinase
Isopiperitenol dehydrogenase
(-)-isopiperitenol dehydrogenase
Isopiperitenol dehydrogenase
Integral membrane HPP family protein Homeodomain leucine zipper protein HB-1

Nitrate transporter 1:2
Unknown protein
Polygalacturonase GH28
Polygalacturonase GH28
Band 7 family
Isochorismate synthase 1, chloroplast precursor
Phosphoribosylanthranilate transferase

|  | VIT_17s0000g05710 | 17 | 6312022 | 6312896 | Unknown |
| :---: | :---: | :---: | :---: | :---: | :---: |
|  | VIT_17s0000g05720 | 17 | 6319018 | 6320270 | No hit |
|  | VIT_17s0000g05730 | 17 | 6343021 | 6345217 | No hit |
|  | VIT_17s0000g05740 | 17 | 6347635 | 6349215 | Pectate lyase, N-terminal |
|  | VIT_17s0000g05750 | 17 | 6350085 | 6359530 | Isochorismate synthase |
|  | VIT_17s0000g05760 | 17 | 6361704 | 6385247 | Nuclear protein ZAP |
|  | VIT_17s0000g05770 | 17 | 6397487 | 6410902 | Nuclear ribonuclease Z |
| pheno_17_2 | VIT_17s0000g07830 | 17 | 8918616 | 8923034 | Asp/Glu racemase; Tetratricopeptide helical |
|  | VIT_17s0000g07840 | 17 | 8924188 | 8925529 | Unknown protein |
|  | VIT_17s0000g07850 | 17 | 8931341 | 8937870 | Nucleobase-ascorbate transporter 6 (NAT6) |
|  | VIT_17s0000g07870 | 17 | 8950730 | 8962366 | Ribosomal protein L29 |
|  | VIT_17s0000g07880 | 17 | 8963993 | 8974679 | Bromo-adjacenty (BAH) domain-containing protein |
|  | VIT_17s0000g07890 | 17 | 8976835 | 8985225 | Rac-like GTP-binding protein ARAC10 (GTPase protein ROP10) |
|  | VIT_17s0000g07900 | 17 | 9005528 | 9011550 | Protease Do 9 |
|  | VIT_17s0000g07910 | 17 | 9011840 | 9014726 | Nodulin MtN21 family |
|  | VIT_17s0000g07920 | 17 | 9030527 | 9032996 | Hypoxia-responsive |
|  | VIT_17s0000g07930 | 17 | 9034815 | 9036877 | Nodulin MtN21 family |
|  | VIT_17s0000g07940 | 17 | 9041797 | 9042384 | ERF/AP2 Gene Family (VvERF024) |
|  | VIT 17s0000g07950 | 17 | 9054392 | 9056553 | Unknown protein |
| pheno_18_1 | VIT_18s0001g01250 | 18 | 1835240 | 1837419 | Senescence-inducible chloroplast stay-green protein 2 |
|  | VIT_18s0001g01270 | 18 | 1856972 | 1863925 | Ankyrin |
|  | VIT_18s0001g01300 | 18 | 1877353 | 1877899 | Wall-associated receptor kinase 5 |
|  | VIT_18s0001g01310 | 18 | 1877900 | 1880259 | Wall-associated receptor kinase 5 |
|  | VIT_18s0001g01320 | 18 | 1885028 | 1894797 | Wall-associated receptor kinase 5 |
|  | VIT_18s0001g01350 | 18 | 1897393 | 1914523 | Toprim domain-containing protein |
|  | VIT_18s0001g01360 | 18 | 1934332 | 1948604 | Toprim domain-containing protein |
|  | VIT_18s0001g01370 | 18 | 1952680 | 1968096 | Wall-associated receptor kinase 5 |
|  | VIT_18s0001g01390 | 18 | 1982983 | 1985445 | GA 20-oxidase |
|  | VIT_18s0001g01410 | 18 | 1996714 | 1997619 | Unknown |
|  | VIT_18s0001g01440 | 18 | 2005999 | 2006364 | No hit |
|  | VIT_18s0001g01460 | 18 | 2028167 | 2028620 | No hit |
|  | VIT_18s0001g01470 | 18 | 2040472 | 2040960 | No hit |
|  | VIT_18s0001g01480 | 18 | 2050288 | 2050581 | No hit |
|  | VIT_18s0001g01500 | 18 | 2060078 | 2060598 | No hit |
|  | VIT_18s0001g01510 | 18 | 2075169 | 2084464 | No hit |


| VIT_18s0001g01530 | 18 | 2086640 | 2087608 |
| :--- | :--- | :--- | :--- |
| VIT_18s0001g01550 | 18 | 2123749 | 2127932 |
| VIT_18s0001g01560 | 18 | 2130873 | 2138916 |
| VIT_18s0001g01580 | 18 | 2149563 | 2165796 |
| VIT_18s0001g01590 | 18 | 2174967 | 2177070 |
| VIT_18s0001g01600 | 18 | 2177493 | 2197513 |
| VIT_18s0001g01630 | 18 | 2200625 | 2201333 |
| VIT_18s0001g01640 | 18 | 2203044 | 2218030 |
| VIT_18s0001g01650 | 18 | 2222531 | 2227556 |
| VIT_18s0001g01660 | 18 | 2227558 | 2229622 |
| VIT_18s0001g01670 | 18 | 2231148 | 2244524 |
| VIT_18s0001g01680 | 18 | 2245944 | 2250220 |
| VIT_18s0001g01690 | 18 | 2256608 | 2258038 |
| VIT_18s0001g01700 | 18 | 2259998 | 2279932 |
| VIT_18s0001g01740 | 18 | 2281218 | 2286726 |
| VIT_18s0001g01760 | 18 | 2291789 | 2294046 |
| VIT_18s0001g01770 | 18 | 2299018 | 2303152 |
| VIT_18s0001g01780 | 18 | 2306657 | 2322950 |
| VIT_18s0001g01810 | 18 | 2324024 | 2335447 |
| VIT_18s0001g01830 | 18 | 2336971 | 2345138 |
| VIT_18s0001g01840 | 18 | 2347198 | 2348370 |
| VIT_18s0001g01850 | 18 | 2352063 | 2359939 |
| VIT_18s0001g01860 | 18 | 2359940 | 2405390 |
| VIT_18s0001g01930 | 18 | 2406583 | 2411640 |
| VIT_18s0001g01950 | 18 | 2413199 | 2414815 |
| VIT_18s0001g01960 | 18 | 2415255 | 2417215 |
| VIT_18s0001g01980 | 18 | 2418945 | 2426660 |
| VIT_18s0001g02000 | 18 | 2438485 | 2442668 |
| VIT_18s0001g02010 | 18 | 2442718 | 2443387 |
| VIT_18s0001g02020 | 18 | 2450437 | 2453364 |
| VIT_18s0001g02030 | 18 | 2460774 | 2467119 |
| VIT_18s0001g02050 | 18 | 2477226 | 2479822 |
| VIT_18s0001g02060 | 18 | 2479868 | 2484109 |
| VIT_18s0001g02080 | 18 | 2485740 | 2489654 |
| VIT_18s0001g02090 | 18 | 2496692 | 2503378 |
| VIT_18s0001g02100 | 18 | 2504341 | 2505039 |


| VIT_18s0001g02120 | 18 | 2513739 | 2517442 |
| :--- | :--- | :--- | :--- |
| VIT_18s0001g02130 | 18 | 2519567 | 2521025 |
| VIT_18s0001g02140 | 18 | 2521402 | 2529629 |
| VIT_18s0001g02160 | 18 | 2543141 | 2554330 |
| VIT_18s0001g02190 | 18 | 2564080 | 2567938 |
| VIT_18s0001g02200 | 18 | 2573308 | 2584831 |
| VIT_18s0001g02220 | 18 | 2592072 | 2601990 |
| VIT_18s0001g02230 | 18 | 2604468 | 2613076 |
| VIT_18s0001g02260 | 18 | 2618439 | 2626112 |
| VIT_18s0001g02270 | 18 | 2627885 | 2629132 |
| VIT_18s0001g02280 | 18 | 2629486 | 2630245 |
| VIT_18s0001g02300 | 18 | 2643266 | 2645234 |
| VIT_18s0001g02340 | 18 | 2663270 | 2675579 |
| VIT_18s0001g02370 | 18 | 2677244 | 2684335 |
| VIT_18s0001g02380 | 18 | 2688559 | 2734881 |
| VIT_18s0001g02420 | 18 | 2734882 | 2737586 |
| VIT_18s0001g02440 | 18 | 2745381 | 2754460 |
| VIT_18s0001g02470 | 18 | 2761252 | 2772231 |
| VIT_18s0001g02480 | 18 | 2777880 | 2785087 |
| VIT_18s0001g02510 | 18 | 2791934 | 2798318 |
| VIT_18s0001g02540 | 18 | 2802829 | 2805078 |
| VIT_18s0001g02550 | 18 | 2813209 | 2821688 |
| VIT_18s0001g02570 | 18 | 2822064 | 2827157 |
| VIT_18s0001g02610 | 18 | 2845415 | 2847110 |
| VIT_18s0001g02630 | 18 | 2855229 | 2860609 |
| VIT_18s0001g02640 | 18 | 2861119 | 2864695 |
| VIT_18s0001g02650 | 18 | 2866342 | 2870749 |
| VIT_18s0001g02670 | 18 | 2875900 | 2876019 |
| VIT_18s0001g02680 | 18 | 2877451 | 2880061 |
| VIT_18s0001g02690 | 18 | 2881165 | 2881615 |
| VIT_18s0001g02700 | 18 | 2881700 | 2884919 |
| VIT_18s0001g02710 | 18 | 2885819 | 2894159 |
| VIT_18s0001g02730 | 18 | 2898842 | 2905274 |
| VIT_18s0001g02740 | 18 | 2905882 | 2907423 |
| VIT_18s0001g02750 | 18 | 2909150 | 2910927 |
| VIT_18s0001g02760 | 18 | 2910928 | 2914749 |

Metal transporter Nramp6
Metal transporter Nramp6
Metal transporter Nramp1
Switching protein 3C ATSWI3C
Switching protein 3C ATSWI3C
8-oxoguanine-DNA glycosylase (OGG1)
Beta-galactosidase / lactase
Beta-galactosidase / lactase Proteasome 20S beta subunit C1 (PBC1) (PRCT)

Unknown
Zinc finger (C3HC4-type ring finger) family protein NAC domain-containing protein (VvNAC08)

Armadillo/beta-catenin repeat
DNA replication factor C complex subunit 5
Kinesin protein (MKRP1)
Aspartic Protease (VvAP40)
Aldehyde Dehydrogenase (VvALDH3H1)
L-ascorbate peroxidase, thylakoid-bound (tAPX)
Unknown protein
Protein disulfide-isomerase A1
ARR9 typeA
Delta-aminolevulinic acid dehydratase, chloroplast precursor
IAA-amino acid hydrolase 6
Caffeic acid methyltransferase
F-box only protein 9
Unknown protein
V-type $\mathrm{H}+$-transporting ATPase 16 kDa proteolipid subunit
No hit
$\mathrm{BTB} / \mathrm{POZ}$ domain-containing protein Unknown protein
Chlorophyll a oxygenase (CAO) Unknown protein

Unknown
Photosystem II 22 kDa protein PSBS Unknown protein

Lipase GDSL

| VIT_18s0001g02770 | 18 | 2914955 | 2920663 |
| :--- | :--- | :--- | :--- |
| VIT_18s0001g02780 | 18 | 2921245 | 2922708 |
| VIT_18s0001g02790 | 18 | 2925094 | 2926968 |
| VIT_18s0001g02810 $^{2}$ | 18 | 2929082 | 2940502 |
| VIT_18s0001g02820 | 18 | 2950570 | 2959607 |
| VIT_18s0001g02830 $^{18}$ | 18 | 2960201 | 2973823 |
| VIT_18s0001g02860 | 18 | 2987834 | 2994441 |
| VIT_18s0001g02950 | 18 | 3042485 | 3045575 |
| VIT_18s0001g02970 | 18 | 3057427 | 3063862 |
| VIT_18s0001g03000 | 18 | 3064354 | 3065865 |
| VIT_18s0001g03020 | 18 | 3069509 | 3079236 |
| VIT_18s0001g03060 | 18 | 3084053 | 3095775 |
| VIT_18s0001g03080 | 18 | 3098022 | 3101041 |
| VIT_18s0001g03090 | 18 | 3102029 | 3103369 |
| VIT_18s0001g03100 | 18 | 3109637 | 3133804 |
| VIT_18s0001g03110 | 18 | 3134535 | 3146362 |
| VIT_18s0001g03130 | 18 | 3149249 | 3169173 |
| VIT_18s0001g03150 | 18 | 3170440 | 3174962 |
| VIT_18s0001g03160 | 18 | 3176998 | 3180139 |
| VIT_18s0001g03170 | 18 | 3180624 | 3188491 |
| VIT_18s0001g03180 | 18 | 3190049 | 3192531 |
| VIT_18s0001g03190 | 18 | 3197133 | 3198315 |
| VIT_18s0001g03200 | 18 | 3198754 | 3201304 |
| VIT_18s0001g03220 | 18 | 3206237 | 3206969 |
| VIT_18s0001g03230 | 18 | 3208995 | 3214247 |
| VIT_18s0001g03240 | 18 | 3219544 | 3220699 |
| VIT_18s0001g03250 | 18 | 3222134 | 3232291 |
| VIT_18s0001g03270 | 18 | 3233803 | 3234804 |
| VIT_18s0001g03290 | 18 | 3244469 | 3246948 |
| VIT_18s0001g03300 | 18 | 3249246 | 3253146 |
| VIT_18s0001g03310 | 18 | 3254531 | 3256064 |
| VIT_18s0001g03330 | 18 | 3261422 | 3268381 |
| VIT_18s0001g03370 | 18 | 3282634 | 3291512 |
| VIT_18s0001g03390 | 18 | 3298924 | 3306685 |
| VIT_18s0001g03420 | 18 | 3307440 | 3307550 |
| VIT_18s0001g03430 | 18 | 3309917 | 3311940 |
|  |  |  |  |


| VIT_18s0001g03440 | 18 | 3313991 | 3315438 |
| :--- | :--- | :--- | :--- |
| VIT_18s0001g03450 | 18 | 3320975 | 3328281 |
| VIT_18s0001g03470 | 18 | 3338337 | 3340959 |
| VIT_18s0001g03490 | 18 | 3349225 | 3350592 |
| VIT_18s0001g03510 | 18 | 3354104 | 3359003 |
| VIT_18s0001g03520 | 18 | 3359393 | 3369207 |
| VIT_18s0001g03540 | 18 | 3380455 | 3383288 |
| VIT_18s0001g03570 | 18 | 3387900 | 3389335 |
| VIT_18s0001g03580 | 18 | 3389546 | 3393993 |
| VIT_18s0001g03610 | 18 | 3401893 | 3411144 |
| VIT_18s0001g03630 | 18 | 3412345 | 3414417 |
| VIT_18s0001g03640 | 18 | 3417193 | 3418012 |
| VIT_18s0001g03650 | 18 | 3418650 | 3419295 |
| VIT_18s0001g03670 | 18 | 3422279 | 3424214 |
| VIT_18s0001g03680 | 18 | 3433101 | 3445210 |
| VIT_18s0001g03720 | 18 | 3445760 | 3445939 |
| VIT_18s0001g03730 | 18 | 3445960 | 3490992 |
| VIT_18s0001g03760 | 18 | 3492878 | 3505322 |
| VIT_18s0001g03790 | 18 | 3515123 | 3530220 |
| VIT_18s0001g03820 | 18 | 3536802 | 3542078 |
| VIT_18s0001g03840 | 18 | 3555522 | 3560233 |
| VIT_18s0001g03850 | 18 | 3562597 | 3563546 |
| VIT_18s0001g03870 | 18 | 3564229 | 3564690 |
| VIT_18s0001g03880 | 18 | 3565751 | 3567556 |
| VIT_18s0001g03910 | 18 | 3577992 | 3582255 |
| VIT_18s0001g03930 | 18 | 3585471 | 3585895 |
| VIT_18s0001g03940 | 18 | 3586978 | 3587567 |
| VIT_18s0001g03950 | 18 | 3587990 | 3596239 |
| VIT_18s0001g03960 | 18 | 3596240 | 3606771 |
| VIT_18s0001g03990 | 18 | 3638981 | 3646405 |
| VIT_18s0001g04010 | 18 | 3648331 | 3660863 |
| VIT_18s0001g04040 | 18 | 3668613 | 3697492 |
| VIT_18s0001g04100 | 18 | 3700504 | 3718045 |
| VIT_18s0001g04130 | 18 | 3720073 | 3720771 |
| VIT_18s0001g04140 | 18 | 3725171 | 3732449 |
| VIT_18s0001g04150 | 18 | 3732776 | 3738425 |
| VIT |  |  |  |

No hit
Glycine-rich protein
Flavonol synthase
Flavonol synthase
flavonol synthase XM 002284374.1 Camphor resistance CrcB Auxin transporter protein 4

Thaumatin ATLP-1
Ubiquitin-fold modifier 1 precursor Auxin-independent growth promoter
Pentatricopeptide (PPR) repeat-containing protein
No hit
No hit
Zinc finger ( C 2 H 2 type) family
Protein kinase
No hit
SET Domain group 37
Holocarboxylase synthetase 1 (HCS1)
Oxysterol binding protein
Phosphatidylinositol-4-phosphate 5-kinase 1
TEL2 (Terminal EAR1-like 2)
Ribosomal protein S10 30S
No hit
Polcalcin
Nitrate reductase 2 (NR2)
No hit
No hit
Pm27 protein
Mekk1
Stearoyl-ACP desaturase
Salt tolerant protein
PHD finger transcription factor Histone acetyl transferase HAM1

Unknown
Vacuolar protein sorting 9 (VPS9)
Avr9 elicitor response protein

| VIT_18s0001g04160 | 18 | 3741241 | 3745559 |
| :--- | :--- | :--- | :--- |
| VIT_18s0001g04180 | 18 | 3749619 | 3757678 |
| VIT_18s0001g04190 | 18 | 3759293 | 3763520 |
| VIT_18s0001g04270 | 18 | 3791368 | 3794765 |
| VIT_18s0001g04290 | 18 | 3798810 | 3805936 |
| VIT_18s0001g04340 | 18 | 3822948 | 3829597 |
| VIT_18s0001g04380 | 18 | 3831159 | 3832784 |
| VIT_18s0001g04400 | 18 | 3842548 | 3845101 |
| VIT_18s0001g04420 | 18 | 3846995 | 3851906 |
| VIT_18s0001g04440 | 18 | 3852065 | 3861217 |
| VIT_18s0001g04470 | 18 | 3861987 | 3867762 |
| VIT_18s0001g04500 | 18 | 3875315 | 3878509 |
| VIT_18s0001g04520 | 18 | 3879307 | 3882849 |
| VIT_18s0001g04540 | 18 | 3883733 | 3898683 |
| VIT_18s0001g04580 | 18 | 3900400 | 3905806 |
| VIT_18s0001g04590 | 18 | 3906073 | 3907918 |
| VIT_18s0001g04600 | 18 | 3913481 | 3917327 |
| VIT_18s0001g04610 | 18 | 3918203 | 3923636 |
| VIT_18s0001g04630 | 18 | 3924182 | 3930381 |
| VIT_18s0001g04640 | 18 | 3931182 | 3931761 |
| VIT_18s0001g04660 | 18 | 3935472 | 3935878 |
| VIT_18s0001g04680 | 18 | 3938582 | 3956444 |
| VIT_18s0001g04700 | 18 | 3956796 | 3960644 |
| VIT_18s0001g04730 | 18 | 3963413 | 3965554 |
| VIT_18s0001g04750 | 18 | 3967437 | 3969192 |
| VIT_18s0001g04760 | 18 | 3976114 | 3980165 |
| VIT_18s0001g04770 | 18 | 3980567 | 3987316 |
| VIT_18s0001g04790 | 18 | 3994178 | 4000375 |
| VIT_18s0001g04800 | 18 | 4003656 | 4004282 |
| VIT_18s0001g04810 | 18 | 4014840 | 4015257 |
| VIT_18s0001g04850 | 18 | 4033185 | 4034494 |
| VIT_18s0001g04860 | 18 | 4035264 | 4040608 |
| VIT_18s0001g04880 | 18 | 4041678 | 4046983 |
| VIT_18s0001g04890 | 18 | 4048925 | 4052950 |
| VIT_18s0001g04910 | 18 | 4062087 | 4068935 |
| VIT_18s0001g04920 | 18 | 4068936 | 4069711 |
|  |  |  |  |

Translation initiation factor eIF-5
Auxin response factor ARF17
Co-chaperone grpE

Pentatricopeptide (PPR) repeat-containing
No hit
Glycine hydroxymethyltransferase
Unknown protein
Rhomboid family KOM (kompeito) Phosphopantothenate--cysteine ligase

Co-chaperone protein DnaJ
Basic Leucine Zipper Transcription Factor (VvbZIP42)
Enhanced EM level EEL (VvABF-2), Basic Leucine Zipper Transcription Factor (VvbZIP43)
No hit
Unknown protein Unknown protein

Binding
Glutaredoxin Unknown protein
Aspartate aminotransferase
No hit
No hit
RPG related protein 1 RR1 GP5 ubiquitin-like
Kelch repeat-containing F-box protein Acetylcholinesterase
Ribosomal protein L30 (RPL30B) 60S
Apoptotic chromatin condensation inducer 1 ACIN1
Aspartic Protease (VvAP42)
Hyperosmotically inducible periplasmic protein putative MADS-box type delta 1b (VviMADSD1b)

