



Heterologous expression of cyanobacterial Orange Carotenoid Protein (OCP2) as a soluble carrier of ketocarotenoids in *Chlamydomonas reinhardtii*

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ABSTRACT

Photosynthetic organisms evolved different mechanisms to protect themselves from high irradiances and photodamage. In cyanobacteria, the photoactive Orange Carotenoid-binding Protein (OCP) acts both as a light sensor and quencher of excitation energy. It binds keto-carotenoids and, when photoactivated, interacts with phycobilisomes, thermally dissipating the excitation energy absorbed by the latter, and acting as efficient singlet oxygen quencher. Here, we report the heterologous expression of an OCP2 protein from the thermophilic cyanobacterium *Fischerella thermalis* (*FtOCP2*) in the model organism for green algae, *Chlamydomonas reinhardtii*. Robust expression of *FtOCP2* was obtained through a synthetic redesigning strategy for optimized expression of the transgene. *FtOCP2* expression was achieved both in UV-mediated mutant 4 strain, previously selected for efficient transgene expression, and in a background strain previously engineered for constitutive expression of an endogenous β -carotene ketolase, normally poorly expressed in this species, resulting into astaxanthin and other ketocarotenoids accumulation. Recombinant *FtOCP2* was successfully localized into the chloroplast. Upon purification it was possible to demonstrate the formation of holoproteins with different xanthophylls and keto-carotenoids bound, including astaxanthin. Moreover, isolated ketocarotenoid-binding *FtOCP2* holoproteins conserved their photoconversion properties. Carotenoids bound to *FtOCP2* were thus maintained in solution even in absence of organic solvent. The synthetic biology approach herein reported could thus be considered as a novel tool for improving the solubility of ketocarotenoids produced in green algae, by binding to water-soluble carotenoids binding proteins.

1. Introduction

Photosynthetic organisms need light to drive photosynthetic reactions, producing NADPH and ATP molecules which are essential to satisfy the metabolic demands of the cells. However, in natural environments, these organisms can easily experience rapid changes in light intensity and quality [1]. High light exposure can cause an over-excitation of the photosynthetic apparatus due to the excessive absorption of photons, that cannot be compensated by the capacity of Calvin-Benson cycle to regenerate ADP and NADP⁺. This condition is extremely harmful for the cell. Indeed, the excess of absorbed light cannot be properly used for charge separation, thus increasing the probability of energy transfer to oxygen molecules and finally leading to the formation of toxic reactive oxygen species (ROS) [1]. Excited and

highly reactive intermediates, like ROS, have a high oxidative potential that may damage the photosynthetic apparatus, leading to photodamage or even cell death [1]. To prevent or minimize generation of such oxidizing molecules, photosynthetic organisms have evolved a wide range of photoprotective mechanisms, among which the fastest is the so called non-photochemical quenching (NPQ), which acts through the thermal dissipation of excess energy [1–3].

Cyanobacteria and higher plants show significant differences at the level of their light harvesting systems, despite a conserved structure of the core complex of Photosystems. In cyanobacteria, intramembranous supramolecular assemblies of phycobiliproteins, called phycobilisomes (PBS), act as light harvesting antennae [4], whereas higher plants evolved transmembrane antenna proteins called Light Harvesting Complexes (LHC), which bind carotenoids and chlorophylls [5,6]. LHC

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complexes or LHC-like proteins can be found in most of the eukaryotic photosynthetic organism, except for glaucophytes, which retained the cyanobacterial PBS, and red algae, which exploit both PBS and LHC proteins [7,8]. Diversity in light harvesting systems caused also a different evolution of NPQ photoprotective mechanisms. Photosynthetic organisms where LHC or LHC-like proteins are present, require some specific subunits to trigger NPQ, as LHCSR (Light Harvesting Complex Stress-related) or their close relative LHCX (Light Harvesting Complex X) and/or PSBS (Photosystem II Subunit S) proteins [9–11]. Either LHCSR or PSBS subunits contain protonatable residues exposed to the lumen, which are responsible for quenching activation upon lumen acidification [12–14]. Differently, in the case of cyanobacteria, NPQ is triggered by an Orange Carotenoid-binding Protein (OCP) [15–20].

Three paralog families of OCP, called OCP1, OCP2, and OCPX have been identified so far: OCPX and OCP1 share a common ancestor, whereas OCP2 and OCP1 are the result of a second divergence [21,22]. Genomic analysis allowed identifying OCP1 as the most widespread OCP protein with members in almost every phylogenetic subclade of cyanobacteria [23]. On the contrary OCP2 is less frequently observed in cyanobacteria genomes and in some cases both OCP1 and OCP2 can be found encoded by the same genome [22]. The third class of OCP proteins, OCPX, is usually present in genomes where OCP1 or OCP2 cannot be found [24]. OCPX is more frequently observed compared to OCP2 especially in filamentous heterocyst-forming cyanobacteria [24]. OCP is a water-soluble protein which binds a single carotenoid molecule, senses blue-light, and interacts with PBS. In particular, OCP has a crucial role in the NPQ mechanism of cyanobacteria because the OCP-PBS interaction allows inducing thermal dissipation of excess energy absorbed by the latter [25,26]. 3-Hydroxyechinenone has been reported as the main carotenoid bound by OCP proteins, even if other types of carotenoids and ketocarotenoids, like echinenone, canthaxanthin and zeaxanthin were also found bound to these soluble protein [25]. Among these ligands ketocarotenoids were specifically reported to be essential for OCP photoactivation and its function in photoprotection [25,27]. Under strong blue-green light, OCP^O (orange form) is converted to OCP^R (red active form), which can trigger NPQ mechanism [28]. OCP2 and OCPX red active forms can spontaneously revert to OCP^O in the dark [22], whereas OCP1 needs a Fluorescence Recovery Protein (FRP) to accelerate OCP^R-to-OCP^O conversion reaction and for detaching from PBSs [29]. Interestingly, OCP2 and FRP genes were never found in genome where OCP1 was present [22]. OCP shows a modularity in structure; it contains an effector N-terminal domain (NTD), a sensor C-terminal domain (CTD) and a carotenoid. Both NTD and CTD present homologous gene in the majority of cyanobacteria genomes named helix carotenoid proteins (HCPs) in the case of NTD and CTDHs (CTD-like homologs) in the case of CTD [23]. The role of these proteins is still not clear at all but it is proposed that these NTD and CTD homologous can mix generating OCP-like protein with different photoprotective properties [15]. In recent years OCP function and role have acquired an increasing interest, both as a singlet oxygen (¹O₂) [15] and energy quencher in living cells, but also as an antioxidant molecule for pharmaceutical and cosmetic applications [30,31]. In biological systems, as plants but also animals or humans, excessive light exposure leads to the formation of ROS, that could damage biomolecules, affecting the integrity and stability of cells and tissues. Pathobiochemistry of several diseases of light-exposed tissues is indeed influenced also by these photooxidative processes [32]. For these reasons, research toward the discovery of new bioavailable antioxidant molecules is extremely active and requires the use of highly hydrophilic antioxidants. Water-soluble carotenoid binding proteins, in fact, might be suitable to this purpose, as they are able to confer hydrophilic properties to carotenoids, which otherwise would be relatively hydrophobic and embedded in biological membranes. In addition, it was recently reported the higher antioxidant power of OCP compared to vitamin C, tocopherol and isolated carotenoids [30]. Bourcier de Carbon and co-workers have reported the possibility to produce high amount of holo-OCP protein in *Escherichia coli* [31] however, to the best of