Unknown
Aspartate transaminase.
Unknown protein
Low affinity sulphate transporter
Sulfate transporter 1.3
No hit

| VIT_18s0001g04930 | 18 | 4070013 | 4071952 |
| :--- | :--- | :--- | :--- |
| VIT_18s0001g04940 | 18 | 4072111 | 4072857 |
| VIT_18s0001g04960 | 18 | 4076661 | 4078865 |
| VIT_18s0001g04970 | 18 | 4087210 | 4096302 |
| VIT_18s0001g04980 | 18 | 4097931 | 4113803 |
| VIT_18s0001g05020 | 18 | 4122072 | 4123281 |
| VIT_18s0001g05030 | 18 | 4130541 | 4134290 |
| VIT_18s0001g05040 | 18 | 4138118 | 4140727 |
| VIT_18s0001g05060 | 18 | 4143615 | 4147853 |
| VIT_18s0001g05080 | 18 | 4148968 | 4155129 |
| VIT_18s0001g05100 | 18 | 4156671 | 4159130 |
| VIT_18s0001g05110 | 18 | 4159695 | 4167493 |
| VIT_18s0001g05130 | 18 | 4176691 | 4179995 |
| VIT_18s0001g05160 | 18 | 4181102 | 4184913 |
| VIT_18s0001g05180 | 18 | 4185500 | 4189977 |
| VIT_18s0001g05220 | 18 | 4205002 | 4212978 |
| VIT_18s0001g05250 | 18 | 4220268 | 4222313 |
| VIT_18s0001g05270 | 18 | 4225485 | 4241449 |
| VIT_18s0001g05300 | 18 | 4254678 | 4257480 |
| VIT_18s0001g05310 | 18 | 4268207 | 4269413 |
| VIT_18s0001g05330 | 18 | 4272391 | 4273949 |
| VIT_18s0001g05340 | 18 | 4274062 | 4275436 |
| VIT_18s0001g05370 | 18 | 4279252 | 4280705 |
| VIT_18s0001g05380 | 18 | 4284177 | 4287720 |
| VIT_18s0001g05400 | 18 | 4299165 | 4300807 |
| VIT_18s0001g05420 | 18 | 4306846 | 4309947 |
| VIT_18s0001g05440 | 18 | 4310333 | 4319306 |
| VIT_18s0001g05480 | 18 | 4326127 | 4329358 |
| VIT_18s0001g05490 | 18 | 4329359 | 4329584 |
| VIT_18s0001g05500 | 18 | 4331692 | 4337441 |
| VIT_18s0001g05530 | 18 | 4338743 | 4341637 |
| VIT_18s0001g05550 | 18 | 4349906 | 4352613 |
| VIT_18s0001g05570 | 18 | 4363673 | 4368991 |
| VIT_18s0001g05580 | 18 | 4373859 | 4374826 |
| VIT_18s0001g05590 | 18 | 4377588 | 4386070 |
| VIT_18s0001g05600 | 18 | 4390485 | 4398813 |
| VIT |  |  |  |

Sulfate transporter 1.3
No hit
Sulfate transporter 1.2
TRNA modification GTPase trmE
Acetyl-CoA carboxylase 2 (ACC2)
Senescence-associated protein
Glucan 1,3-beta-glucosidase precursor Pollen Ole e 1 allergen and extensin
2,3-bisphosphoglycerate-dependent phosphoglycerate mutase
SEC14 cytosolic factor, putative
No hit
DNA2-NAM7 helicase
Unknown protein
Glycosyl hydrolase family 3 protein
Beta-D-xylosidase
WD-40 repeat
Dehydration Responsive Element-Binding Transcription Factor (VvDREB27)
ERGIC and golgi 3
Trehalose-6-phosphate phosphatase
F-box domain containing protein
No hit
Transcription termination factor mitochondrial mTERF
Unknown protein
TTL1 (tetratricopetide-repeat thioredoxin-like 1)
Unknown
UNE2 (unfertilized embryo sac 2); carbohydrate transporter
Methyltransferase type 11
Unknown protein
No hit
PUMILIO 7 (APUM7)
Annexin 1 (ANN1)
Splicing factor, arginine/serine-rich 2
Hexose transporter HT2
Unknown protein
CCR4-NOT transcription complex subunit 6
Synaptonemal complex protein 1 (ZYP1A)

| VIT_18s0001g05620 | 18 | 4399508 | 4405646 |
| :--- | :--- | :--- | :--- |
| VIT_18s0001g05640 | 18 | 4407516 | 4416666 |
| VIT_18s0001g05670 | 18 | 4425461 | 4426613 |
| VIT_18s0001g05680 $^{18}$ | 18 | 4433417 | 4440032 |
| VIT_18s0001g05690 | 18 | 4442502 | 4445540 |
| VIT_18s0001g05710 | 18 | 4445541 | 4452656 |
| VIT_18s0001g05720 | 18 | 4453327 | 4457820 |
| VIT_18s0001g05730 | 18 | 4458540 | 4462662 |
| VIT_18s0001g05740 | 18 | 4465943 | 4475083 |
| VIT_18s0001g05780 | 18 | 4483440 | 4486850 |
| VIT_18s0001g05790 | 18 | 4491794 | 4493362 |
| VIT_18s0001g05800 | 18 | 4494490 | 4503957 |
| VIT_18s0001g05840 | 18 | 4520619 | 4529938 |
| VIT_18s0001g05860 | 18 | 4531169 | 4532101 |
| VIT_18s0001g05870 | 18 | 4535173 | 4537082 |
| VIT_18s0001g05900 | 18 | 4544137 | 4544831 |
| VIT_18s0001g05910 | 18 | 4550436 | 4552401 |
| VIT_18s0001g05950 | 18 | 4564102 | 4565743 |
| VIT_18s0001g05970 | 18 | 4566745 | 4574723 |
| VIT_18s0001g05990 | 18 | 4574889 | 4586041 |
| VIT_18s0001g06000 | 18 | 4586571 | 4589855 |
| VIT_18s0001g06020 | 18 | 4594155 | 4595268 |
| VIT_18s0001g06030 | 18 | 4597940 | 4599044 |
| VIT_18s0001g06040 | 18 | 4600121 | 4600746 |
| VIT_18s0001g06050 | 18 | 4600747 | 4602556 |
| VIT_18s0001g06060 | 18 | 4605365 | 4607009 |
| VIT_18s0001g06080 | 18 | 4610459 | 4612060 |
| VIT_18s0001g06090 | 18 | 4613633 | 4615316 |
| VIT_18s0001g06120 | 18 | 4629508 | 4631173 |
| VIT_18s0001g06130 | 18 | 4631192 | 4631628 |
| VIT_18s0001g06140 | 18 | 4643431 | 4644462 |
| VIT_18s0001g06150 | 18 | 4652464 | 4653648 |
| VIT_18s0001g06170 | 18 | 4658419 | 4659571 |
| VIT_18s0001g06180 | 18 | 4659693 | 4663192 |
| VIT_18s0001g06200 | 18 | 4674500 | 4678247 |
| VIT_18s0001g06220 | 18 | 4684266 | 4686537 |
|  |  |  |  |

[^1]| VIT_18s0001g06250 | 18 | 4700787 | 4704775 |
| :--- | :--- | :--- | :--- |
| VIT_18s0001g06270 | 18 | 4705961 | 4722205 |
| VIT_18s0001g06290 | 18 | 4724387 | 4727239 |
| VIT_18s0001g06300 | 18 | 4728333 | 4729667 |
| VIT_18s0001g06310 | 18 | 4733340 | 4737514 |
| VIT_18s001g06320 $^{18}$ | 18 | 4738323 | 4739992 |
| VIT_18s0001g06330 | 18 | 4742712 | 4744730 |
| VIT_18s0001g06350 | 18 | 4751168 | 4752609 |
| VIT_18s0001g06360 | 18 | 4760150 | 4762878 |
| VIT_18s0001g06370 | 18 | 4765814 | 4772318 |
| VIT_18s0001g06390 | 18 | 4773069 | 4780063 |
| VIT_18s0001g06400 | 18 | 4788251 | 4789021 |
| VIT_18s0001g06410 | 18 | 4791622 | 4794578 |
| VIT_18s0001g06420 | 18 | 4796509 | 4805339 |
| VIT_18s0001g06430 | 18 | 4806981 | 4808947 |
| VIT_18s0001g06440 | 18 | 4810193 | 4815522 |
| VIT_18s0001g06460 | 18 | 4815751 | 4855137 |
| VIT_18s0001g06500 | 18 | 4859577 | 4898877 |
| VIT_18s0001g06520 | 18 | 4900037 | 4954270 |
| VIT_18s001g06560 | 18 | 4962807 | 4963515 |
| VIT_18s0001g06580 | 18 | 4966021 | 4966915 |
| VIT_18s0001g06590 | 18 | 4970021 | 4973908 |
| VIT_18s0001g06600 | 18 | 4974906 | 4979125 |
| VIT_18s0001g06610 | 18 | 4982073 | 4983574 |
| VIT_18s0001g06630 | 18 | 4993758 | 4999430 |
| VIT_18s0001g06640 | 18 | 5000284 | 5003505 |
| VIT_18s001g06650 | 18 | 5009725 | 5011125 |
| VIT_18s0001g06670 | 18 | 5020431 | 5021751 |
| VIT_18s0001g06690 | 18 | 5029717 | 5032277 |
| VIT_18s0001g06710 | 18 | 5035063 | 5039911 |
| VIT_18s0001g06760 | 18 | 5054380 | 5055720 |
| VIT_18s0001g06770 | 18 | 5058487 | 5059001 |
| VIT_18s0001g06790 | 18 | 5062066 | 5066242 |
| VIT_18s0001g06820 | 18 | 5077418 | 5081480 |
| VIT_18s0001g06840 | 18 | 5092164 | 5094226 |
| VIT_18s0001g06850 | 18 | 5102796 | 5105019 |
|  |  |  |  |

3-deoxy-D-arabino-heptulosonate 7-phosphate synthase Transcription factor jumonji ( jmjC ) domain-containing protein

Ribosomal protein S6 (RPS6B) 40S
Purine permease 1 PUP1
SnRK2-8
Cupin, RmlC-type
14-3-3 protein GF14 omega (GRF2)
VQ motif-containing protein
Alcohol dehydrogenase
L-ascorbate peroxidase, chloroplast
Auxin-independent growth promoter
No hit
Ribosomal protein 60S
No hit
Homeobox-leucine zipper protein ATHB-6
No hit
Queuine tRNA-ribosyltransferase
Unknown protein
Isoamylase protein.
No hit
Plastocyanin domain-containing protein
Pigment defective 322
Unknown protein
Plastid-targeted protein 2
No hit
Zinc finger (C3HC4-type ring finger) basic helix-loop-helix (bHLH) family

Ring-H2 finger protein ATL1N
Gibberellin 20 oxidase 2
Oxidoreductase, 2OG-Fe(II) oxygenase
Clathrin assembly protein 16
No hit
Protein TRANSPARENT TESTA 12 (DDTFR18) MATE efflux family protein ripening responsive

Peroxidase GvPx2b class III
Peroxidase GvPx2b class III

|  | VIT_18s0001g06890 | 18 | 5121954 | 5123595 | Peroxidase GvPx2b, class III [Vitis vinifera] |
| :---: | :---: | :---: | :---: | :---: | :---: |
|  | VIT_18s0001g06910 | 18 | 5139237 | 5140617 | Purine permease 1 PUP1 |
|  | VIT_18s0001g06940 | 18 | 5163243 | 5164681 | Purine permease 1 (PUP1) |
|  | VIT_18s0001g06950 | 18 | 5169728 | 5171215 | Purine permease 1 (PUP1) |
|  | VIT_18s0001g06970 | 18 | 5184425 | 5218375 | DnaJ homolog, subfamily A, member 3 |
|  | VIT_18s0001g06980 | 18 | 5220072 | 5221115 | Pentatricopeptide repeat-containing protein |
|  | VIT_18s0001g07000 | 18 | 5240693 | 5244484 | Pentatricopeptide (PPR) repeat-containing protein |
|  | VIT_18s0001g07010 | 18 | 5244635 | 5245588 | GLUTATHIONE S-TRANSFERASE TAU 8 |
|  | VIT_18s0001g07020 | 18 | 5247639 | 5250783 | No hit |
|  | VIT_18s0001g07060 | 18 | 5256010 | 5263653 | Phosphoribosylaminoimidazole-succinocarboxamide synthase. |
|  | VIT_18s0001g07070 | 18 | 5266578 | 5271872 | Phosphoenolpyruvate carboxylase kinase 2 (PEPKR2) |
|  | VIT_18s0001g07080 | 18 | 5275135 | 5280005 | Protein transport protein SFT1 |
|  | VIT_18s0001g07090 | 18 | 5290562 | 5293561 | Unknown protein |
|  | VIT_18s0001g07100 | 18 | 5294004 | 5304799 | Pyridoxamine 5'-phosphate oxidase |
|  | VIT_18s0001g07110 | 18 | 5310450 | 5311506 | No hit |
|  | VIT_18s0001g07120 | 18 | 5311507 | 5314031 | Transport inhibitor response 1 protein |
|  | VIT_18s0001g07130 | 18 | 5323899 | 5335911 | GEM-like 1 |
|  | VIT_18s0001g07140 | 18 | 5343590 | 5344691 | Zinc Finger Homeodomain Transcription Factor (VvZHD10) |
| pheno_18_2 | VIT_18s0001g12820 | 18 | 10924591 | 10933193 | Dihydroflavonol 4-reductase |
|  | VIT_18s0001g12830 | 18 | 10937041 | 10940329 | 1,4-beta-mannan endohydrolase |
|  | VIT_18s0001g12840 | 18 | 10940330 | 10945165 | ADP-glucose pyrophosphorylase large subunit CagpL2 |
|  | VIT_18s0001g12850 | 18 | 10951026 | 10951788 | Ribosomal protein L31 |
|  | VIT_18s0001g12860 | 18 | 10961548 | 10963014 | Unknown protein |
|  | VIT_-18s0001g12870 | 18 | 10968173 | 10971522 | CYP722A1 |
|  | VIT_18s0001g12880 | 18 | 10974280 | 10990318 | Jasmonate O-methyltransferase |
|  | VIT_18s0001g12890 | 18 | 10990319 | 10995000 | Jasmonate O-methyltransferase |
|  | VIT_18s0001g12900 | 18 | 10999096 | 11003121 | S-adenosyl-L-methionine:salicylic acid carboxyl methyltransferase |
|  | VIT_18s0001g12910 | 18 | 11011979 | 11016761 | Inositol-pentakisphosphate 2-kinase 1 ATIPK1 |
|  | VIT_18s0001g12920 | 18 | 11018628 | 11019094 | No hit |
|  | VIT_18s0001g12930 | 18 | 11019643 | 11020221 | Heavy-metal-associated domain-containing protein |
|  | VIT_18s0001g12940 | 18 | 11025678 | 11029403 | Dihydrofolate reductase-thymidylate synthase |
|  | VIT_18s0001g12950 | 18 | 11035878 | 11042058 | SEC14 cytosolic factor |
|  | VIT_18s0001g12960 | 18 | 11043696 | 11049454 | Wound-responsive protein |
|  | VIT_18s0001g12970 | 18 | 11056806 | 11060964 | Ethylene-responsive protein |
|  | VIT_18s0001g12980 | 18 | 11070674 | 11090733 | Cyclin-dependent protein kinase CYCT1;4 |
|  | VIT_-18s0001g12990 | 18 | 11099778 | 11102242 | Anthranilate N-benzoyltransferase protein 1 |


| VIT_18s0001g13000 | 18 | 11105764 | 11116980 |
| :--- | :--- | :--- | :--- |
| VIT_18s0001g13010 | 18 | 11126023 | 11129236 |
| VIT_18s0001g13020 | 18 | 11137583 | 11137919 |
| VIT_18s0001g13030 | 18 | 11137920 | 11138195 |
| VIT_18s0001g13040 | 18 | 11138196 | 11139020 |
| VIT_18s0001g13050 | 18 | 11143793 | 11148102 |
| VIT_18s0001g13060 | 18 | 11153980 | 11161250 |
| VIT_18s0001g13070 | 18 | 11161718 | 11165469 |
| VIT_18s0001g13080 | 18 | 11166531 | 11167946 |
| VIT_18s0001g13090 | 18 | 11169585 | 11174073 |
| VIT_18s0001g13100 | 18 | 11175122 | 11179283 |
| VIT_18s0001g13110 | 18 | 11184119 | 11185662 |
| VIT_18s0001g13120 | 18 | 11188323 | 11195903 |
| VIT_18s0001g13130 | 18 | 11196526 | 11198302 |
| VIT_18s0001g13140 | 18 | 11199295 | 11200843 |
| VIT_18s0001g13150 | 18 | 11201495 | 11206144 |
| VIT_18s0001g13160 | 18 | 11209086 | 11221435 |
| VIT_18s0001g13180 | 18 | 11241300 | 11245481 |
| VIT_18s0001g13190 | 18 | 11247397 | 11252857 |
| VIT_18s0001g13200 | 18 | 11256653 | 11261569 |
| VIT_18s0001g13210 | 18 | 11268578 | 11271421 |
| VIT_18s0001g13220 | 18 | 11275604 | 11278396 |
| VIT_18s0001g13230 | 18 | 11286706 | 11293739 |
| VIT_18s0001g13240 | 18 | 11293740 | 11296751 |
| VIT_18s0001g13250 | 18 | 11297040 | 11299715 |
| VIT_18s0001g13260 | 18 | 11307834 | 11309051 |
| VIT_18s0001g13270 | 18 | 11311350 | 11316053 |
| VIT_18s0001g13280 | 18 | 11318782 | 11330916 |
| VIT_18s0001g13290 | 18 | 11331768 | 11334686 |
| VIT_18s0001g13300 | 18 | 11335834 | 11336703 |
| VIT_18s0001g13310 | 18 | 11338330 | 11346082 |
| VIT_18s0001g13320 | 18 | 11347342 | 11348103 |
| VIT_18s0001g13330 | 18 | 11348587 | 11353628 |
| VIT_18s0001g13340 | 18 | 11355046 | 11359625 |
| VIT_18s0001g13350 | 18 | 11374221 | 11381621 |
| VIT_18s0001g13360 | 18 | 11384285 | 11387145 |
|  |  |  |  |

Adaptor-related protein complex AP-4, mu 1 Mitogen-activated Protein Kinase (VvMPK11)

$$
\begin{aligned}
& \text { No hit } \\
& \text { No hit }
\end{aligned}
$$

Basic Leucine Zipper Transcription Factor (VvbZIP47)
Protein translocase Tic20 C 3 H 2 C 3 ring-finger protein NAD-dependent epimerase/dehydratase

Unknown protein
Proteasome 26S AAA-ATPase subunit (RPT4)
Glycosyl transferase family 1 protein
Peroxidase
GTP-binding protein hflX BEL1-like homeodomain 10

No hit
BEL1 homeotic protein 3
Tetrapyrrole methylase
Ubiquitin-conjugating enzyme E2 W
MAPK (MPK9)
Cytokinin dehydrogenase 5 precursor
Lectin jacalin
Pentatricopeptide (PPR) repeat-containing protein Beta-galactosidase BG1 [Vitis vinifera]

Thioredoxin H-type 1
Thioredoxin H-type 1
ATP-dependent protease La (LON) domain-containing protein
Papain cysteine peptidase XBCP3
RanBPM
GTP1/OBG family
Unknown
GTP1/OBG family protein
ERF/AP2 Gene Family (VvERF039),Dehydration Responsive Element-Binding Transcription Factor (VvDREB29) Purple acid phosphatase 32- ATPAP32/PAP32 Purple acid phosphatase 32- ATPAP32/PAP32
Proton-dependent oligopeptide transport (POT) family protein
Nodulin MtN21 family

| VIT_18s0001g13370 | 18 | 11397795 | 11406180 |
| :--- | :--- | :--- | :--- |
| VIT_18s0001g13380 | 18 | 11409762 | 11411534 |
| VIT_18s0001g13390 | 18 | 11419071 | 11420557 |
| VIT_18s0001g13400 | 18 | 11425591 | 11427332 |
| VIT_18s0001g13410 | 18 | 11429258 | 11475180 |
| VIT_18s0001g13420 | 18 | 11486984 | 11488357 |
| VIT_18s0001g13430 | 18 | 11496666 | 11502715 |
| VIT_18s0001g13440 | 18 | 11503598 | 11504539 |
| VIT_18s0001g13450 | 18 | 11505093 | 11505323 |
| VIT_18s0001g13460 | 18 | 11506606 | 11512370 |
| VIT_18s0001g13490 | 18 | 11525758 | 11527691 |
| VIT_18s0001g13500 | 18 | 11528060 | 11539323 |
| VIT_18s0001g13510 | 18 | 11539928 | 11541250 |
| VIT_18s0001g13520 | 18 | 11546677 | 11548061 |
| VIT_18s0001g13530 | 18 | 11552151 | 11558455 |
| VIT_18s0001g13540 | 18 | 11559472 | 11560071 |
| VIT_18s0001g13550 | 18 | 11561321 | 11567816 |
| VIT_18s0001g13560 | 18 | 11577367 | 11581885 |
| VIT_18s0001g13570 | 18 | 11584918 | 11587466 |
| VIT_18s0001g13580 | 18 | 11600569 | 11607735 |
| VIT_18s0001g13590 | 18 | 11608623 | 11612061 |
| VIT_18s0001g13600 | 18 | 11612878 | 11613828 |
| VIT_18s0001g13610 | 18 | 11613829 | 11620092 |
| VIT_18s0001g13620 | 18 | 11630815 | 11630958 |
| VIT_18s0001g13630 | 18 | 11631001 | 11633191 |
| VIT_18s0001g13640 | 18 | 11636802 | 11640737 |
| VIT_18s0001g13650 | 18 | 11640899 | 11647357 |
| VIT_18s0001g13660 | 18 | 11656750 | 11691457 |
| VIT_18s0001g13670 | 18 | 11698060 | 11699721 |
| VIT_18s0001g13680 | 18 | 11710329 | 11711606 |
| VIT_18s0001g13690 | 18 | 11723026 | 11723485 |
| VIT_18s0001g13700 | 18 | 11726587 | 11729541 |
| VIT_18s0001g13710 | 18 | 11729964 | 11734394 |
| VIT_18s0001g13720 | 18 | 11734395 | 11741936 |
| VIT_18s0001g13730 | 18 | 11741937 | 11743502 |
| VIT 18s0001g13740 | 18 | 11745491 | 11746246 |
|  |  |  |  |

Transcription initiation factor TFIIF beta subunit (TFIIF-beta)
Papain cysteine proteinase isoform I
Papain cysteine proteinase isoform II
Papain cysteine proteinase isoform I
V-type $\mathrm{H}+$-transporting ATPase subunit I
SHI-related sequence 5
Coatomer alpha subunit
SLAH1 (SLAC1 homologue 1)
SLAH1 (SLAC1 homologue 1)
putative MADS-box Apetala 3a (VviAP3a)
Maturase K
forkhead-associated domain-containing protein
Her2-p6
Zinc finger (B-box type)
SH3 domain-containing protein 2
Glycine-rich protein
Zinc knuckle (CCHC-type) family Unknown protein
Calcium ion binding protein
Kinesin motor protein
Leucine-rich repeat protein kinase

## No hit

Abhydrolase domain-containing protein 5
No hit
V-type $\mathrm{H}+$-transporting ATPase 16 kDa proteolipid subunit Amine oxidase Amine oxidase
EMB2757/TAN (embryo defective 2757)
Glyoxal oxidase-related
Histone H3
No hit
Cell growth defect factor -2 BPC6/BBR/BPC6/BPC6 BPC6/BBR/BPC6/BPC6 ВРС6/BBR/BPC6/BPC6
Basic Leucine Zipper Transcription Factor (VvbZIP48)

| VIT_18s0001g13750 | 18 | 11750154 | 11761059 |
| :--- | :--- | :--- | :--- |
| VIT_18s0001g13760 | 18 | 11761760 | 11763067 |
| VIT_18s0001g13770 | 18 | 11767560 | 11769317 |
| VIT_18s0001g13780 | 18 | 11774368 | 11775896 |
| VIT_18s0001g13790 | 18 | 11779263 | 11781156 |
| VIT_18s0001g13800 | 18 | 11784344 | 11786071 |
| VIT_18s0001g13810 | 18 | 11789356 | 11791909 |
| VIT_18s0001g13820 | 18 | 11794593 | 11796161 |
| VIT_18s0001g13830 | 18 | 11801940 | 11803508 |
| VIT_18s0001g13850 | 18 | 11832431 | 11833905 |
| VIT_18s0001g13870 | 18 | 11862607 | 11871541 |
| VIT_18s0001g13880 | 18 | 11876078 | 11892231 |
| VIT_18s0001g13890 | 18 | 11893711 | 11897887 |
| VIT_18s0001g13900 | 18 | 11898378 | 11909253 |
| VIT_18s0001g13910 | 18 | 11909699 | 11914734 |
| VIT_18s0001g13920 | 18 | 11917645 | 11919060 |
| VIT_18s0001g13930 | 18 | 11920505 | 11928962 |
| VIT_18s0001g13940 | 18 | 11933806 | 11936210 |
| VIT_18s0001g13950 | 18 | 11940712 | 11949407 |
| VIT_18s0001g13960 | 18 | 11959623 | 11960345 |
| VIT_18s0001g13970 | 18 | 11974983 | 11980862 |
| VIT_18s0001g13980 | 18 | 11987475 | 11987922 |
| VIT_18s0001g13990 | 18 | 11989161 | 11997757 |
| VIT_18s0001g14000 | 18 | 12002927 | 12003389 |
| VIT_18s0001g14010 | 18 | 12004532 | 12014082 |
| VIT_18s0001g14020 | 18 | 12032185 | 12069047 |
| VIT_18s0001g14030 | 18 | 12073618 | 12075722 |
| VIT_18s0001g14040 | 18 | 12086327 | 12090970 |
| VIT_18s0001g14060 | 18 | 12117278 | 12118772 |
| VIT_18s0001g14070 | 18 | 12133743 | 12135762 |
| VIT_18s0001g14080 | 18 | 12136487 | 12140512 |
| VIT_18s0001g14090 | 18 | 12141515 | 12144267 |
| VIT_18s0001g14100 | 18 | 12146146 | 12150550 |
| VIT_18s0001g14110 | 18 | 12151849 | 12158339 |
| VIT_18s0001g14120 | 18 | 12160472 | 12174229 |
| VIT_18s0001g14130 | 18 | 12179540 | 12181647 |
|  |  |  |  |

Regulator of chromosome condensation (RCC1)
CYP71AT2v2
Cytochrome P450, family 83 , subfamily B, polypeptide 1 Cytochrome P450, family 83, subfamily B, polypeptide 1 Cytochrome P450, family 83, subfamily B, polypeptide 1 CYP71AT2v2
Heat shock protein-related
Cytochrome P450, family 83, subfamily B, polypeptide 1 Cytochrome P450, family 83 , subfamily B, polypeptide 1 Cytochrome P450, family 83, subfamily B, polypeptide 1

Calmodulin binding protein Zinc finger (CCCH-type) family protein Unknown protein
Unknown protein
Unknown protein
Unknown protein
Auxin response factor 5 (Transcription factor MONOPTEROS) Pentatricopeptide (PPR) repeat-containing

RNA polymerase Rpa43 subunit
SAUR_E
Myosin-related
Auxin responsive SAUR protein
No hit
Auxin-induced SAUR
Sulfate adenylyltransferase
$5^{\prime}-3$ ' exoribonuclease 2
Lysine decarboxylase
Endo-1,4-beta-glucanase
Transposon protein, CACTA, En/Spm sub-class
Unknown
Diphthine synthase (DPH5) Molecular chaperone DnaJ
ABA-responsive protein (HVA22)HVA22H Transcription factor E2F/dimerisation partner (TDP) E2F2 Translation initiation factor eIF-3 subunit 3 Zinc finger (C2H2 type) family

| VIT_18s0001g14140 | 18 | 12191289 | 12193271 |
| :--- | :--- | :--- | :--- |
| VIT_18s0001g14150 | 18 | 12197073 | 12197669 |
| VIT_18s0001g14160 | 18 | 12205115 | 12205631 |
| VIT_18s0001g14170 | 18 | 12212398 | 12215228 |
| VIT_18s0001g14180 | 18 | 12216225 | 12219253 |
| VIT_18s0001g14190 | 18 | 12223875 | 12228196 |
| VIT_18s0001g14200 | 18 | 12229450 | 12229914 |
| VIT_18s0001g14210 | 18 | 12230572 | 12231893 |
| VIT_18s0001g14230 | 18 | 12252526 | 12257027 |
| VIT_18s0001g14240 | 18 | 12257459 | 12261168 |
| VIT_18s0001g14250 | 18 | 12262636 | 12268135 |
| VIT_18s0001g14260 | 18 | 12273788 | 12276090 |
| VIT_18s0001g14270 | 18 | 12276091 | 12276597 |
| VIT_18s0001g14280 | 18 | 12281671 | 12286855 |
| VIT_18s0001g14290 | 18 | 12292329 | 12294516 |
| VIT_18s0001g14300 | 18 | 12299065 | 12301674 |
| VIT_18s0001g14310 | 18 | 12303077 | 12305027 |
| VIT_18s0001g14320 | 18 | 12306644 | 12309420 |
| VIT_18s0001g14330 | 18 | 12310077 | 12311494 |
| VIT_18s0001g14340 | 18 | 12313857 | 12316276 |
| VIT_18s0001g14350 | 18 | 12335581 | 12336788 |
| VIT_18s0001g14360 | 18 | 12337145 | 12340985 |
| VIT_18s0001g14370 | 18 | 12341546 | 12342790 |
| VIT_18s0001g14380 | 18 | 12343090 | 12350501 |
| VIT_18s0001g14390 | 18 | 12362497 | 12370721 |
| VIT_18s0001g14400 | 18 | 12377314 | 12378089 |
| VIT_18s0001g14410 | 18 | 12383680 | 12402362 |
| VIT_18s0001g14420 | 18 | 12403268 | 12404144 |
| VIT_18s0001g14430 | 18 | 12431386 | 12431715 |
| VIT_18s0001g14440 | 18 | 12432955 | 12439459 |
| VIT_18s0001g14450 | 18 | 12453766 | 12456583 |
| VIT_18s0001g14460 | 18 | 12457152 | 12463238 |
| VIT_18s0001g14470 | 18 | 12465257 | 12466480 |
| VIT_18s0001g14480 | 18 | 12468330 | 12470542 |
| VIT_18s0001g14490 | 18 | 12488747 | 12490747 |
| VIT_18s0001g14500 | 18 | 12501841 | 12507813 |
|  |  |  |  |

No hit
No hit
Heat-and acid-stable phosphoprotein
Cyclin-dependent protein kinase regulator CYCB2 4 Armadillo/beta-catenin repeat family protein

Unknown
No hit
Unknown protein
Hexokinase
Adaptor-related protein complex 2, sigma 1 sub
Unknown protein
No hit
Gibberellin-regulated protein 1 (GASA1)
ER lumen protein retaining receptor
Invertase-like protein
Cytomatrix protein
flavanone-3-hydroxylase 2 (F3H2) [Vitis vinifera]
Unknown protein
No hit
No hit
No hit
Tubulin beta-1 chain
No hit
Brain and reproductive organ-expressed protein
Transducin protein
Peptidyl-prolyl cis-trans isomerase ROC5 (rotamase CYP 5)
Unknown protein
Unknown
Unknown

## Molecular chaperone DnaJ

Ferredoxin:nadp+ Oxidoreductase PETH
Unknown protein
No hit
Thaumatin SCUTL1
Thaumatin
SHD (shepherd)

| VIT_18s0001g14510 | 18 | 12509258 | 12513293 |
| :--- | :--- | :--- | :--- |
| VIT_18s0001g14520 | 18 | 12526253 | 12526939 |
| VIT_18s0001g14530 | 18 | 12533609 | 12536412 |
| VIT_18s0001g14540 | 18 | 12545280 | 12566106 |
| VIT_18s0001g14550 | 18 | 12579175 | 12591899 |
| VIT_18s0001g14560 | 18 | 12608249 | 12608970 |
| VIT_18s0001g14580 | 18 | 12620422 | 12628019 |
| VIT_18s0001g14590 | 18 | 12634521 | 12635111 |
| VIT_18s0001g14600 | 18 | 12645229 | 12650755 |
| VIT_18s0001g14610 | 18 | 12668387 | 12671744 |
| VIT_18s0001g14620 | 18 | 12678950 | 12679433 |
| VIT_18s0001g14630 | 18 | 12682841 | 12691610 |
| VIT_18s0001g14640 | 18 | 12698089 | 12699502 |
| VIT_18s0001g14650 | 18 | 12699590 | 12707069 |
| VIT_18s0001g14660 | 18 | 12710695 | 12716340 |
| VIT_18s0001g14670 | 18 | 12719099 | 12724188 |
| VIT_18s0001g14680 | 18 | 12725219 | 12753021 |
| VIT_18s0001g14690 | 18 | 12763424 | 12766855 |
| VIT_18s0001g14700 | 18 | 12767403 | 12770616 |
| VIT_18s0001g14710 | 18 | 12772148 | 12773056 |
| VIT_18s0001g14720 | 18 | 12773057 | 12773368 |
| VIT_18s0001g14730 | 18 | 12776655 | 12781411 |
| VIT_18s0001g14740 | 18 | 12783608 | 12784580 |
| VIT_18s0001g14750 | 18 | 12787199 | 12809768 |
| VIT_18s0001g14760 | 18 | 12821158 | 12822999 |
| VIT_18s0001g14770 | 18 | 12823491 | 12840711 |
| VIT_18s0001g14780 | 18 | 12841218 | 12848820 |
| VIT_18s0001g14790 | 18 | 12853195 | 12855945 |
| VIT_18s0001g14800 | 18 | 12859139 | 12862545 |
| VIT_18s0001g14810 | 18 | 12863066 | 12871408 |
| VIT_18s0001g14840 | 18 | 12887405 | 12890179 |
| VIT_18s0001g14850 | 18 | 12893670 | 12898347 |
| VIT_18s0001g14860 | 18 | 12900823 | 12904890 |
| VIT_18s0001g14870 | 18 | 12908766 | 12912392 |
| VIT_18s0001g14880 | 18 | 12914221 | 12923327 |
| VIT_18s0001g14890 | 18 | 12936974 | 12937903 |
|  |  |  |  |

Ribosomal protein L28
Unknown protein
Ubiquitin-protein ligase CIP8 (COP1-interacting protein 8) DEAD-box ATP-dependent RNA helicase 28

Unknown protein
Unknown
Kinesin family member 18/19 ATSYTC/NTMC2T1.3/NTMC2TYPE1.3/SYTC Pentatricopeptide (PPR) repeat-containing protein

Clavata1 receptor kinase (CLV1)
Ribosomal protein S16
ATSYTB/NTMC2T1.2/NTMC2TYPE1.2/SYTB
No hit
DNA-directed RNA polymerase I subunit A12 ABC Transporter (VvMDR3 - VvABCB3) Rac-like GTP-binding protein RAC1

Vacuolar protein sorting 35
Protein kinase
Unknown
Ketol-acid reductoisomerase precursor
Ankyrin repeat family protein
Glutaredoxin
Protein binding
Zinc finger (FYVE type) VPS19 Lipase 3 (EXL3) family II extracellular Lipase 3 (EXL3) family II extracellular Lipase 3 (EXL3) family II extracellular Lipase 3 (EXL3) family II extracellular Lipase 3 (EXL3) family II extracellular Lipase 3 (EXL3) family II extracellular Nickel ion transporter Unknown protein

Subtilisin-type protease precursor Unknown protein
ABA-responsive element-binding protein 3 (AREB3), Basic Leucine Zipper Transcription Factor (VvbZIP49)

| VIT_18s0001g14900 | 18 | 12941612 | 12948377 |
| :--- | :--- | :--- | :--- |
| VIT_18s0001g14910 | 18 | 12954423 | 12956365 |
| VIT_18s0001g14920 | 18 | 12959804 | 12961455 |
| VIT_18s0001g14930 | 18 | 12964045 | 12967757 |
| VIT_18s0001g14940 | 18 | 12969264 | 12971832 |
| VIT_18s0001g14950 | 18 | 12978238 | 12979279 |
| VIT_18s0001g14960 | 18 | 12980373 | 12984313 |
| VIT_18s0001g14970 | 18 | 12985919 | 12987853 |
| VIT_18s0001g14980 | 18 | 12992480 | 12996562 |
| VIT_18s0001g14990 | 18 | 12997417 | 13013251 |
| VIT_18s0001g15000 | 18 | 13026931 | 13036065 |
| VIT_18s0001g15010 | 18 | 13037311 | 13042185 |
| VIT_18s0001g15020 | 18 | 13043063 | 13050564 |
| VIT_18s0001g15050 | 18 | 13073854 | 13087034 |
| VIT_18s0001g15060 | 18 | 13104442 | 13105749 |
| VIT_18s0001g15070 | 18 | 13107851 | 13109297 |
| VIT_18s0001g15080 | 18 | 13112575 | 13112793 |
| VIT_18s0001g15090 | 18 | 13133931 | 13152578 |
| VIT_18s0001g15100 | 18 | 13176556 | 13178357 |
| VIT_18s0001g15110 | 18 | 13195418 | 13195879 |
| VIT_18s0001g15120 | 18 | 13199060 | 13200303 |
| VIT_18s0001g15130 | 18 | 13201886 | 13203120 |
| VIT_18s0001g15140 | 18 | 13208994 | 13210315 |
| VIT_18s0001g15150 | 18 | 13231125 | 13232440 |
| VIT_18s0001g15160 | 18 | 13277786 | 13278346 |
| VIT_18s0001g15170 | 18 | 13278347 | 13278514 |
| VIT_18s0001g15180 | 18 | 13279241 | 13280607 |
| VIT_18s0001g15190 | 18 | 13303082 | 13304738 |
| VIT_18s0001g15200 | 18 | 13306083 | 13306887 |
| VIT_18s0001g15220 | 18 | 13312158 | 13313392 |
| VIT_18s0001g15230 | 18 | 13322054 | 13323283 |
| VIT_18s0001g15240 | 18 | 13365469 | 13366663 |
| VIT_18s0001g15250 | 18 | 13389043 | 13389957 |
| VIT_18s0001g15260 | 18 | 13397962 | 13399162 |
| VIT_18s0001g15270 | 18 | 13408234 | 13413047 |
| VIT_18s0001g15280 | 18 | 13418393 | 13429810 |
|  |  |  |  |

La domain-containing protein
Mannitol dehydrogenase
Unknown protein
Unknown protein
Pentatricopeptide (PPR) repeat-containing EMB2170 (embryo defective 2170) Pentatricopeptide (PPR) repeat-containing Pentatricopeptide (PPR) repeat-containing 3-methyl-2-oxobutanoate dehydrogenase

Dual-specific kinase DSK1
ACT domain containing protein (ACR4)
F-box and leucine-rich repeat protein 1 DnaJ homolog, subfamily A, member 4

Protein phosphatase 2C
No hit
No hit
No hit
RAB GTPase RAB18
flavin-containing monooxygenase, putative Flavin-containing monooxygenases

Unknown protein
WD40 repeat protein
Unknown
Unknown protein
S-adenosylmethionine sythetase 2
S-adenosylmethionine sythetase 2
No hit
flavin-containing monooxygenase, putative
SIK1 (serine/threonine kinase 1)
Unknown protein
Unknown
Blue (type 1) copper domain
Unknown
myb domain protein 52
SCL1 (scarecrow-like 1)
RNA exonuclease 1

| VIT_18s0001g15290 | 18 | 13430862 | 13443967 |
| :--- | :--- | :--- | :--- |
| VIT_18s0001g15300 | 18 | 13444139 | 13444288 |
| VIT_18s0001g15310 | 18 | 13444700 | 13445878 |
| VIT_18s0001g15320 | 18 | 13446395 | 13458670 |
| VIT_18s0001g15330 | 18 | 13476403 | 13478586 |
| VIT_18s0001g15340 | 18 | 13488810 | 13490580 |
| VIT_18s0001g15350 | 18 | 13493089 | 13494429 |
| VIT_18s0001g15360 | 18 | 13495083 | 13497819 |
| VIT_18s0001g15370 | 18 | 13498468 | 13504540 |
| VIT_18s0001g15380 | 18 | 13511653 | 13519868 |
| VIT_18s0001g15390 | 18 | 13521135 | 13522636 |
| VIT_18s0001g15400 | 18 | 13526007 | 13535458 |
| VIT_18s0001g15410 | 18 | 13539135 | 13541625 |
| VIT_18s0001g15420 | 18 | 13551359 | 13554243 |
| VIT_18s0001g15430 | 18 | 13563113 | 13563819 |
| VIT_18s0001g15450 | 18 | 13581959 | 13584888 |
| VIT_18s0001g15460 | 18 | 13616618 | 13618309 |
| VIT_18s0001g15470 | 18 | 13620924 | 13622663 |
| VIT_18s0001g15510 | 18 | 13646454 | 13647443 |
| VIT_18s0001g15520 | 18 | 13650247 | 13655312 |
| VIT_18s0001g15530 | 18 | 13662369 | 13665624 |
| VIT_18s0001g15540 | 18 | 13666500 | 13667021 |
| VIT_18s0001g15550 | 18 | 13667832 | 13671220 |
| VIT_18s0001g15560 | 18 | 13676072 | 13681500 |
| VIT_18s0001g15570 | 18 | 13682439 | 13700040 |
| VIT_18s0001g15580 | 18 | 13701036 | 13706265 |
| VIT_18s0001g15600 | 18 | 13720692 | 13730593 |
| VIT_18s0001g15610 | 18 | 13733152 | 13734200 |
| VIT_18s0001g15620 | 18 | 13735971 | 13739523 |
| VIT_18s0001g15630 | 18 | 13793649 | 13799974 |
| VIT_18s0001g15640 | 18 | 13803789 | 13805200 |
| VIT_18s0001g15650 | 18 | 13805201 | 13806648 |
| VIT_18s0001g15660 | 18 | 13810659 | 13811793 |
| VIT_18s0001g15670 | 18 | 13812058 | 13816979 |
| VIT_18s0001g15680 | 18 | 13833441 | 13835421 |
| VIT_18s0001g15690 | 18 | 13835422 | 13835790 |
|  |  |  |  |

Ribosome maturation protein SDO1
No hit

## Thioredoxin M-type

 SEU3B proteinNodulin MtN3 family
Pentatricopeptide (PPR) repeat-containing protein Pentatricopeptide (PPR) repeat-containing protein

Thylakoid lumenal 29.8 kDa protein Exostosin family protein No hit
Gaiacol peroxidase
KOW domain-containing transcription factor family protein
Alcohol dehydrogenase 1
Auxin efflux carrier protein 6
Alliin lyase
Alcohol dehydrogenase 3 Stearyl acyl carrier protein desaturase Leaf senescence related protein-like

Unknown
Leaf senescence protein
Pentatricopeptide (PPR) repeat-containing
No hit
Serine carboxypeptidase II
Unknown protein
Zinc finger (CCCH-type) family protein Glycogenin glucosyltransferase (glycogenin)

Vacuolar protein sorting 45
Unknown
Receptor kinase TRKe
MAP kinase activating protein
Pathogenesis-related
Pathogenesis related protein
Pathogen-related
Cytoplasm protein
Cellulase
Endo-1,4-beta-glucanase

| 18s0001g15700 | 18 | 13836500 | 13838302 |
| :---: | :---: | :---: | :---: |
| VIT_18s0001g15710 | 18 | 13848186 | 13855092 |
| VIT_18s0001g15720 | 18 | 13856940 | 13861020 |
| VIT_18s0001g15730 | 18 | 13865318 | 13866466 |
| VIT_18s0001g15740 | 18 | 13869407 | 13877879 |
| VIT_18s0001g15 | 18 | 13901947 | 13917102 |
| VIT_18s0001g15760 | 18 | 13920267 | 9 |
| VIT_18s0001g15790 | 18 | 13930846 | 13931145 |
| VIT_18s0001g15800 | 18 | 13945801 | 13947522 |
| VIT_18s0166g00010 | 18 | 13949617 | 13952330 |
| VIT_18s0166g00020 | 18 | 13966224 | 13967049 |
| VIT_18s0166g00030 | 18 | 13981658 | 13985444 |
| VIT_18s0166g00040 | 18 | 13989688 | 13993683 |
| VIT_18s0166g00050 | 18 | 13995521 | 14000593 |
| VIT_18s0166g00060 | 18 | 14002147 | 14002275 |
| VIT_18s0166g00070 | 18 | 14002488 | 14002937 |
| VIT_18s0166g00080 | 18 | 14023042 | 14027197 |
| VIT_18s0166g00090 | 18 | 14033533 | 14034350 |
| VIT_18s0166g00100 | 18 | 14041093 | 14041505 |
| VIT_18s0166g00110 | 18 | 14041566 | 14042290 |
| VIT_18s0166g00120 | 18 | 14071620 | 14072689 |
| VIT_18s0166g00130 | 18 | 14076141 | 14077530 |
| VIT_18s0166g00140 | 18 | 14096408 | 14097042 |
| VIT_18s0166g00150 | 18 | 14109274 | 14109357 |
| VIT_18s0166g00160 | 18 | 14116931 | 14117425 |
| VIT_18s0166g00170 | 18 | 14122252 | 14123294 |
| VIT_18s0166g00180 | 18 | 14130785 | 14134673 |
| VIT_18s0166g00190 | 18 | 14139015 | 14140400 |
| VIT_18s0166g00210 | 18 | 14160272 | 14171887 |
| VIT_18s0166g00220 | 18 | 14204751 | 14205007 |
| VIT_18s0166g00230 | 18 | 14207946 | 14210222 |
| VIT_18s0166g00240 | 18 | 14210516 | 14228120 |
| VIT_18s0166g00250 | 18 | 14230226 | 14231594 |
| VIT_18s0166g00260 | 18 | 14233967 | 14234837 |
| VIT_18s0166g00270 | 18 | 14276150 | 14276380 |
| VIT_18s0166g00280 | 18 | 14293747 | 142944 |

GLTP3 (glycolipid transfer protein 3)
Electron carrier/ oxidoreductase
Leucine Rich Repeat receptor-like kinase Dof zinc finger protein DOF3.5

No hit
Protein kinase Xa21
Copine BON3 (BONZAI 3)
V-type H+-transporting ATPase 16 kDa proteolipid subunit Protein kinase Xa21
Protein kinase Xa21
RWP-RK domain-containing protein
No hit
Protein kinase Xa21 EFR (EF-TU receptor)

No hit
ABC Transporter (VvWBC1 - VvABCG1)
VQ motif-containing protein
No hit
E4/E8 binding protein-1
Concanavalin A lectin
Substrate carrier, Mitochondrial
Oligopeptidase A
No hit
No hit
Unknown
Zinc finger protein ATRZ-1A
U-box domain-containing protein
Regulator of chromosome condensation (RCC1)
No hit
S-receptor kinase
RNA recognition motif (RRM)-containing protein
No hit
No hit
Mutator-like transposase-like protein
Transcription factor related

| VIT_18s0166g00290 | 18 | 14296029 | 14329589 |
| :--- | :--- | :--- | :--- |
| VIT_18s0166g00300 | 18 | 14365849 | 14379716 |
| VIT_18s0166g00310 | 18 | 14379717 | 14382427 |
| VIT_18s0166g00320 | 18 | 14382428 | 14384232 |
| VIT_18s0076g00380 | 18 | 14405761 | 14410018 |
| VIT_18s0076g00370 | 18 | 14415354 | 14416016 |
| VIT_18s0076g00360 | 18 | 14424417 | 14427563 |
| VIT_18s0076g00350 | 18 | 14479728 | 14481025 |
| VIT_18s0076g00340 | 18 | 14491505 | 14493905 |
| VIT_18s0076g00330 | 18 | 14494814 | 14503181 |
| VIT_18s0076g00320 | 18 | 14527632 | 14528344 |
| VIT_18s0076g00310 | 18 | 14550818 | 14563944 |
| VIT_18s0076g00300 | 18 | 14575052 | 14575648 |
| VIT_18s0076g00290 | 18 | 14619070 | 14619498 |
| VIT_18s0076g00280 | 18 | 14659006 | 14685257 |
| VIT_18s0076g00270 | 18 | 14725759 | 14726714 |
| VIT_18s0076g00260 | 18 | 14726740 | 14745454 |
| VIT_18s0076g00250 | 18 | 14786911 | 14789327 |
| VIT_18s0076g00240 | 18 | 14807850 | 14815102 |
| VIT_18s0076g00230 | 18 | 14827587 | 14833371 |
| VIT_18s0076g00220 | 18 | 14843815 | 14846194 |
| VIT_18s0076g00210 | 18 | 14868749 | 14920388 |
| VIT_18s0076g00200 | 18 | 14926786 | 14927997 |
| VIT_18s0076g00190 | 18 | 14937941 | 14939662 |

Protein kinase
Histidinol dehydrogenase, chloroplast precursor ABC transporter F member 4 ABC Transporter (VvGCN1 - VvABCF1) ABC Transporter (VvGCN5 - VvABCF5)
Ribosomal protein P2 (RPP2A) acidic 60S
EMB1075 (embryo defective 1075) carboxy-lyase
F-box protein (FBX3)
CYP707A3
Basic Leucine Zipper Transcription Factor (VvbZIP50)
Nodulin
Translation initiation factor eIF-5B Hexokinase-like protein No hit
Protein tyrosine phosphatase No hit
Unknown protein
Sucrose-proton symporter 2 SUC2 (SUT2-2) Preprotein translocase Sec Y subunit No hit
Sucrose-proton symporter 2 SUC2 (SUT2-3)
Alpha/beta hydrolase fold
DELLA protein RGL2 (RGA-like protein 2)

Supplementary Table 10. Definition of "molecular veraison". Number of modulated genes $(|\log 2 F C|>2)$, in red, across the time points during berry development. In black the number of days before phenological veraison.

|  | Days | t 0 |  | t 1 |  | t 2 |  | t 3 |  | t 4 |  | t 5 |
| ---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  |  |  |  |  |  |  |  |  |  |  |  |  |
| 2012 |  | -36 | 27 | -28 | 0 | -18 | 127 | -9 | 4 | 0 | 0 |  |
| 2013 |  | -20 | 11 | -14 | 52 | -7 | 26 | 0 | 0 |  | 0 |  |
| 2014 |  | -20 | 6 | -14 | 38 | -6 | 58 | 0 | 0 |  | 0 |  |

Supplementary Table 11

| Meta-QTL | Gene ID | Chr | Start <br> (bp) | End(bp) | Annotation_V1 | Transcriptomic candidates |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| pheno_3_1 | VIT_03s0038g00670 | 3 | 592539 | 594503 | fructose-bisphosphate aldolase, chloroplast precursor | Palumbo et al 2014, Massonet et al 2017 |
|  | VIT_03s0038g00760 | 3 | 644491 | 647420 | Arginine decarboxylase (Fragment) | Fasoli et al 2018 |
|  | VIT_03s0038g00860 | 3 | 689247 | 693308 | Basic Leucine Zipper Transcription Factor (VvbZIP05) | Fasoli et al 2018 |
|  | VIT_03s0038g01090 | 3 | 862995 | 863398 | Auxin responsive SAUR protein | Palumbo et al 2014, Massonet et al 2017 |
|  | VIT_03s0038g01110 | 3 | 866357 | 866897 | Auxin-responsive SAUR31 | Palumbo et al 2014, Massonet et al 2017 |
|  | VIT_03s0038g01310 | 3 | 921733 | 927965 | Auxin responsive SAUR protein | Fasoli et al 2018 |
|  | VIT_03s0038g01380 | 3 | 971266 | 975571 | Calcium-binding EF hand | Palumbo et al 2014, Massonet et al 2017, Fasoli et al 2018 |
|  | VIT_03s0038g01410 | 3 | 984045 | 985466 | Aquaporin PIP PIP1A | Palumbo et al 2014, Massonet et al 2017, Fasoli et al 2018 |
|  | VIT_03s0038g01510 | 3 | 1041388 | 1041967 | Unknown protein | Fasoli et al 2018 |
|  | VIT_03s0038g01830 | 3 | 1283253 | 1285733 | Proline-rich protein 4 | Palumbo et al 2014, Massonet et al 2017, Fasoli et al 2018 |


|  | VIT_03s0038g02050 | 3 | 1412118 | 1412819 | Dirigent protein pDIR7 | Palumbo et al 2014, Massonet et al 2017 |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | VIT_03s0038g02090 | 3 | 1441262 | 1442535 | Phospholipase C. | Palumbo et al 2014, Massonet et al 2017, Fasoli et al 2018 |
|  | VIT_03s0038g02110 | 3 | 1449330 | 1451521 | Co-chaperone-curved DNA binding protein A | Fasoli et al 2018 |
|  | VIT_03s0038g02130 | 3 | 1468239 | 1469371 | Cold shock protein-1 | Fasoli et al 2018 |
|  | VIT_03s0038g02150 | 3 | 1482091 | 1483271 | Unknown | Palumbo et al 2014, Massonet et al 2017 |
|  | VIT_03s0038g02170 | 3 | 1492918 | 1494778 | Thaumatin | Palumbo et al 2014, Massonet et al 2017 Fasoli et al 2018 |
|  | VIT_03s0038g02190 | 3 | 1502133 | 1506299 | Nodulin | Fasoli et al 2018 |
|  | VIT_03s0038g02220 | 3 | 1528083 | 1528899 | Cofilin | Palumbo et al 2014, Massonet et al 2017 |
|  | VIT_03s0038g02230 | 3 | 1535564 | 1538193 | Subtilisin-like serine protease 2 | Palumbo et al 2014, Massonet et al 2017 |
| pheno_3_2 | VIT_03s0180g00010 | 3 | 5934501 | 5937492 | Cysteine proteinase | $\begin{array}{r} \hline \text { Fasoli et al } \\ 2018 \end{array}$ |
|  | VIT_03s0180g00040 | 3 | 5973785 | 5975813 | Cyclin D3_2 | Fasoli et al 2018 |
|  | VIT_03s0180g00090 | 3 | 6023040 | 6024531 | Receptor kinase | Fasoli et al 2018 |


| VIT_03s0180g00200 | 3 | 6124465 | 6126202 | Limonoid UDP-glucosyltransferase (VvGT2) | Palumbo et al 2014, Massonet et al 2017, Fasoli et al 2018 |
| :---: | :---: | :---: | :---: | :---: | :---: |
| VIT_03s0180g00280 | 3 | 6255489 | 6257208 | Indole-3-acetate beta-glucosyltransferase | Fasoli et al 2018 |
| VIT_03s0180g00320 | 3 | 6281986 | 6283705 | indole-3-acetate beta-glucosyltransferase (VvGT3) | Palumbo et al 2014, Massonet et al 2017 |
| VIT_03s0091g00210 | 3 | 6507392 | 6509263 | Ethylene-responsive protein | Palumbo et al 2014, Massonet et al 2017 |
| VIT_03s0091g00240 | 3 | 6521904 | 6537905 | Haloacid dehalogenase hydrolase | Fasoli et al 2018 |
| VIT_03s0091g00260 | 3 | 6548677 | 6549577 | Zinc finger protein 4 | Palumbo et al 2014, Massonet et al 2017, Fasoli et al 2018 |
| VIT_03s0091g00420 | 3 | 6785458 | 6787059 | No hit | Palumbo et al 2014, Massonet et al 2017, Fasoli et al 2018 |
| VIT_03s0091g00450 | 3 | 6811798 | 6813280 | Progesterone 5-beta-reductase | Palumbo et al 2014, Massonet et al 2017 |
| VIT_03s0091g00490 | 3 | 6859815 | 6860409 | Unknown protein | Palumbo et al 2014, Massonet et al 2017 |
| VIT_03s0091g00500 | 3 | 6871858 | 6872341 | Unknown protein | Palumbo et al 2014, Massonet et al 2017 |
| VIT_03s0091g00580 | 3 | 7035451 | 7039010 | Protein Mpv17 | Fasoli et al $2018$ |


| VIT_03s0091g00670 | 3 | 7099015 | 7099742 |
| :--- | :--- | :--- | :--- |
| VIT_03s0091g00680 | 3 | 7103281 | 7113666 |
| VIT_03s0091g00870 | 3 | 7342165 | 7357074 |
| VIT_03s0091g01010 | 3 | 7535063 | 7536574 |
| VIT_03s0091g01060 | 3 | 7673917 | 7675754 |
| VIT_03s0091g01130 | 3 | 7741682 | 7743818 |
|  |  |  |  |
| VIT_03s0091g01240 | 3 | 7903958 | 7907570 |
|  |  |  |  |
| VIT_03s0091g01290 | 3 | 7951175 | 7955690 |
| VIT_03s0088g00320 | 3 | 8343365 | 8347681 |
| VIT_03s 00088 s 0088 g 00050 | 3 | 8035215 | 8039037 |
|  |  | 8315170 | 8315924 |


| Lateral organ boundaries protein 38 | Palumbo et al 2014, Massonet et al 2017, Fasoli et al 2018 |
| :---: | :---: |
| NPY2 (naked pins in yuc mutants 2) | Fasoli et al 2018 |
| Adenylylsulfate kinase 1 (AKN1) | Fasoli et al 2018 |
| LIM domain protein WLIM1 | Palumbo et al 2014, Massonet et al 2017, Fasoli et al 2018 |
| Cyclin delta-2 | Fasoli et al 2018 |
| Meprin and TRAF homology domain-containing protein | Palumbo et al 2014, Massonet et al 2017 |
| Serine carboxypeptidase S10 / Anthocyanin Acyl-transferase | Palumbo et al 2014, Massonet et al 2017, Fasoli et al 2018 |
| Serine carboxypeptidase S10 | Palumbo et al 2014, Massonet et al 2017 |
| Serine carboxypeptitase 1 | Palumbo et al 2014, Massonet et al 2017 |
| Phytosulfokines PSK2 | Palumbo et al 2014, Massonet et al 2017 |
| Peptidase M50 | Palumbo et al 2014, Massonet et al 2017 |


|  | VIT_03s0088g00710 | 3 | 8916223 | 8916893 | Pathogenesis-related protein 1 precursor (PRP 1) | Palumbo et al 2014, Massonet et al 2017 |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | VIT_03s0088g00810 | 3 | 9044529 | 9045211 | Pathogenesis-related protein 1 precursor (PRP 1) | Palumbo et al 2014, Massonet et al 2017 |
|  | VIT_03s0088g01180 | 3 | 9438885 | 9442060 | Proline iminopeptidase | Fasoli et al 2018 |
|  | VIT_03s0088g01250 | 3 | 9538993 | 9552353 | D-threo-aldose 1-dehydrogenase | Palumbo et al 2014, Massonet et al 2017 |
| pheno_5_1 | VIT_05s0029g01140 | 5 | 17405622 | 17411752 | Sucrose-phosphate synthase | Palumbo et al 2014, Massonet et al 2017, Fasoli et al 2018 |
|  | VIT_05s0062g00240 | 5 | 18818659 | 18820264 | Xyloglucan endotransglucosylase/hydrolase 23 | Palumbo et al 2014, Massonet et al 2017 |
|  | VIT_05s0062g00250 | 5 | 18831568 | 18833016 | Xyloglucan endotransglucosylase/hydrolase 15 | Palumbo et al 2014, Massonet et al 2017 |
|  | VIT_05s0062g00270 | 5 | 18843664 | 18845204 | UDP-glucose:flavonoid 7-O-glucosyltransferase | Fasoli et al 2018 |
|  | VIT_05s0062g00430 | 5 | 18991958 | 18993513 | UDP-glucose:flavonoid 7-O-glucosyltransferase | Palumbo et al 2014, Massonet et al 2017 |
|  | VIT_05s0062g00480 | 5 | 19056125 | 19057777 | Xyloglucan endo-transglycosylase, C-terminal | Palumbo et al 2014, Massonet et al 2017, Fasoli et al 2018 |
|  | VIT_05s0062g00520 | 5 | 19088785 | 19090792 | UDP-glucose:flavonoid 7-O-glucosyltransferase | Palumbo et al 2014, Massonet et al 2017 |


|  |  |  | Palumbo et al <br> 2014, Massonet <br> et al 2017, |
| :--- | :--- | :--- | :--- | :--- | :--- |
| Fasoli et al |  |  |  |
| 2018 |  |  |  |


|  | VIT_11s0016g03940 | 11 | 3224068 | 3225265 | Heat shock transcription factor C1 | Palumbo et al 2014, Massonet et al 2017 |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | VIT_11s0016g03980 | 11 | 3249496 | 3250787 | Unknown protein | Fasoli et al 2018 |
| pheno_12_1 | VIT_12s0035g02080 | 12 | 23970815 | 23972655 | Unknown protein | Fasoli et al 2018 |
|  | VIT_12s0035g02090 | 12 | 23983677 | 23999372 | Leucine-rich repeat family protein | Fasoli et al 2018 |
|  | VIT_12s0035g02120 | 12 | 24046092 | 24050103 | Unknown | Fasoli et al 2018 |
|  | VIT_12s0035g02150 | 12 | 24097742 | 24106206 | ferric reduction oxidase 7 FRO7 | Palumbo et al 2014, Massonet et al 2017, Fasoli et al 2018 |
| pheno_14_1 | VIT_14s0083g00580 | 14 | 22620849 | 22623743 | S-adenosyl-L-methionine decarboxylase | Fasoli et al 2018 |
|  | VIT_14s0083g00620 | 14 | 22672469 | 22675655 | NIK1 (NSP- interacting kinase 1) | Palumbo et al 2014, Massonet et al 2017 |
|  | VIT_14s0083g00640 | 14 | 22696160 | 22698346 | Constans 2 (COL2) | Palumbo et al 2014, Massonet et al 2017 |
|  | VIT_14s0083g00910 | 14 | 23106621 | 23115040 | Auxin-independent growth promoter | Fasoli et al 2018 |
|  | VIT_14s0083g00940 | 14 | 23137130 | 23143930 | Auxin-independent growth promoter | Palumbo et al 2014, Massonet et al 2017, Fasoli et al 2018 |
|  | VIT_14s0083g01030 | 14 | 23320331 | 23341036 | putative MADS-box Fruitfull 2 (VviFUL2) | Palumbo et al 2014, Massonet et al 2017, Fasoli et al 2018 |


| VIT_14s0083g01100 | 14 | 23425741 | 23428785 |
| :--- | :--- | :--- | :--- | :--- |
| VIT_14s0083g01110 | 14 | 23435436 | 23438457 |
| VIT_14s0083g01140 | 14 | 23478460 | 23480345 |
| VIT_14s0083g01160 | 14 | 23527926 | 23532692 |
| VIT_14s0083g01170 | 14 | 23552658 | 23569757 |
| VIT_14s0083g01210 | 14 | 23631468 | 23634185 |
| VIT_14s0083g01220 | 14 | 23647671 | 23648618 |
|  |  |  |  |
| VIT_14s0068g00010 | 14 | 23691896 | 23694505 |
| VIT_14s0068g00030 | 14 | 23710282 | 23713253 |
| VIT_14s0068g00040 | 14 | 23730955 | 23731566 |
|  |  |  |  |


| Alpha-1,4-glucan-protein synthase 1 | Palumbo et al 2014, Massonet et al 2017 |
| :---: | :---: |
| Brassinosteroid-6-oxidase | Palumbo et al 2014, Massonet et al 2017 |
| B12D | Palumbo et al 2014, Massonet et al 2017 |
| COBRA protein | Fasoli et al 2018 |
| Mitochondrial substrate carrier family protein | Fasoli et al 2018 |
| feronia receptor-like kinase | Palumbo et al 2014, Massonet et al 2017 |
| feronia receptor-like kinase | Palumbo et al 2014, Massonet et al 2017 |
| feronia receptor-like kinase | Palumbo et al 2014, Massonet et al 2017 |
| feronia receptor-like kinase | Palumbo et al 2014, Massonet et al 2017 |
| No hit | Palumbo et al 2014, Massonet et al 2017 |
| No hit | Palumbo et al 2014, Massonet et al 2017 |


|  |  |  |  | Palumbo et al <br> 2014, Massonet <br> et al 2017 |
| :--- | :--- | :--- | :--- | :--- | :--- |
| VIT_14s0068g00140 |  |  |  |  |


|  | VIT_16s0022g01770 | 16 | 15467138 | 15473529 | Phosphopyruvate hydratase. | Palumbo et al 2014, Massonet et al 2017, Fasoli et al 2018 |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | VIT_16s0022g01870 | 16 | 15736355 | 15738229 | Unknown protein | Fasoli et al 2018 |
|  | VIT_16s0022g02080 | 16 | 16041486 | 16043235 | Exostosin family protein | Fasoli et al 2018 |
|  | VIT_16s0022g02100 | 16 | 16060136 | 16061359 | Embryo-specific 3 | Fasoli et al 2018 |
|  | VIT_16s0022g02200 | 16 | 16203664 | 16206181 | Subtilase | Palumbo et al 2014, Massonet et al 2017 |
|  | VIT_16s0022g02230 | 16 | 16240572 | 16248680 | Leucine-rich repeat receptor protein kinase EXS | Fasoli et al 2018 |
|  | VIT_16s0022g02340 | 16 | 16470141 | 16475595 | fructokinase-2 | Fasoli et al 2018 |
| pheno_16_2 | VIT_16s0100g00220 | 16 | 17132026 | 17132339 | No hit | Fasoli et al 2018 |
|  | VIT_16s0100g00270 | 16 | 17150479 | 17150928 | Peptidoglycan-binding LysM domain-containing protein | Palumbo et al 2014, Massonet et al 2017 |
|  | VIT_16s0100g00290 | 16 | 17162424 | 17180217 | L-idonate dehydrogenase | Palumbo et al 2014, Massonet et al 2017 |
|  | VIT_16s0100g00350 | 16 | 17248816 | 17261155 | ABC Transporter (VvTAP3 - VvABCB23) | $\begin{array}{r} \text { Fasoli et al } \\ 2018 \\ \hline \end{array}$ |
| pheno_17_1 | VIT_17s0000g04750 | 17 | 5279676 | 5281368 | UDP-glycosyltransferase 89B2 | Fasoli et al 2018 |
|  | VIT_17s0000g04840 | 17 | 5357937 | 5362286 | Sterile alpha motif (SAM) domain-containing | Palumbo et al 2014, Massonet et al 2017 |
|  | VIT_17s0000g04890 | 17 | 5409996 | 5423899 | D-aminoacyl-tRNA deacylase GEKO1 | Fasoli et al 2018 |
|  | VIT_17s0000g05020 | 17 | 5637669 | 5644801 | Squamosa promoter-binding protein 6 (SPL6) | Palumbo et al 2014, Massonet et al 2017 |


| VIT_17s0000g05030 | 17 | 5645732 | 5648492 | Alpha-1,4-glucan-protein synthase 1 | Fasoli et al 2018 |
| :---: | :---: | :---: | :---: | :---: | :---: |
| VIT_17s0000g05040 | 17 | 5654593 | 5656105 | No hit | Fasoli et al 2018 |
| VIT_17s0000g05050 | 17 | 5659282 | 5660704 | COBRA-like protein 4 | Fasoli et al 2018 |
| VIT_17s0000g05070 | 17 | 5676169 | 5679862 | Phytochelatin synthetase | Palumbo et al 2014, Massonet et al 2017, Fasoli et al 2018 |
| VIT_17s0000g05110 | 17 | 5731853 | 5734268 | CYP78A4 | Palumbo et al 2014, Massonet et al 2017 |
| VIT_17s0000g05240 | 17 | 5869290 | 5885095 | Nuclear transport factor 2 (NTF2) | Fasoli et al 2018 |
| VIT_17s0000g05250 | 17 | 5885327 | 5886300 | Unknown protein | Palumbo et al 2014, Massonet et al 2017 |
| VIT_17s0000g05460 | 17 | 6088077 | 6090457 | Nodulin | Palumbo et al 2014, Massonet et al 2017 |
| VIT_17s0000g05550 | 17 | 6183508 | 6195130 | Proton-dependent oligopeptide transport (POT) family protein | Palumbo et al 2014, Massonet et al 2017 |
| VIT_17s0000g05580 | 17 | 6213229 | 6221132 | Isopiperitenol dehydrogenase | Palumbo et al 2014, Massonet et al 2017, Fasoli et al 2018 |
| VIT_17s0000g05600 | 17 | 6255629 | 6256773 | (-)-isopiperitenol dehydrogenase | Fasoli et al 2018 |
| VIT_17s0000g05610 | 17 | 6257443 | 6258438 | Isopiperitenol dehydrogenase | Fasoli et al 2018 |

$\left.\begin{array}{llllll} & & & \begin{array}{r}\text { Palumbo et al } \\ \text { 2014, Massonet } \\ \text { et al } 2017\end{array} \\ \text { Fasoli et al }\end{array}\right)$

| VIT_18s0001g03610 | 18 | 3401893 | 3411144 | Auxin-independent growth promoter | Palumbo et al 2014, Massonet et al 2017 |
| :---: | :---: | :---: | :---: | :---: | :---: |
| VIT_18s0001g03670 | 18 | 3422279 | 3424214 | Zinc finger (C2H2 type) family | Palumbo et al 2014, Massonet et al 2017 |
| VIT_18s0001g03880 | 18 | 3565751 | 3567556 | Polcalcin | Fasoli et al 2018 |
| VIT_18s0001g04150 | 18 | 3732776 | 3738425 | Avr9 elicitor response protein | Fasoli et al 2018 |
| VIT_18s0001g04340 | 18 | 3822948 | 3829597 | Glycine hydroxymethyltransferase | Fasoli et al 2018 |
| VIT_18s0001g04680 | 18 | 3938582 | 3956444 | RPG related protein 1 RR1 | Palumbo et al 2014, Massonet et al 2017 |
| VIT_18s0001g04790 | 18 | 3994178 | 4000375 | Aspartic Protease (VvAP42) | Palumbo et al 2014, Massonet et al 2017 |
| VIT_18s0001g04920 | 18 | 4068936 | 4069711 | No hit | Palumbo et al 2014, Massonet et al 2017 |
| VIT_18s0001g04930 | 18 | 4070013 | 4071952 | Sulfate transporter 1.3 | Palumbo et al 2014, Massonet et al 2017 |
| VIT_18s0001g05020 | 18 | 4122072 | 4123281 | Senescence-associated protein | Palumbo et al 2014, Massonet et al 2017 |
| VIT_18s0001g05040 | 18 | 4138118 | 4140727 | Pollen Ole e 1 allergen and extensin | Palumbo et al 2014, Massonet et al 2017 |
| VIT_18s0001g05180 | 18 | 4185500 | 4189977 | Beta-D-xylosidase | Palumbo et al 2014, Massonet et al 2017, Fasoli et al 2018 |


| VIT_18s0001g05220 | 18 | 4205002 | 4212978 | WD-40 repeat | Fasoli et al 2018 |
| :---: | :---: | :---: | :---: | :---: | :---: |
| VIT_18s0001g05300 | 18 | 4254678 | 4257480 | Trehalose-6-phosphate phosphatase | Fasoli et al 2018 |
| VIT_18s0001g05570 | 18 | 4363673 | 4368991 | Hexose transporter HT2 | Fasoli et al 2018 |
| VIT_18s0001g05910 | 18 | 4550436 | 4552401 | Cis-zeatin O-beta-D-glucosyltransferase | Palumbo et al 2014, Massonet et al 2017 |
| VIT_18s0001g05990 | 18 | 4574889 | 4586041 | UDP-glycosyltransferase 85A1 | Palumbo et al 2014, Massonet et al 2017 |
| VIT_18s0001g06060 | 18 | 4605365 | 4607009 | UDP-glycosyltransferase 85A1 | Palumbo et al 2014, Massonet et al 2017, Fasoli et al 2018 |
| VIT_18s0001g06180 | 18 | 4659693 | 4663192 | Phosphate-induced protein 1 | Palumbo et al 2014, Massonet et al 2017 |
| VIT_18s0001g06220 | 18 | 4684266 | 4686537 | F-box protein (FBW2) | Palumbo et al 2014, Massonet et al 2017 |
| VIT_18s0001g06370 | 18 | 4765814 | 4772318 | L-ascorbate peroxidase, chloroplast | Palumbo et al 2014, Massonet et al 2017 |
| VIT_18s0001g06430 | 18 | 4806981 | 4808947 | Homeobox-leucine zipper protein ATHB-6 | Palumbo et al 2014, Massonet et al 2017, Fasoli et al 2018 |
| VIT_18s0001g06580 | 18 | 4966021 | 4966915 | Plastocyanin domain-containing protein | Palumbo et al 2014, Massonet et al 2017 |


|  | VIT_18s0001g06690 | 18 | 5029717 | 5032277 | Gibberellin 20 oxidase 2 | Palumbo et al 2014, Massonet et al 2017 |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | VIT_18s0001g06820 | 18 | 5077418 | 5081480 | MATE efflux family protein ripening responsive | Fasoli et al 2018 |
|  | VIT_18s0001g07090 | 18 | 5290562 | 5293561 | Unknown protein | Fasoli et al 2018 |
| pheno_18_2 | VIT_18s0001g12830 | 18 | 10937041 | 10940329 | 1,4-beta-mannan endohydrolase | Palumbo et al 2014, Massonet et al 2017, Fasoli et al 2018 |
|  | VIT_18s0001g12840 | 18 | 10940330 | 10945165 | ADP-glucose pyrophosphorylase large subunit CagpL2 | Fasoli et al 2018 |
|  | VIT_18s0001g12960 | 18 | 11043696 | 11049454 | Wound-responsive protein | Fasoli et al 2018 |
|  | VIT_18s0001g12990 | 18 | 11099778 | 11102242 | Anthranilate N-benzoyltransferase protein 1 | Fasoli et al 2018 |
|  | VIT_18s0001g13010 | 18 | 11126023 | 11129236 | Mitogen-activated Protein Kinase (VvMPK11) | Fasoli et al 2018 |
|  | VIT_18s0001g13200 | 18 | 11256653 | 11261569 | Cytokinin dehydrogenase 5 precursor | Palumbo et al 2014, Massonet et al 2017 |
|  | VIT_18s0001g13230 | 18 | 11286706 | 11293739 | Beta-galactosidase BG1 [Vitis vinifera] | Palumbo et al 2014, Massonet et al 2017 |
|  | VIT_18s0001g13360 | 18 | 11384285 | 11387145 | Nodulin MtN21 family | Palumbo et al 2014, Massonet et al 2017 |
|  | VIT_18s0001g13610 | 18 | 11613829 | 11620092 | Abhydrolase domain-containing protein 5 | Fasoli et al 2018 |
|  | VIT_18s0001g13770 | 18 | 11767560 | 11769317 | Cytochrome P450, family 83, subfamily B, polypeptide 1 | Fasoli et al 2018 |
|  | VIT_18s0001g13780 | 18 | 11774368 | 11775896 | Cytochrome P450, family 83, subfamily B, polypeptide 1 | Fasoli et al 2018 |
|  | VIT_18s0001g13790 | 18 | 11779263 | 11781156 | Cytochrome P450, family 83, subfamily B, polypeptide 1 | Fasoli et al $2018$ |


| VIT_18s0001g13970 | 18 | 11974983 | 11980862 |
| :--- | :--- | :--- | :--- |
| VIT_18s0001g14040 | 18 | 12086327 | 12090970 |
|  |  |  |  |
| VIT_18s0001g14130 | 18 | 12179540 | 12181647 |
|  |  |  |  |
| VIT_18s0001g14260 | 18 | 12273788 | 12276090 |
|  |  |  |  |
| VIT_18s0001g14270 | 18 | 12276091 | 12276597 |
|  |  |  |  |
| VIT_18s0001g14360 | 18 | 12337145 | 12340985 |
| VIT_18s0001g14440 | 18 | 12432955 | 12439459 |
| VIT_18s0001g14450 | 18 | 12453766 | 12456583 |
| VIT_18s 0001 s 00014780 | 18 | 12841218 | 12848820 |


| Myosin-related | Palumbo et al 2014, Massonet et al 2017, Fasoli et al 2018 |
| :---: | :---: |
| Endo-1,4-beta-glucanase | Palumbo et al 2014, Massonet et al 2017 |
| Zinc finger (C2H2 type) family | Palumbo et al 2014, Massonet et al 2017 |
| No hit | Palumbo et al 2014, Massonet et al 2017, Fasoli et al 2018 |
| Gibberellin-regulated protein 1 (GASA1) | Palumbo et al 2014, Massonet et al 2017, Fasoli et al 2018 |
| Tubulin beta-1 chain | Palumbo et al 2014, Massonet et al 2017 |
| Molecular chaperone DnaJ | Palumbo et al 2014, Massonet et al 2017 |
| Ferredoxin:nadp+ Oxidoreductase PETH | Palumbo et al 2014, Massonet et al 2017 |
| No hit | Fasoli et al 2018 |
| Lipase 3 (EXL3) family II extracellular | Fasoli et al 2018 |


| VIT_18s0001g14980 | 18 | 12992480 | 12996562 | 3-methyl-2-oxobutanoate dehydrogenase | Palumbo et al 2014, Massonet et al 2017 |
| :---: | :---: | :---: | :---: | :---: | :---: |
| VIT_18s0001g15000 | 18 | 13026931 | 13036065 | ACT domain containing protein (ACR4) | Palumbo et al 2014, Massonet et al 2017, Fasoli et al 2018 |
| VIT_18s0001g15120 | 18 | 13199060 | 13200303 | Unknown protein | Palumbo et al 2014, Massonet et al 2017, Fasoli et al 2018 |
| VIT_18s0001g15360 | 18 | 13495083 | 13497819 | Thylakoid lumenal 29.8 kDa protein | Fasoli et al 2018 |
| VIT_18s0001g15390 | 18 | 13521135 | 13522636 | Gaiacol peroxidase | Fasoli et al 2018 |
| VIT_18s0001g15410 | 18 | 13539135 | 13541625 | Alcohol dehydrogenase 1 | Palumbo et al 2014, Massonet et al 2017, Fasoli et al 2018 |
| VIT_18s0001g15460 | 18 | 13616618 | 13618309 | Stearyl acyl carrier protein desaturase | Palumbo et al 2014, Massonet et al 2017 |
| VIT_18s0001g15520 | 18 | 13650247 | 13655312 | Leaf senescence protein | Palumbo et al 2014, Massonet et al 2017, Fasoli et al 2018 |
| VIT_18s0001g15720 | 18 | 13856940 | 13861020 | Leucine Rich Repeat receptor-like kinase | Palumbo et al 2014, Massonet et al 2017, Fasoli et al 2018 |
| VIT_18s0001g15730 | 18 | 13865318 | 13866466 | Dof zinc finger protein DOF3.5 | Fasoli et al 2018 |
| VIT_18s0166g00250 | 18 | 14230226 | 14231594 | No hit | Fasoli et al 2018 |


| VIT_18s0076g00330 | 18 | 14494814 | 14503181 | Basic Leucine Zipper Transcription Factor (VvbZIP50) | Fasoli et al <br> 2018 |  |  |
| :--- | ---: | ---: | ---: | ---: | ---: | ---: | ---: |
| VIT_18s0076g00310 | 18 | 14550818 | 14563944 |  |  | Translation initiation factor eIF-5B | Fasoli et al |
| 2018 |  |  |  |  |  |  |  |

Supplementary Table 12

| NAME | $\begin{aligned} & \text { COD } \\ & \text { E } \end{aligned}$ | NAME | $\begin{aligned} & \text { COD } \\ & \text { E } \end{aligned}$ | NAME | $\begin{aligned} & \text { COD } \\ & \mathbf{E} \end{aligned}$ | NAME | $\begin{aligned} & \text { COD } \\ & \text { E } \end{aligned}$ |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| ABBOTTABOTTI | 943 | BERLA_GROSA | 1069 | CAPIBIANCHI | 1159 | CORNIOLA | 1103 |
| ABBUOTO | 1 | BERTINORA | 1030 | CAPOBIANCO_BIANCO | 507 | CORSO_BIANCO | 1046 |
| ABELLO | 677 | BERZAMINO | 692 | CAPOBIANCO_NERO | 508 | CORVINA | 82 |
| ABRUSCO | 2 | BIANCHETTA_DI_BACEDASC O | 693 | CAPOLONGO_1 | 1045 | CORVINA_ASOLO | 624 |
| AGHEDENE | 678 | BIANCHETTA_DI_DIOLO | 694 | CAPOLUNGO | 509 | CORVINONE | 83 |
| AGLIANICO | 3 | BIANCHETTA_TREVIGIANA | 32 | CAPRETTA | 510 | COVE | 84 |
| AGLIANICO_GRANA_GROSSA | 944 | BIANCO_DALESSANO | 33 | CAPRUGNONE | 1160 | CREPALLOCCHI | 722 |
| AHMEUR_MONARDE | 1146 | BIANCOLELLA | 34 | CARDINAL | 433 | CROATINA | 85 |
| AIREN | 676 | BIANCONE | 37 | CARICAGIOLA | 66 | CROVASSA | 86 |
| ALBANA | 6 | BIANCONE_DELLANTELLA | 1034 | CARIGNANO | 67 | CUNEUTE | 723 |
| ALBANA_BORDINI | 1061 | BIANCUZZO | 695 | CARMENERE | 50 | DALL | 724 |
| ALBANA_DEL_PANIERE | 679 | BICANE | 424 | CARRICANTE | 68 | DAMASCHINO | 87 |
| ALBANELLO | 7 | BIGOLONA | 696 | CASAVECCHIA | 366 | DAMIANO | 726 |
| ALBARANZEULI_BIANCO | 10 | BLANC_DE_MORGEX | 35 | CASENTINO | 1161 | DANUTA | 484 |


| ALBAROLA | 30 | BOGGIONE_NERO | 1151 | CASETTA | 362 | DATTIER_DE_BEYROUTH | 431 |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| ALBAROSSA | 8 | BOGGIONE_ROSSO | 698 | CATALANESCA_BIANCA | 425 | DELIGHT | 493 |
| ALEANTE_RIVALTO | 690 | BOMBINO_BIANCO | 38 | CATARRATTO_BIANCO_COMUNE | 70 | DIAMANT | 1113 |
| ALEATICO | 11 | BOMBINO_NERO | 39 | CATARRATTONE | 1093 | DIMIAT | 1105 |
| ALICANTE_BOUSCHET | 12 | BONAMICO | 48 | CAVECIA | 710 | DINDARELLA | 88 |
| ALIONZA | 16 | BONARDA | 40 | CAVRARA_NERA | 1068 | DIOLINOIR | 89 |
| ALPHONSE_LAVALLEE | 430 | BONDA | 41 | CENERENTE | 512 | DOLCIAME | 91 |
| AMERICA | 984 | BORBOTTONE | 700 | CENTESIMINO | 299 | DON_MARIANO | 435 |
| ANCELLOTTA | 17 | BOSCHERA | 42 | CERASOLO | 712 | DOUX_DHENRY | 92 |
| ANSONICA | 18 | BOSCO | 43 | CESANESE_DEI_CASTELLI_ROMA NI | 513 | DURAPECCIO_DI_SLORENZO | 729 |
| ANTINELLO | 979 | BOVALE | 45 | CHARDONNAY | 244 | DURELLA | 93 |
| APIRENA_BRUNI | 495 | BRACCIOLA_NERA | 46 | CHASSELAS_BLANC | 426 | EARLY_MUSCAT | 479 |
| APRILLA | 946 | BRACHETTO | 47 | CHENIN | 298 | EMERALD_RIESLING | 667 |
| ARAMONE | 703 | BRAMBANA | 701 | CIAVARELLA_FRANCESE | 514 | EMPEROR | 1107 |
| ARGUMANNU | 684 | BRESSANA | 502 | CICCHIOLA | 986 | ERBALUCE | 94 |
| ARIS | 641 | BRUGNOLA | 503 | CICINESE_GROSSO | 715 | ERBISEDDA | 521 |
| ARNEIS | 19 | BRUGNOLINO | 1037 | CILIEGIOLO | 74 | ERVI | 95 |
| ARSILICO | 685 | BUBBIA | 702 | CIMINNITA | 429 | FALANGHINA | 96 |
| ARVESINIADU | 20 | BURGHESANA | 1067 | CLAIRETTE | 75 | FERTILIA | 97 |
| AURORA_2 | 1116 | BUSSANELLO | 49 | CLARETTE | 1071 | FIANO_ROSA_LOCOROTONDO_2 | 1013 |
| AVANA | 22 | CABERNET_FRANC | 51 | CLOTILDE_PROSPERI | 449 | FLAVIS | 99 |
| AVARENGO | 23 | CABERNET_SAUVIGNON | 52 | COCCIATOSTA | 716 | FOGARINA | 731 |
| AXINA_DE_FRANCIA | 687 | CACCIADEBITI | 705 | COCOCCIOLA | 76 | FOGLIA_TONDA | 100 |
| AXINA_DE_TRES_BIAS | 688 | CACCIO_DI_FERMO_1 | 951 | CODA_DI_VOLPE_BIANCA | 77 | FOGLIONA | 1076 |
| BACCHUS | 642 | CADDIU | 57 | CODELUNGHE | 717 | FORASTERA | 101 |
| BARBERA | 24 | CAGNULARI | 58 | COLOMBANA_NERA | 78 | FORCELLA | 1042 |

| BARBERA_BIANCA | 25 | CALABRESE | 59 | COLORINO_FORTE | 1167 | FORCESE_DASCOLI | 1043 |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| BARBERA_SELVATICO | 1150 | CALAMARO | 708 | COPETA | 985 | FORGIARIN | 102 |
| BARESANA | 422 | CALORIA | 60 | CORBINELLA_PADOVANA | 518 | FORSELLINA | 103 |
| BARSAGLINA | 27 | CANAIOLO_NERO | 62 | CORBINONA_13_GT | 1074 | FORTANA | 104 |
| BECUET | 290 | CANINA_1_1 | 1157 | CORINTO_BIANCO | 477 | FOSCA | 732 |
| BEGUGNOL | 689 | CANINA_NERA | 64 | CORINTO_NERO | 488 | FRACIDELLA | 733 |
| BELLINO | 432 | CANNAMELE | 709 | CORNACCHIA | 1170 | FRANCONIA | 106 |
| BELLISE | 1029 | CANNER_SEEDLESS | 483 | CORNALLIN | 79 | FRAPPATO | 107 |
| BELLONE | 28 | CAPENA | 506 | CORNAREA | 80 | FREISA | 108 |
| NAME | $\begin{aligned} & \text { COD } \\ & \text { E } \end{aligned}$ | NAME | $\begin{aligned} & \text { COD } \\ & \text { E } \end{aligned}$ | NAME | $\begin{aligned} & \text { COD } \\ & \text { E } \end{aligned}$ | NAME | $\begin{aligned} & \text { COD } \\ & \text { E } \end{aligned}$ |
| FRENCH_COLOMBARD | 1025 | INCTERZI_N1 | 134 | MALVASIA_DI_CANDIA_AROMA TICA | 160 | MOSCATO_DADDA | 443 |
| FUBIANO | 109 | INVERNENGA | 135 | MALVASIA_DI_CASORZO | 161 | MOSCATO_DAMBURGO | 444 |
| FULIGNO | 738 | INVERNESCA | 455 | MALVASIA_DI_SCHIERANO | 164 | MOSCATO_DI_SCANZO | 185 |
| FUMIN | 110 | INZOLIA_IMPERIALE | 471 | MALVASIA_FINA | 666 | MOSCATO_DI_SPAGNA | 1183 |
| FURMINT | 647 | INZOLIA_PARCHITANA | 1110 | MALVASIA_FURMHANN | 547 | MOSCATO_DI_TERRACINA | 460 |
| GABBA_VOLPE | 525 | ITALIA | 473 | MALVASIA_ISTRIANA | 165 | MOSCATO_GIALLO | 186 |
| GAGLIOPPO | 111 | ITALICA | 136 | MALVASIA_NERA_DI_BRINDISI_1 | 166 | MOSCATO_NERO | 559 |
| GALLICO | 966 | JULY_MUSCAT | 472 | MALVASIA_NERA_LUNGA | 368 | MOSCATO_NERO_D`ACQUI | 187 |
| GAMARET | 112 | JUVARELLO | 542 | MAMMOLETTA | 1178 | MOSCATO_ROSA_2 | 188 |
| GAMAY | 243 | KADARKA | 674 | MAMMOLO | 169 | MOSCATO_VIOLETTO | 557 |
| GAMBUGLIANA | 739 | KALILI | 486 | MANTONICO_BIANCO | 978 | MULLER_THURGAU | 197 |
| GARANOIR | 115 | KERNER | 137 | MANTONICONE | 977 | MUSCAT_OTTONEL | 637 |
| GAROFANA | 747 | KINGS_RUBY | 481 | MANZONI_BIANCO | 170 | NASCETTA | 367 |
| GATTA | 740 | LACRIMA | 138 | MANZONI_MOSCATO | 292 | NASCO | 198 |
| GERUSALEMME | 1094 | LACRIMA_DI_MARIA | 456 | MANZONI_ROSA | 293 | NEBBIERA | 199 |

GIRO
GIUSTOLISI
GOLDRIESLING
GOLDTRAMINER
GORGOTTESCO
GOSEN
GRAPPI_DE_LA_BORDOGNE
GRAY_RIESLING
GRECHETTO_DI_TODI
GRECO_DI_TUFO
GRECO_NERO_1
GRENACHE_NOIR
GRIGNOLINO
GRILLO
GRILLONE
GROPPELLA
GROPPELLO_DI_MOCASINA
GROPPELLO_DI_REVO
GROPPELLO_GENTILE
GROS BOUSCHET
GROSPARIOL
GROSSOLANO
gUARDAVALLE
harslevelu
IMPIGNO

| 114 | LAGREIN | 139 | MARCHIONE_2 |
| :---: | :---: | :---: | :---: |
| 742 | LAMBRUSCA_DI_ALESSAND RIA | 140 | MARCONE |
| 651 | LAMBRUSCO_A_FOGLIA_FRA ST | 141 | MARSANNE |
| 365 | LAMBRUSCO_DI_SORBARA | 142 | MARSIGLIANA_NERA_1 |
| 1176 | LAMBRUSCO_GRASPAROSSA | 143 | MARUGA |
| 364 | LAMBRUSCO_MAESTRI | 144 | MARUGGIO_1 |
| 629 | LAMBRUSCO_MARANI | 145 | MARZEMINA_2 |
| 668 | LAMBRUSCO_MONTERICCO | 146 | MARZEMINO |
| 1048 | LAMBRUSCO_OLIVA | 149 | MATILDE |
| 120 | LAMBRUSCO_SALAMINO | 147 | MATURANO |
| 122 | LAMBRUSCO_VIADANESE | 148 | MAYOLET |
| 654 | LATTUARIO_NERO | 437 | MAZZESE |
| 123 | LICRONAXIU | 1051 | MEDRULINU |
| 124 | LINNEO_NERO | 439 | MELARA_1 |
| 746 | LIVORNESE_BIANCA | 289 | MERERA |
| 1078 | LUGLIOLA_ROSSA | 753 | MERLOT |
| 125 | LUGLIOLA_ROSSA | 753 | MICHELE_PALIERI |
| 291 | LUMASSINA | 150 | MINNELLA_BIANCA_2 |
| 127 | LUPECCIO_(pisa) | 634 | MOLINARA |
| 530 | MACERATINO | 151 | MOLINELLI |
| 745 | MADALONA | 754 | MONDEUSE |
| 751 | MADDALENA_BRUNI | 457 | MONICA |
| 128 | MAGLIOCCO_CANINO_2 | 152 | MONICA_BIANCA |
| 636 | MAGLIOCCO_DOLCE | 970 | MONTE_MADONNA_1_GT |
| 130 | MAIOLICA_2 | 153 | MONTEPULCIANO |

987 NEBBIOLO ..... 200
545 NEGRETTO ..... 202
171 NEGRETTO_DI_SALUZZO ..... 778
440 NEGRO_AMARO ..... 204
758 NER DALA ..... 209
548 NERA GROSSA ..... 569
052 NERELLO_MASCALESE ..... 205
174 NERETTA_CUNEESE ..... 206
421 NERETTO DI BAIRO ..... 207
549 NERO_DI_VASCA ..... 572
175 NERONA ..... 1181
176 NEYRET ..... 210
770 NIBIO ..... 779
177 NIEDDERA ..... 211
769 NIEDDU_MANNU_1 ..... 212
54 NIGRA ..... 213
441 NOCCHIANELLO DI GAZZARETTO 1 ..... 780
178 NOCCHIANELLO_NERO ..... 781
179 NOCCHIANELLO_PELOSO ..... 782
768 NOCERA ..... 214
1014 NOCERONE DI MILAZZO 1 ..... 445
180 NOSIOLA ..... 215
553 NOTARDOMENICO_2 ..... 216
772 NURAGUS ..... 217
181 OLIVELLA NERA 2 ..... 1095

INC_BRUNI_185
INC_MALVASIA_X_TREBBIANO_TOSC ANO
INC_MANZONI_2_14
INCBIANCO_FEDIT_51_C_S_C
INCBRUNI_54
INCMANZONI_2_15
INCMATHIASZ_210
INCOGNITA
INCPIROVANO_105_VOLTA
NAME
PALLAGRELLO NERO

PALOMBA
Palomino
PAMPANARO GRANDE
PANFINONE
PANSE_BLANCHE
PANSE_PRECOCE
PANSE_ROSA_DIMALAGA
PASCALE
Passau
pattaresco
PAVANA
PECHINO
PECORELLO
PECORINO

535 MALBECH
537 MALBO GENTILE
538 MALIGIA_1
131 MALVASIA
132 MALVASIA_BIANCA_2
133 MALVASIA BIANCA_DI_CAN
3 DIA
423 MALVASIA_BIANCA_LUNGA
541 MALVASIA_DEL_LAZIO
1090 MALVASIA_DELLE_LIPARI COD NAME

190 PRECOCISSIMA MALIANI
787 PRIE_ROUGE
1024 PRIMITIVO
1060 PRIMITIVO 31 Pr
788 PRODEST
462 PROSECCO
463 PROSECCO_LUNGO
1102 PRUNESTA
223 PRUNESTA_BIANCA
$22{ }^{\mathrm{P}} \mathrm{A}$
PRUNESTA ROSSO VIOLACE A

783 QUAGLIANO
226 QUEEN
789 RABOSO_PIAVE
227 RABOSO_VERONESE
228 RAGUSANO

53 MONTONICO_BIANCO
154 MONTONICO_NERO
544 MONTONICO_PINTO
155 MONTU
156 MONUKKA
157 MORELLINO
158 mOSCATO
159 MOSCATO_BCASALESE
162 MOSCATO_BIANCO
COD NAME

807 SANTA MARIA
252 SANTO_STEFANO_BIANCO
253 SAUVIGNON
1012 SAUVIGNON_GROS
254 SAUVIGNON_VERT
255 SCACCIADEBITI
256 SCACCO
258 SCHIAVA
1112 SCHIAVA_GENTILE
475 SCHIAVA_GRIGIA
259 SCHIAVA_GROSSA
451 SChiAvotto
580 SCHIOCCOLETTO
260 SCHIOPPETTINO
982 SCIAGLIN
182 OLIVETTA BIANCA ..... 1111
998 OLIVETTA VIBONESE ..... 1109
1009 ORA ..... 485
194 ORSINA ..... 574
1097 ORTRUGO ..... 219
774 OSELETA ..... 220
1081 OSERIE_DU_FARODE ..... 786
565 OTTAVIANELLO ..... 221
184 PALLAGRELLO BIANCO ..... 189
COD NAME ..... COD
283 TINTILIA ..... 191
193 TINTORIA_2 ..... 833
300 TOCAI_FRIULANO ..... 301
829 TORBATO ..... 318
672 TRAMINER_AROMATICO ..... 319
706 TREBBIANO_ABRUZZESE_(biotipo_Sbag ..... 320
830 TREBBIANO_ABRUZZESE_(biotipo_Svag ..... 321
284 TREBBIANO GIALLO ..... 322
285 TREBBIANO_MODENESE ..... 323
286 TREBBIANO_ROMAGNOLO ..... 324
287 TREBBIANO_SPOLETINO ..... 325
594 TREBBIANO_TOSCANO ..... 326
1194 TREVISANA NERA ..... 328
302 TROISCINA ..... 1020
303 TUCCANESE ..... 992

PEDEVENDA
PEDICELLO_ROSSO PEDRO_XIMENES pelaverga PELAVERGA_PICCOLO PELAVERGA S GRATO PENSICATO
PERERA
PERGOLESE DI TIVOLI PERLA_DI_CSABA

PERLETTE
PERLONA
PERRICONE
PETIT_ROUGE
PETIT_VERDOT
PETITE_ARVINE
PICCOLA NERA PICOLIT

PICULIT_NERI
PIEDIROSSO
PIGNOLA
PIGNOLO
PINELLA
PINOT_BIANCO
PIZZUTELLO BIANCO
PIZZUTELLO NERO

| 229 | RAMONDA | 1190 | SCIASCINOSO |
| :---: | :---: | :---: | :---: |
| 575 | REBO | 261 | SCIASCINOSO |
| 1027 | RECANTINA | 590 | SCIMISCIA_1 |
| 230 | RED_GLOBE_1 | 480 | SCONOSCIUTA_MARTINETTI |
| 231 | RED_OHANEZ | 1115 | SELEZIONE_BRUNI_54 |
| 577 | REFOSCO_DAL_PEDROSSO | 262 | SEMIDANO |
| 792 | REFOSCO_DI_RAUSCEDO | 1015 | SEMILLON |
| 232 | REFOSCO_NOSTRANO | 263 | SENNEN |
| 447 | REGINA_DEI_VIGNETI | 468 | SGAVETTA |
| 464 | RETAGLIADO_BIANCO | 264 | SGIUSEPPE_BIANCO |
| 494 | RIBOLLA_GIALLA | 265 | SGIUSEPPE_NERO |
| 436 | RIESLING | 266 | SILA |
| 233 | RIESLING_BLAU | 675 | SIRIO |
| 234 | RIESLING_ITALICO | 267 | SLORENZO |
| 55 | RIMINESE | 810 | SLUNARDO |
| 235 | ROIETTO_GRISO | 815 | SMARTINO |
| 236 | RONDINELLA | 269 | SMICHELE |
| 237 | ROSA_TARDIVA | 816 | SOMARELLO_ROSSO |
| 238 | ROSETTA | 1191 | SOPERGA |
| 239 | ROSSA_DI_BITONTO | 452 | SPAGNA_BIANCA |
| 240 | ROSSESE | 271 | SULTANA |
| 241 | ROSSESE_BIANCO | 294 | SULTANINA_BIANCA |
| 242 | ROSSIGNOLA | 272 | SURBANO |
| 245 | ROSSO_DI_LECCE | 993 | SUSUMANIELLO |
| 465 | ROSSOLA_NERA | 273 | SYLVANER_VERDE |
| 448 | ROUSSIN | 275 | TAGLIAFERRO |


| 304 | TURCA | 329 |
| ---: | :--- | ---: |
| 304 | UCELUT | 330 |
| 295 | ULIVELLO | 1084 |
| 1126 | UNKNOWN | 823 |
| 596 | UNKNOWN | 534 |
| 305 | UNKNOWN | 183 |
| 306 | UVA_ACETO | 839 |
| 363 | UVA_CARNE | 841 |
| 307 | UVA_CARRIERI | 842 |
| 591 | UVA_DEI_VECCHI | 844 |
| 277 | UVA_DI_COLOGNE | 845 |
| 962 | UVA_DI_CRIMEA | 846 |
| 308 | UVA_DI_MORNICO | 837 |
| 593 | UVA_DI_TROIA | 332 |
| 278 | UVA_GATTA | 849 |
| 279 | UVA_LONGANESI | 331 |
| 280 | UVA_MELONA | 459 |
| 598 | UVA_POLCE | 1139 |
| 309 | UVA_PRUGNA | 868 |
| 836 | UVA_RARA | 333 |
| 1099 | UVA_REALE | 613 |
| 1098 | UVA_TOSCA | 334 |
| 820 | UVALINO | 361 |
| 310 | VALENTINO | 336 |
| 311 | VECCIUTELLO | 36 |
| 601 | VEGA |  |

304 TURCA ..... 329U04 UCIVELOT1084
126 UNKNOWN534839
363 UVARN842844846
308 UVA DI MORNICO33233159
98 UVA POLCE868
836 UVA_RARA613
098 UVA TOSCA361
601 VEGA ..... 336

| PLASSA | 249 | RUBI_RED | 454 | TANNAT | 312 | VELTLINER | 337 |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| PLISSONA | 803 | RUCHE | 276 | TAZZELENGHE | 313 | VELTLINER_ROSSO_PRECOCE | 1086 |
| POLLERA_NERA | 250 | S_ANNA_DI_LIPSIA | 469 | TEMPRANILLO | 314 | VERANO | 618 |
| POLLINO | 804 | S. TERESA | 995 | TERBASCI_cp2 | 602 | VERDACCHIO | 858 |
| POLOGHELLO | 581 | SAGRANTINO | 281 | TEROLDEGO | 315 | VERDANA_BIANCA | 975 |
| PORCINA | 582 | SAN_PIETRO | 827 | TERRANO | 316 | VERDANE | 1141 |
| PORCINA_BIANCA | 805 | SANGIOVESE | 119 | TIMORASSO | 317 | VERDEA | 338 |
| PORTOGHESE | 251 | SANGIOVESE_DEL_VERRUCC HIO | 592 | TINTIGLIA | 603 | VERDECA | 339 |
| NAME | $\begin{aligned} & \text { COD } \\ & \text { E } \end{aligned}$ |  |  |  |  |  |  |
| VERDELLO_DI_BRACCIANO | 859 |  |  |  |  |  |  |
| VERDESE | 341 |  |  |  |  |  |  |
| VERDICCHIO_BIANCO | 342 |  |  |  |  |  |  |
| VERDISO | 344 |  |  |  |  |  |  |
| VERDUSCHIA | 345 |  |  |  |  |  |  |
| VERDUZZO_FRIULANO | 346 |  |  |  |  |  |  |
| VERDUZZO_TREVIGIANO | 347 |  |  |  |  |  |  |
| VERMENTINO | 350 |  |  |  |  |  |  |
| VERMENTINO_NERO | 351 |  |  |  |  |  |  |
| VERNACCIA_DI_ORISTANO | 352 |  |  |  |  |  |  |
| VERNACCIA_DI_SGIMIGNANO | 353 |  |  |  |  |  |  |
| VERNACCIA_NERA | 354 |  |  |  |  |  |  |
| VESPAIOLA | 355 |  |  |  |  |  |  |
| VESPOLINA | 356 |  |  |  |  |  |  |
| VESPRINO | 963 |  |  |  |  |  |  |
| VICTORIA | 458 |  |  |  |  |  |  |


| VIEN_DE_NUS | 357 |
| :--- | ---: |
| VIOGNIER | 1021 |
| VITOUSKA | 358 |
| VUILLERMIN | 359 |
| WILDBACHER | 360 |
| WURZER | 645 |
| ZAPPOLINO | 862 |
| ZELEN | 1144 |
| ZIBIBBO | 470 |
| ZIMAVACCA | 1096 |
| ZINGARELLO | 622 |
| ZIZAK | 964 |
| ZUCCACCIO | 864 |

Supplementary Table 13. Country of origin, skin colour and usage of the 530 accessions used for the genetic characterization.

| NAME | ORIGIN | COLOUR | USAGE | NAME | ORIGIN | COLOUR | USAGE | NAME | ORIGIN | COLOUR | USAGE |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| ABBOTTABOTTI | CI | blanc | wine | BIANCUZZO | IT | blanc | wine | CATARRATTO_BIANCO_COMUNE | IS | blanc | wine |
| ABBUOTO | CI | noir | w/t | BICANE | WE | blanc | w/t | CATARRATTONE | IS | blanc | table |
| ABELLO | CI | blanc | wine | BIGOLONA | IT | blanc | wine | CAVECIA | NI | blanc | wine |
| ABRUSCO | CI | noir | wine | BLANC_DE_MORGEX | NI | blanc | wine | CAVRARA_NERA | NI | noir | wine |
| AGHEDENE | IT | blanc | wine | BOGGIONE_NERO | IT | noir | w/t | CENERENTE | CI | noir | wine |
| AGLIANICO | CI | noir | wine | BOGGIONE_ROSSO | IT | noir | wine | CESANESE_DEI_CASTELLI_ROMANI | CI | noir | wine |
| AGLIANICO_GRANA_GROSSA | SI | noir | wine | BOMBINO_BIANCO | CI | blanc | wine | CHARDONNAY | WE | blanc | wine |
| AHMEUR_MONARDE | WE | rose/noir | table | BOMBINO_NERO | SI | noir | w/t | CHASSELAS_BLANC | WE | blanc | w/t |
| ALBANA | CI | blanc | wine | BONAMICO | CI | noir | wine | CHENIN | WE | blanc | wine |
| ALBANA_BORDINI | CI | noir | wine | BONARDA | CI | noir | wine | CICINESE_GROSSO | CI | noir | wine |
| ALBANA_DEL_PANIERE | CI | blanc | wine | BONDA | NI | noir | wine | CILIEGIOLO | IT | noir | w/t |
| ALBANELLO | IS | blanc | wine | BORBOTTONE | IT | blanc | wine | CIMINNITA | WE | blanc | $\mathrm{w} / \mathrm{t} / \mathrm{r}$ |


| ALBARANZEULI_BIANCO | IS | blanc | wine | BOSCHERA | NI | blanc | wine | CLAIRETTE | WE | blanc | w/t |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| ALBAROLA | CI | blanc | w/t | BOSCO | NI | blanc | w/t | CLOTILDE_PROSPERI | IT | blanc | table |
| ALBAROSSA | NI | noir | wine | BOVALE | WE | noir | wine | COCCIATOSTA | IT | blanc | wine |
| ALEANTE_RIVALTO | CI | noir | wine | BRACCIOLA_NERA | CI | noir | wine | COCOCCIOLA | CI | blanc | w/t |
| ALEATICO | CI | noir | w/t | BRACHETTO | NI | noir | wine | CODA_DI_VOLPE_BIANCA | SI | blanc | w/t |
| ALICANTE_BOUSCHET | WE | noir | wine | BRAMBANA | IT | noir | wine | CODELUNGHE | IT | noir | wine |
| ALIONZA | CI | blanc | w/t | BUBBIA | IT | noir | w/t | COPETA | SI | noir | w/t |
| ALPHONSE_LAVALLEE | WE | noir | w/t/r | BURGHESANA | IT | noir | wine | CORBINELLA_PADOVANA | NI | noir | wine |
| AMERICA | AM | noir/blanc | wine | BUSSANELLO | NI | blanc | wine | CORBINONA_13_GT | NI | noir | wine |
| ANCELLOTTA | CI | noir | wine | CABERNET_FRANC | WE | noir | wine | CORINTO_BIANCO | EE | blanc | table |
| ANSONICA | CI | blanc | w/t | CABERNET_SAUVIGNON | WE | noir | wine | CORINTO_NERO | EE | noir | table |
| ANTINELLO | SI | blanc | wine | CACCIADEBITI | CI | blanc | wine | CORNACCHIA | CI | noir | wine |
| APRILLA | IT | noir | wine | CACCIO_DI_FERMO_1 | CI | blanc/noir | wine | CORNALLIN | NI | noir | wine |
| ARGUMANNU | IS | blanc | wine | CADDIU | IS | noir | w/t | CORNAREA | NI | noir | wine |
| ARNEIS | NI | blanc | wine | CAGNULARI | WE | noir | wine | CORNIOLA | IT | blanc | table |
| ARSILICO | CI | noir | wine | CALABRESE | SI | noir | wine | CORVINA | NI | noir | wine |
| ARVESINIADU | IS | blanc | w/t | CALAMARO | IT | noir | wine | CORVINA_ASOLO | NI | noir | wine |
| AVANA | NI | noir | w/t | CALORIA | CI | noir | wine | CORVINONE | NI | noir | wine |
| AVARENGO | NI | noir | w/t | CANAIOLO_NERO | CI | noir | wine | cove | NI | blanc | wine |
| AXINA_DE_FRANCIA | IS | noir | wine | CANINA_1_1 | CI | noir | wine | CREPALLOCCHI | CI | noir | wine |
| AXINA_DE_TRES_BIAS | IT | noir | wine | CANNAMELE | SI | noir | wine | CROATINA | NI | noir | wine |
| BARBERA | NI | noir | wine | CANNER_SEEDLESS | AM | blanc | table | CROVASSA | NI | noir | wine |
| BARBERA_BIANCA | NI | blanc | wine | CAPENA | CI | blanc | wine | CUNEUTE | CI | noir | wine |
| BARESANA | IT | blanc | table | CAPIBIANCHI | CI | noir/blanc | wine | DALL | UKN | ukn | ukn |
| BARSAGLINA | CI | noir | wine | CAPOBIANCO_BIANCO | CI | blanc | wine | DAMASCHINO | EE | blanc | wine |
| BECUET | WE | noir | wine | CAPOBIANCO_NERO | CI | noir | wine | DAMIANO | IT | blanc | wine |
| BEGUGNOL | UKN | blanc | wine | CAPOLONGO_1 | CI | blanc | wine | DATTIER_DE_BEYROUTH | WE | blanc | table |
| BELLINO | WE | noir | w/t | CAPOLUNGO | CI | blanc | wine | DELIGHT | AM | blanc | table |
| BELLONE | CI | blanc | wine | CAPRETTA | SI | blanc | wine | DIAMANT | EE | blanc | table |
| BERTINORA | IT | blanc | wine | CARDINAL | AM | rouge/noir | w/t | DIMIAT | WE | blanc | w/t |
| BERZAMINO | IT | noir | wine | CARICAGIOLA | IS | noir | wine | DINDARELLA | NI | noir | wine |
| BIANCHETTA_DI_BACEDASCO | CI | blanc | wine | CARIGNANO | CI | noir | wine | DIOLINOIR | CNE | noir | wine |
| BIANCHETTA_DI_DIOLO | CI | blanc | wine | CARMENERE | WE | noir | wine | DOLCIAME | IT | blanc | wine |
| BIANCHETTA_TREVIGIANA | NI | blanc | wine | CARRICANTE | IS | blanc | wine | DON_MARIANO | WE | noir | table |
| BIANCO_DALESSANO | SI | noir | wine | CASAVECCHIA | SI | noir | wine | DOUX_DHENRY | WE | noir | w/t |
| BIANCOLELLA | SI | blanc | wine | CASENTINO | CI | noir | wine | DURAPECCIO_DI_SLORENZO | IT | blanc | wine |


| BIANCONE | CI | blanc | wine | CASETTA | NI | noir | wine | DURELLA |
| :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- |
| BIANCONE_DELLANTELLA | CI | blanc | wine | CATALANESCA_BIANCA | SI | blanc | w/t | EARLY_MUSCAT |$\quad$ wine


| NAME | ORIGIN | COLOUR | USAGE | NAME | ORIGIN | COLOUR | USAGE |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| ERBALUCE | NI | blanc | w/t | INC_MALVASIA_X_TREBBIANO_TOSCANO | CI | blanc | wine |
| FALANGHINA | SI | blanc | wine | INC_MANZONI_2_14 | NI | noir | wine |
| FERTILIA | NI | noir | wine | INCBIANCO_FEDIT_51_C_S_G | CI | blanc | wine |
| FIANO_ROSA_LOCOROTONDO_2 | SI | blanc | wine | INCBRUNI_54 | CI | blanc | w/t |
| FLAVIS | NI | blanc | wine | INCMANZONI_2_15 | NI | noir | wine |
| FOGARINA | CI | noir | wine | INCMATHIASZ_210 | EE | blanc | table |
| FOGLIA_TONDA | CI | noir | wine | INCTERZI_N1 | NI | noir | wine |
| FOGLIONA | IT | blanc | w/t | INVERNENGA | WE | blanc | w/t |
| FORASTERA | WE | blanc | wine | INVERNESCA | IT | blanc | table |
| FORCESE_DASCOLI | CI | blanc | wine | INZOLIA_IMPERIALE | WE | noir | table |
| FORGIARIN | NI | noir | wine | INZOLIA_PARCHITANA | WE | blanc | table |
| FORSELLINA | NI | noir | wine | ITALIA | IT | blanc | table |
| FORTANA | CI | noir | wine | ITALICA | NI | blanc | wine |
| FOSCA | CI | blanc | wine | JULY_MUSCAT | AM | blanc | w/t |
| FRACIDELLA | CI | blanc | wine | KADARKA | EE | noir | wine |
| FRANCONIA | WE | noir | wine | KALILI | WE | blanc | table |
| FRAPPATO | IS | noir | wine | KERNER | CNE | blanc | wine |
| FREISA | NI | noir | wine | LACRIMA | CI | noir | wine |
| FUBIANO | NI | blanc | wine | LACRIMA_DI_MARIA | SI | blanc | table |
| FULIGNO | CI | noir | wine | LAGREIN | NI | noir | wine |
| FUMIN | NI | noir | wine | LAMBRUSCA_DI_ALESSANDRIA | NI | noir | wine |
| GABBA_VOLPE | UKN | blanc | wine | LAMBRUSCO_A_FOGLIA_FRAST | CI | noir | wine |
| GAGLIOPPO | SI | noir | wine | LAMBRUSCO_DI_SORBARA | CI | noir | wine |
| GALLICO | IT | ukn | wine | LAMBRUSCO_GRASPAROSSA | CI | noir | wine |
| GAMARET | WE | noir | wine | LAMBRUSCO_MAESTRI | CI | noir | wine |
| GAMAY | WE | noir | wine | LAMBRUSCO_MARANI | CI | noir | wine |
| GAROFANA | CI | noir | wine | LAMBRUSCO_MONTERICCO | CI | noir | wine |
| GATTA | IT | blanc/noir | wine | LAMBRUSCO_OLIVA | CI | noir | wine |
| GERUSALEMME | IT | blanc | table | LAMBRUSCO_SALAMINO | CI | noir | wine |


| GIRO | WE | blanc/noir/rouge | wine | LAMBRUSCO_VIADANESE | CI | noir | wine |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| GIUSTOLISI | IT | blanc | wine | LATTUARIO_NERO | EE | noir | table |
| GOLDTRAMINER | NI | blanc | wine | LINNEO_NERO | IT | noir | table |
| GOSEN | NI | noir | wine | LIVORNESE_BIANCA | CI | blanc | wine |
| GRANOIR | CNE | noir | wine | LUGLIOLA_ROSSA | CI | noir | wine |
| GRECHETTO_DI_TODI | CI | blanc | wine | LUMASSINA | NI | blanc | wine |
| GRECO_DI_TUFO | SI | blanc | w/t | MACERATINO | CI | blanc | w/t |
| GRECO_NERO_1 | SI | noir | w/t | MADALONA | IT | noir | wine |
| GRENACHE_NOIR | WE | noir | wine | MADDALENA_BRUNI | IT | blanc | table |
| GRIGNOLINO | NI | noir | wine | MAGLIOCCO_CANINO_2 | SI | noir | wine |
| GRILLO | SI | blanc | wine | MAGLIOCCO_DOLCE | SI | noir | wine |
| GRILLONE | IT | noir | wine | MAIOLICA_2 | CI | noir | wine |
| GROPPELLA | NI | noir | wine | MALBECH | WE | noir | wine |
| GROPPELLO_DI_MOCASINA | NI | noir | wine | MALBO_GENTILE | CI | noir | wine |
| GROPPELLO_DI_REVO | NI | noir | wine | MALIGIA_1 | UKN | blanc | wine |
| GROPPELLO_GENTILE | CI | noir | wine | MALVASIA | WE | blanc | wine |
| GROS_BOUSCHET | WE | noir | wine | MALVASIA_BIANCA_2 | SI | blanc | w/t |
| GROSPARIOL | IT | blanc | wine | MALVASIA_BIANCA_DI_CANDIA | CI | blanc | wine |
| GUARDAVALLE | SI | blanc | wine | MALVASIA_BIANCA_LUNGA | CI | blanc | wine |
| HARSLEVELU | EE | blanc | w/t | MALVASIA_DEL_LAZIO | CI | blanc | wine |
| IMPIGNO | SI | blanc | wine | MALVASIA_DELLE_LIPARI | SI | blanc | wine |


| NAME | ORIGIN | COLOUR | USAGE | NAME | ORIGIN | COLOUR | USAGE |
| :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- |
| MALVASIA_DI_CANDIA_AROMATICA | CI | blanc | w/t | MULLER_THURGAU | CNE | blanc | wine |
| MALVASIA_DI_CASORZO | NI | noir | wine | NASCETTA | NI | blanc | wine |
| MALVASIA_DI_SCHIERANO | NI | noir | wine | NASCO | IS | blanc | wine |
| MALVASIA_FURMHANN | CI | blanc | wine | NEBBIERA | NI | noir | wine |
| MALVASIA_ISTRIANA | EE | blanc | wine | NEBBIOLO | NI | noir | wine |
| MALVASIA_NERA_DI_BRINDISI_1 | CI | noir | w/t | NEGRETTO | CI | noir | wine |
| MALVASIA_NERA_LUNGA | NI | noir | wine | NEGRETTO_DI_SALUZZO | NI | noir | w/t |
| MAMMOLETTA | CI | noir | w/t | NEGRO_AMARO | SI | noir | wine |
| MAMMOLO | CI | noir | wine | NER_DALA | NI | noir | wine |
| MANTONICO_BIANCO | SI | blanc | wine | NERELLO_MASCALESE | SI | noir | wine |


| MANTONICONE | SI | noir | wine | NERETTA_CUNEESE | NI | noir |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| MANZONI_BIANCO | NI | blanc | wine | NERETTO_DI_BAIRO | NI | noir |
| MANZONI_MOSCATO | NI | noir | wine | NERO_DI_VASCA | CI | noir |
| MANZONI_ROSA | NI | rouge | wine | NEYRET | NI | noir |
| MARCHIONE_2 | SI | blanc | wine | NIBIO | IT | noir |
| MARSANNE | WE | blanc | wine | NIEDDERA | IS | noir |
| MARUGA | CI | noir | wine | NIEDDU_MANNU_1 | IS | noir |
| MARUGGIO_1 | UKN | blanc | wine | NIGRA | NI | noir |
| MARZEMINA_2 | NI | noir | wine | NOCCHIANELLO_DI_GAZZARETTO_1 | IT | blanc |
| MARZEMINO | CI | noir | w/t | NOCCHIANELLO_NERO | IT | noir |
| MATILDE | IT | blanc | table | NOCCHIANELLO_PELOSO | IT | blanc |
| MATURANO | UKN | blanc | wine | NOCERA | SI | noir |
| MAYOLET | NI | rouge | wine | NOCERONE_DI_MILAZZO_1 | SI | noir |
| MAZZESE | CI | noir | wine | NOSIOLA | NI | blanc |
| MELARA_1 | CI | blanc | wine | NOTARDOMENICO_2 | SI | noir |
| MERERA | IT | noir | wine | NURAGUS | IS | blanc |
| MERLOT | WE | noir | wine | OLIVELLA_NERA_1 | CI | noir |
| MICHELE_PALIERI | IT | noir | table | OLIVELLA_NERA_2 | CI | noir |
| MINNELLA_BIANCA_2 | IS | blanc | wine | OLIVETTA_BIANCA | CI | blanc |
| MOLINARA | NI | noir | wine | OLIVETTA_VIBONESE | CI | noir |
| MONICA | WE | noir | wine | ORA | WE | blanc |
| MONTE_MADONNA_1_GT | IT | blanc | wine | ORTRUGO | CI | blanc |
| MONTEPULCIANO | IT | noir | wine | OSELETA | NI | noir |
| MONTONICO_BIANCO | SI | blanc | w/t | OSERIE_DU_FARODE | WE | rose |
| MONTONICO_NERO | IT | noir | wine | OTTAVIANELLO | SI | noir |
| MONTONICO_PINTO | IT | noir | wine | PALLAGRELLO_BIANCO | SI | blanc |
| MONTU | CI | ukn | ukn | PALLAGRELLO_NERO | SI | noir |
| MONUKKA | EE | blanc/noir | t/r | PALOMBA | IT | noir |
| MORELLINO | CI | noir | wine | PALOMINO | WE | noir/blanc |
| MOSCATO_BCASALESE | NI | blanc | wine | PAMPANARO_GRANDE | CI | blanc |
| MOSCATO_BIANCO | IT | blanc | wine | PANFINONE | CI | blanc |
| MOSCATO_DADDA | IT | noir | table | PANSE_BLANCHE | WE | blanc |
| MOSCATO_DAMBURGO | CNE | noir | w/t | PANSE_PRECOCE | WE | blanc |
| MOSCATO_DI_SCANZO | NI | noir | wine | PANSE_ROSA_DIMALAGA | WE | rose |
| MOSCATO_DI_TERRACINA | CI | blanc | w/t | PASCALE | IS | noir |
| MOSCATO_GIALLO | IT | blanc | w/t | PASSAU | NI | noir |


| MOSCATO_NERO | NI | noir | wine | PATTARESCO | IT | noir | wine |
| :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- |
| MOSCATO_NERO_D_ACQUI | NI | noir | wine | PAVANA | NI | noir | wine |
| MOSCATO_ROSA_2 | NI | rouge | wine | PECHINO | IT | blanc | wine |
| MOSCATO_VIOLETTO | NI | noir | wine | PECORELLO | SI | blanc | wine |


| NAME | ORIGIN | COLOUR | USAGE | NAME | ORIGIN | COLOUR | USAGE |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| PEDEVENDA | NI | blanc | wine | REFOSCO_NOSTRANO | NI | noir | wine |
| PEDICELLO_ROSSO | CI | noir | wine | REGINA_DEI_VIGNETI | EE | blanc | table |
| PEDRO_XIMENES | WE | blanc | wine | RETAGLIADO_BIANCO | IS | blanc | w/t |
| PELAVERGA | NI | noir | table | RIBOLLA_GIALLA | NI | blanc | w/t |
| PELAVERGA_PICCOLO | NI | noir | wine | RIESLING | CNE | blanc | wine |
| PELAVERGA_S_GRATO | NI | noir | wine | RIESLING_ITALICO | IT | blanc | wine |
| PERERA | NI | blanc | wine | ROIETTO_GRISO | IT | gris | table |
| PERLA_DI_CSABA | EE | blanc | w/t | RONDINELLA | NI | noir | wine |
| PERLETTE | AM | blanc | t/r | ROSA_TARDIVA | IT | noir | table |
| PERLONA | IT | blanc | table | ROSSA_DI_BITONTO | SI | rose | table |
| PERRICONE | IS | noir | wine | ROSSESE | NI | noir | wine |
| PETIT_ROUGE | NI | noir | wine | ROSSESE_BIANCO | NI | blanc | wine |
| PETIT_VERDOT | WE | noir | wine | ROSSIGNOLA | NI | noir | wine |
| PETITE_ARVINE | CNE | blanc | wine | ROSSO_DI_LECCE | SI | red | wine |
| PICCOLA_NERA | NI | noir | wine | ROSSOLA_NERA | NI | noir | wine |
| PICOLIT | NI | blanc | wine | ROUSSIN | CNE | noir | wine |
| PICULIT_NERI | NI | noir | wine | RUBI_RED | AM | rouge | table |
| PIEDIROSSO | SI | noir | wine | RUCHE | NI | noir | wine |
| PIGNOLA | NI | noir | wine | S_ANNA_DI_LIPSIA | IT | blanc | table |
| PIGNOLO | NI | noir | wine | SAGRANTINO | CI | noir | wine |
| PINELLA | NI | blanc | wine | SALVATICO | CI | noir | w/t |
| PINOT_BIANCO | WE | blanc | wine | SAN_PIETRO | WE | blanc | wine |
| PIZZUTELLO_BIANCO | IT | blanc | table | SANGIOVESE | NI | noir | wine |
| PIZZUTELLO_NERO | WE | noir | table | SANGIOVESE_DEL_VERRUCCHIO | IT | noir | wine |
| PLASSA | NI | noir | wine | SANTA_MARIA | CI | blanc | wine |
| PLISSONA | IT | noir | wine | SAUVIGNON | WE | blanc | wine |
| POLLERA_NERA | CI | noir | wine | SAUVIGNON_GROS | WE | gris | wine |


| POLLINO | IT | blanc | wine | SCACCIADEBITI | CI | blanc | wine |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| POLOGHELLO | CI | blanc | wine | SCACCO | IT | blanc | wine |
| PORCINA_BIANCA | IT | blanc | wine | SCHIAVA | NI | noir | wine |
| PORTOGHESE | CNE | ukn | wine | SCHIAVA_GENTILE | CI | noir | wine |
| PRECOCISSIMA_MALIANI | IT | noir | wine | SCHIAVA_GRIGIA | NI | noir | wine |
| PRIE_ROUGE | NI | rouge | wine | SCHIAVA_GROSSA | NI | noir | w/t |
| PRIMITIVO | CI | noir | wine | SCHIOPPETTINO | NI | noir | wine |
| PRODEST | NI | noir | wine | SCIAGLIN | NI | blanc | wine |
| PROSECCO | NI | blanc | wine | SCIASCINOSO | CI | noir | wine |
| PROSECCO_LUNGO | NI | blanc | wine | SCIMISCIA_1 | NI | blanc | wine |
| PRUNESTA | SI | noir | w/t | SELEZIONE_BRUNI_54 | NI | blanc | wine |
| PRUNESTA_BIANCA | SI | blanc | w/t | SEMIDANO | IS | blanc | wine |
| PRUNESTA_ROSSO_VIOLACEA | SI | noir | table | SEMILLON | WE | blanc | wine |
| QUAGLIANO | NI | noir | table | SENNEN | NI | noir | wine |
| QUEEN | AM | noir | table | SGAVETTA | CI | noir | wine |
| RABOSO_PIAVE | NI | noir | wine | SGIUSEPPE_NERO | IT | noir | wine |
| RABOSO_VERONESE | CI | noir | wine | SILA | SI | blanc | wine |
| RAGUSANO | SI | blanc | wine | SIRIO | NI | blanc | wine |
| REBO | CI | noir | wine | SLORENZO | IT | noir | wine |
| RECANTINA | NI | noir | wine | SLUNARDO | SI | blanc | wine |
| RED_GLOBE_1 | AM | rouge | table | SMARTINO | IT | noir | wine |
| RED_OHANEZ | AM | red | table | SMICHELE | IT | noir | wine |
| REFOSCO_DAL_PEDROSSO | IT | noir | wine | SOMARELLO_ROSSO | SI | blanc | wine |


| NAME | ORIGIN | COLOUR | USAGE | NAME | ORIGIN | COLOUR | USAGE |
| :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- |
| SOPERGA | NI | noir | wine | VELTLINER | CNE | blanc | wine |
| SPAGNA_BIANCA | WE | blanc | wine | VERANO | CI | red | wine |
| STERESA | IT | blanc | w/t | VERDACCHIO | IT | blanc | wine |
| SULTANA | EE | blanc | table | VERDANA_BIANCA | IT | blanc | wine |
| SULTANINA_BIANCA | EE | blanc | w/t/r | VERDEA | CI | blanc | w/t |
| SURBANO | UKN | ukn | ukn | VERDECA | SI | blanc | wine |
| SUSUMANIELLO | SI | noir | wine | VERDELLO_DI_BRACCIANO | CI | blanc | wine |
| SYLVANER_VERDE | CNE | blanc | wine | VERDICCHIO_BIANCO | IT | blanc | wine |


| TAGLIAFERRO | CI | noir | wine | VERDISO | NI | blanc | w/t |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| TANNAT | WE | noir | wine | VERDUSCHIA | CI | blanc | wine |
| TAZZELENGHE | NI | noir | wine | VERDUZZO_FRIULANO | NI | blanc | wine |
| TEMPRANILLO | WE | noir | wine | VERDUZZO_TREVIGIANO | NI | blanc | wine |
| TERBASCI_cp2 | EE | noir | wine | VERMENTINO | CI | blanc | wine |
| TEROLDEGO | NI | noir | wine | VERMENTINO_NERO | CI | noir | wine |
| TERRANO | CI | noir | wine | VERNACCIA_DI_ORISTANO | IS | blanc | wine |
| TIMORASSO | NI | blanc | wine | VERNACCIA_DI_SGIMIGNANO | CI | blanc | wine |
| TINTILIA | SI | noir | wine | VERNACCIA_NERA | CI | noir | wine |
| TINTORIA_2 | WE | noir | wine | VESPAIOLA | NI | blanc | wine |
| TOCAI_FRIULANO | NI | blanc | wine | VESPOLINA | NI | noir | wine |
| TORBATO | WE | blanc | wine | VESPRINO | IT | blanc | wine |
| TRAMINER_AROMATICO | NI | rouge | wine | VICTORIA | EE | blanc | table |
| TREBBIANO_ABRUZZESE_1 | CI | blanc | wine | VIEN_DE_NUS | WE | noir | wine |
| TREBBIANO_ABRUZZESE_2 | CI | blanc | wine | VITOUSKA | EE | blanc | wine |
| TREBBIANO_GIALLO | CI | blanc | wine | VUILLERMIN | CNE | noir | wine |
| TREBBIANO_MODENESE | CI | blanc | wine | WILDBACHER | CNE | noir | wine |
| TREBBIANO_SPOLETINO | CI | blanc | wine | ZAPPOLINO | IT | noir | wine |
| TREBBIANO_TOSCANO | IT | blanc | wine | ZELEN | EE | blanc | wine |
| TREVISANA_NERA | NI | noir | w/t | ZIBIBBO | SI | blanc | w/t |
| TROISCINA | IT | ukn | wine | ZIMAVACCA | CI | blanc | table |
| TUCCANESE | SI | noir | wine | ZINGARELLO | CI | noir | wine |
| TURCA | NI | noir | wine |  |  |  |  |
| UCELUT | NI | blanc | wine |  |  |  |  |
| ULIVELLO | CI | noir | wine |  |  |  |  |
| UVA_ACETO | IT | blanc | wine |  |  |  |  |
| UVA_CARNE | IT | noir/rouge | wine |  |  |  |  |
| UVA_CARRIERI | IT | blanc | wine |  |  |  |  |
| UVA_DEI_VECCHI | CI | blanc | wine |  |  |  |  |
| UVA_DI_COLOGNE | WE | blanc | wine |  |  |  |  |
| UVA_DI_CRIMEA | IT | noir | wine |  |  |  |  |
| UVA_DI_TROIA | SI | noir | wine |  |  |  |  |
| UVA_GATTA | IT | noir | wine |  |  |  |  |
| UVA_LONGANESI | CI | noir | wine |  |  |  |  |
| UVA_MELONA | IS | rouge | table |  |  |  |  |
| UVA_PRUGNA | IT | blanc/noir | table |  |  |  |  |


| UVA_RARA | NI | noir | wine |
| :--- | :--- | :--- | :--- |
| UVA_REALE | CI | blanc | wine |
| UVA_TOSCA | CI | noir | wine |
| UVALINO | NI | noir | wine |
| VALENTINO | NI | noir | wine |
| VECCIUTELLO | CI | noir | wine |

Supplementary Table 14. Description of the 132 cultivars included in the GWAS diversity panel.

| NAME | Genotyped by | Country | Color | Usage | STR-3 | STR-6 |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| AHMEUR_MONARDE | CCC | WE | rose/noir | table | A-STR | a |
| ALBANA | CCC | CI | blanc | wine | ADMIXED | g |
| ALBARANZEULI_BIANCO | Laucou | IS | blanc | wine | A-STR | a |
| ALEANTE_RIVALTO | CCC | CI | noir | wine | A-STR | g |
| ALEATICO | Laucou | CI | noir | w/t | A-STR | g |
| ALICANTE_BOUSCHET | Laucou | WE | noir | wine | A-STR | g |
| ALPHONSE_LAVALLEE | Laucou | WE | noir | w/t/r | ADMIXED | b |
| ANCELLOTTA | Laucou | CI | noir | wine | B-STR | d |
| ANSONICA | Laucou | CI | blanc | w/t | A-STR | g |
| BELLONE | Laucou | CI | blanc | wine | A-STR | f |
| AXINA_DE_TRES_BIAS | CCC | IT | noir | wine | A-STR | g |
| BARBERA_BIANCA | CCC | NI | blanc | wine | A-STR | g |
| BARESANA | Laucou | IT | noir/rose | table | A-STR | a |
| CREPALLOCCHI | CCC | CI | noir | wine | B-STR | d |
| BIANCO_DALESSANO | Laucou | SI | noir | wine | A-STR | g |
| BOMBINO_BIANCO | Laucou | CI | blanc | wine | A-STR | f |
| BICANE | Laucou | WE | blanc | w/t | ADMIXED | g |
| BONAMICO | Laucou | CI | noir | wine | ADMIXED | g |
| BRACHETTO | Laucou | NI | noir | wine | ADMIXED | g |
| CABERNET_FRANC | Laucou | WE | noir | wine | ADMIXED | g |
| CAGNULARI | CCC | WE | noir | wine | ADMIXED | g |
| CALABRESE | Laucou | SI | noir | wine | ADMIXED | g |
| CANNER_SEEDLESS | CCC | AM | blanc | table | A-STR | a |
| CARDINAL | Laucou | AM | rouge/noir | w/t | ADMIXED | b |
| CARIGNANO | Laucou | CI | noir | wine | A-STR | g |
| FORTANA | CCC | CI | noir | wine | ADMIXED | d |
| CATALANESCA_BIANCA | Laucou | SI | blanc | w/t | A-STR | a |
| INC_MALVASIA_X_TREBBIANO_TOSCANO | CCC | CI | ukn | wine | A-STR | f |
| LAMBRUSCO_A_FOGLIA_FRAST | CCC | CI | noir | wine | B-STR | d |
| CHARDONNAY | Laucou | WE | blanc | wine | ADMIXED | g |
| CHASSELAS_BLANC | Laucou | WE | blanc | w/t | B-STR | g |
| CILIEGIOLO | Laucou | IT | noir | w/t | ADMIXED | g |
| CLAIRETTE | Laucou | WE | blanc | w/t | ADMIXED | g |


| COCOCCIOLA | Laucou | CI | blanc | w/t | A-STR | g |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| LAMBRUSCO_DI_SORBARA | Laucou | CI | noir | wine | B-STR | d |
| CORINTO_NERO | CCC | EE | noir | table | A-STR | g |
| LAMBRUSCO_MAESTRI | Laucou | CI | noir | wine | B-STR | d |
| LAMBRUSCO_MARANI | Laucou | CI | noir | wine | B-STR | d |
| CORNIOLA | Laucou | IT | blanc/noir | table | A-STR | a |
| LAMBRUSCO_SALAMINO | Laucou | CI | noir | wine | B-STR | d |
| CROATINA | Laucou | NI | noir | wine | ADMIXED | g |
| CROVASSA | CCC | NI | noir | wine | ADMIXED | g |
| DATTIER_DE_BEYROUTH | Laucou | WE | blanc | table | A-STR | a |
| DIMIAT | Laucou | WE | blanc | w/t | A-STR | g |
| DON_MARIANO | Laucou | WE | noir | table | A-STR | a |
| EARLY_MUSCAT | CCC | AM | blanc | w/t | ADMIXED | b |
| FALANGHINA | Laucou | SI | blanc | wine | ADMIXED | g |
| FLAVIS | CCC | NI | blanc | wine | B-STR | g |
| FORASTERA | Laucou | WE | blanc | wine | ADMIXED | g |
| GAMARET | Laucou | WE | noir | wine | B-STR | g |
| GAMAY | Laucou | WE | noir | wine | ADMIXED | g |
| GIUSTOLISI | Laucou | IT | blanc | wine | A-STR | g |
| GRECO_DI_TUFO | Laucou | SI | blanc | w/t | ADMIXED | g |
| GRILLO | Laucou | SI | blanc | wine | A-STR | g |
| GROSPARIOL | CCC | IT | blanc | wine | B-STR | d |
| HARSLEVELU | Laucou | EE | blanc | w/t | A-STR | g |
| IMPIGNO | Laucou | SI | blanc | wine | A-STR | g |
| INZOLIA_IMPERIALE | CCC | WE | noir | table | A-STR | a |
| ITALIA | Laucou | IT | blanc | table | ADMIXED | g |
| JULY_MUSCAT | Laucou | AM | blanc | w/t | ADMIXED | b |
| KADARKA | Laucou | EE | blanc/noir/gris/rose | wine | A-STR | g |
| KALILI | CCC | WE | ukn | table | A-STR | g |
| KERNER | Laucou | CNE | blanc | wine | B-STR | g |
| LAGREIN | Laucou | NI | noir | wine | B-STR | g |
| LAMBRUSCA_DI_ALESSANDRIA | CCC | NI | noir | wine | ADMIXED | g |
| ARNEIS | Laucou | NI | blanc | wine | B-STR | d |
| AVARENGO | Laucou | NI | noir | w/t | B-STR | d |
| CASETTA | Laucou | NI | noir | wine | B-STR | d |
| LAMBRUSCO_VIADANESE | CCC | CI | noir | wine | ADMIXED | g |


| LUMASSINA | Laucou | NI | blanc | wine | ADMIXED | g |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| MALBO_GENTILE | CCC | CI | noir | wine | ADMIXED | g |
| MALIGIA_1 | CCC | UKN | blanc | wine | ADMIXED | g |
| MALVASIA_FURMHANN | CCC | CI | blanc | wine | ADMIXED | c |
| MALVASIA_ISTRIANA | Laucou | EE | blanc | wine | ADMIXED | c |
| MAMMOLETTA | CCC | CI | noir | w/t | ADMIXED | g |
| CATARRATTO_BIANCO_COMUNE | Laucou | IS | blanc | wine | ADMIXED | f |
| PERRICONE | Laucou | IS | noir | wine | A-STR | f |
| MATILDE | Laucou | IT | blanc | table | ADMIXED | b |
| MONTONICO_NERO | CCC | IT | noir | wine | ADMIXED | g |
| MONUKKA | Laucou | EE | blanc/noir | t/r | A-STR | a |
| MOSCATO_DI_TERRACINA | Laucou | CI | blanc | w/t | A-STR | g |
| MOSCATO_GIALLO | Laucou | IT | blanc | w/t | A-STR | g |
| MULLER_THURGAU | Laucou | CNE | blanc | wine | B-STR | g |
| FORSELLINA | CCC | NI | noir | wine | B-STR | d |
| MONTEPULCIANO | Laucou | IT | noir | wine | A-STR | f |
| FREISA | Laucou | NI | noir | wine | B-STR | d |
| NIBIO | Laucou | IT | noir | wine | ADMIXED | g |
| NIEDDERA | Laucou | IS | noir | wine | ADMIXED | g |
| OLIVETTA_BIANCA | Laucou | CI | blanc | table | A-STR | a |
| OLIVETTA_VIBONESE | CCC | CI | noir | table | ADMIXED | a |
| ORA | Laucou | WE | blanc | table | A-STR | g |
| FUMIN | Laucou | NI | noir | wine | B-STR | d |
| PANSE_PRECOCE | Laucou | WE | blanc | w/t | A-STR | g |
| GRIGNOLINO | Laucou | NI | noir | wine | B-STR | d |
| PEDRO_XIMENES | Laucou | WE | blanc | wine | A-STR | a |
| PERLA_DI_CSABA | Laucou | EE | blanc | w/t | ADMIXED | g |
| PETIT_VERDOT | Laucou | WE | noir | wine | B-STR | g |
| PIEDIROSSO | Laucou | SI | noir | wine | ADMIXED | g |
| PRIMITIVO | Laucou | CI | noir | wine | ADMIXED | g |
| PICOLIT | Laucou | NI | blanc | wine | B-STR | d |
| RED_GLOBE_1 | Laucou | AM | rouge | table | A-STR | a |
| RED_OHANEZ | Laucou | AM | red | table | A-STR | a |
| RIESLING | Laucou | CNE | blanc | wine | B-STR | g |
| RIESLING_ITALICO | Laucou | IT | blanc | wine | B-STR | g |
| ROSSIGNOLA | Laucou | NI | noir | wine | ADMIXED | g |


| PIGNOLO | CCC | NI | noir | wine | B-STR | d |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| SAGRANTINO | Laucou | CI | noir | wine | ADMIXED | g |
| CODA_DI_VOLPE_BIANCA | Laucou | SI | blanc | w/t | ADMIXED | f |
| SCHIAVA_GRIGIA | CCC | NI | noir | wine | ADMIXED | g |
| RABOSO_PIAVE | Laucou | NI | noir | wine | B-STR | d |
| SCIMISCIA_1 | CCC | NI | blanc | wine | B-STR | g |
| SELEZIONE_BRUNI_54 | CCC | NI | blanc | wine | A-STR | g |
| SEMIDANO | Laucou | IS | blanc | wine | A-STR | g |
| MERLOT | Laucou | WE | noir | wine | B-STR | e |
| SULTANA | Laucou | EE | blanc | table | A-STR | a |
| TEMPRANILLO | Laucou | WE | noir | wine | A-STR | g |
| TERBASCI_cp2 | CCC | EE | noir | wine | A-STR | a |
| TORBATO | CCC | WE | blanc | wine | A-STR | a |
| SEMILLON | Laucou | WE | blanc | wine | B-STR | e |
| TREBBIANO_SPOLETINO | Laucou | CI | blanc | wine | ADMIXED | g |
| CHENIN | Laucou | WE | blanc | wine | B-STR | e |
| UVA_DI_TROIA | Laucou | SI | noir | wine | A-STR | g |
| RAGUSANO | CCC | SI | blanc | wine | A-STR | f |
| VELTLINER | CCC | CNE | blanc | wine | ADMIXED | g |
| TINTILIA | CCC | SI | noir | wine | ADMIXED | f |
| VERDEA | Laucou | CI | blanc | w/t | ADMIXED | g |
| VERDISO | Laucou | NI | blanc | w/t | B-STR | d |
| VERDUZZO_FRIULANO | Laucou | NI | blanc | wine | B-STR | g |
| VERMENTINO_NERO | Laucou | CI | noir | wine | ADMIXED | g |
| VERNACCIA_DI_SGIMIGNANO | Laucou | CI | blanc | wine | ADMIXED | g |
| VICTORIA | Laucou | EE | blanc | table | A-STR | g |
| ZAPPOLINO | CCC | IT | noir | wine | ADMIXED | g |

Supplementary Table 15. Concentration and quality ratio as estimated with the Nanodrop of the 37 additional samples of the CCC used to reach the 132 for the diversity panel.

| NAME | Concentration (ng/ul) | A260/A280 | A260/A230 |
| :--- | ---: | ---: | ---: |
| ALBANA | 36.42 | 1.813 | 1.472 |
| BARBERA_BIANCA | 18.69 | 1.827 | 1.296 |
| CAGNULARI | 12.62 | 1.742 | 0.869 |
| CROVASSA | 17.40 | 1.862 | 1.857 |
| FLAVIS | 26.10 | 1.749 | 1.175 |
| FORSELLINA | 34.12 | 1.808 | 1.770 |
| FORTANA | 7.30 | 1.537 | 0.342 |
| LAMBRUSCA_DI_ALESSANDRIA | 8.74 | 1.710 | 0.736 |
| LAMBRUSCO_A_FOGLIA_FRAST | 16.60 | 1.726 | 1.219 |
| LAMBRUSCO_VIADANESE | 18.67 | 1.804 | 1.530 |
| MALBO_GENTILE | 10.91 | 1.761 | 0.991 |
| TINTILIA | 32.32 | 1.792 | 1.450 |
| PIGNOLO | 25.51 | 1.789 | 1.179 |
| SCHIAVA_GRIGIA | 27.59 | 1.695 | 0.758 |
| SCIMISCIA_1 | 16.83 | 1.666 | 1.075 |
| TORBATO | 21.46 | 1.657 | 0.689 |
| VELTLINER | 18.54 | 1.773 | 1.188 |
| INZOLIA_IMPERIALE | 28.85 | 1.777 | 1.460 |
| EARLY_MUSCAT | 9.49 | 1.615 | 0.522 |
| CANNER_SEEDLESS | 26.32 | 1.743 | 1.137 |
| KALILI | 54.39 | NA | NA |
| CORINTO_NERO | 39.41 | NA |  |
| INC_MALVASIA_X_TREBBIANO_TOSCANO | 8.14 | 1.763 | 0.876 |
| MALIGIA_1 | 33.03 | 1.803 | 1.623 |
| MALVASIA_FURMHANN | 16.55 | 1.797 | 1.101 |
| SELEZIONE_BRUNI_54 | 12.84 | 1.674 | 0.706 |
| TERBASCI_cp2 | 11.54 | 1.602 | 0.532 |
| AXINA_DE_TRES_BIAS | 26.89 | 1.777 | 1.084 |
| ALEANTE_RIVALTO | 13.19 | 1.600 | 0.642 |
| CREPALLOCCHI | 24.79 | 1.782 | 1.299 |
| GROSPARIOL | 15.14 | 1.810 | 1.420 |
| ZAPPOLINO | 13.90 | 1.669 | 0.711 |
| RAGUSANO | 10.06 | 1.492 | 0.349 |
| MONTONICO_NERO | 16.30 | 1.709 | 0.896 |
| OLIVETTA_VIBONESE |  | 1.774 | 0.984 |
| AHMEUR_MONARDE | 1.577 | 0.453 |  |
| MAMMOLETTA |  | 1.726 | 1.005 |
|  |  |  |  |

Supplementary Table 16. Decay of linkage disequilibrium at 0.2 estimated with the 10 K SNPs on the diversity panel and corrected by kinship using the package LDcorSV (Mangin et al., 2012).

| Chromosome | bp |
| :--- | ---: |
| $\mathbf{1}$ | 85845 |
| $\mathbf{2}$ | 124009 |
| $\mathbf{3}$ | 73080 |
| $\mathbf{4}$ | 122093 |
| $\mathbf{5}$ | 82377 |
| $\mathbf{6}$ | 85783 |
| $\mathbf{7}$ | 63387 |
| $\mathbf{8}$ | 84859 |
| $\mathbf{9}$ | 53133 |
| $\mathbf{1 0}$ | 45839 |
| $\mathbf{1 1}$ | 68616 |
| $\mathbf{1 2}$ | 53121 |
| $\mathbf{1 3}$ | 62514 |
| $\mathbf{1 4}$ | 81352 |
| $\mathbf{1 5}$ | 63304 |
| $\mathbf{1 6}$ | 37407 |
| $\mathbf{1 7}$ | 104773 |
| $\mathbf{1 8}$ | 107425 |
| $\mathbf{1 9}$ | 77366 |
| Mean | 77699 |

Supplementary Table 17. Results of GWAS run on each year independently. No significant results were found with QTCAT software. In red are highlighted SNPs that are common between either two approaches or two traits within the same approach. ${ }^{*} F D R<0.05$; ${ }^{* *} F D R<0.01$ as obtained from the approach under which the association was found.

|  |  | Emmax | Gapit |
| :---: | :---: | :---: | :---: |
| FB | 2008 | Chr5_2541834* | Chr5_2541834** |
|  |  | Chr5_4058168* | Chr5_4058168* |
|  |  | Chr5_4194452* | Chr5_4194452 ** |
|  | 2011 | Chr5_1134974* |  |
|  |  | Chr5_2541834** |  |
|  | 2012 | Chr7_10408692 ** |  |
|  |  | Chr15_9638997 ** |  |
|  |  | Chr15_10198185 ** |  |
|  | 2013 | Chr16_2792486* |  |
| VB | 2010 |  | Chr15_15612783 * |
|  | 2013 |  | Chr18_29062596 ** |
| F-V | 2013 |  | Chr12_7294026* |
|  |  |  | Chr12_4097019 * |
|  |  |  | Chr17_2824109* |
|  |  |  | Chr17_2677298* |
|  |  |  | Chr18_29062596 ** |



 EMMAX.

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Thanks to the awesome \#rstats community from Twitter and the StackOverflow users for providing assistance for any kind of $R /$ statistics/computer science related problem.


[^0]:    No hit
    No hit
    Ankyrin repeat
    Ankyrin
    Protein kinase Xa21
    No hit
    No hit
    Protein kinase Xa21
    Receptor kinase TRKe
    Ankyrin repeat
    Ankyrin repeat protein
    No hit
    Seven in absentia SINA4
    No hit
    Ankyrin repeat
    CTV. 20
    Ankyrin repeat protein PUMILIO 5 (APUM5)
    F-box family protein
    Protease Do-like 2, chloroplastic DEGP2
    Retrotransposon protein
    Unknown protein Ribosomal protein S8 (RPS8B) 40S Protein transport protein Sec23A

    PRD1
    Pentatricopeptide repeat protein
    No hit
    Pentatricopeptide repeat-containing protein
    Vacuolar ATP synthase subunit F
    Inner membrane import protein Tic22, Chloroplast
    Xyloglucan endotransglucosylase/hydrolase 23
    Xyloglucan endotransglucosylase/hydrolase 15
    Unknown
    UDP-glucose:flavonoid 7-O-glucosyltransferase
    UDP-glucoronosyl/UDP-glucosyl transferase UGT75C1
    UDP-glucoronosyl/UDP-glucosyl transferase UGT75C1

[^1]:    Carrier protein, Mitochondrial
    Unknown protein myb domain protein 93 Cryptochrome 1
    Protein phosphatase 2C
    Hydrolase, alpha/beta fold
    14-3-3 protein GF14 epsilon (GRF10) Adenosine 5 '-phosphosulfate reductase 3

    DNA-binding protein
    Leucine-rich repeat family protein Unknown protein
    Dehydration-responsive protein
    RNA recognition motif (RRM)-containing protein
    Transparent testa 1
    WPP domain-associated protein
    No hit
    Cis-zeatin O-beta-D-glucosyltransferase Cis-zeatin O-beta-D-glucosyltransferase

    No hit
    UDP-glycosyltransferase 85A1
    No hit
    Phosphate-induced protein 1 Erg-1
    Phosphotyrosyl phosphatase activator PTPA (GB:X73478) Phosphotyrosyl phosphatase activator PTPA (GB:X73478)

    UDP-glycosyltransferase 85A1
    UDP-glycosyltransferase 85A1 Cis-zeatin O-beta-D-glucosyltransferase Cis-zeatin O-beta-D-glucosyltransferase

    No hit
    Phosphate-induced protein 1
    Phosphate-induced protein 1
    Phosphate-induced protein 1
    Phosphate-induced protein 1
    Phosphotyrosyl phosphatase activator (PTPA)
    F-box protein (FBW2)

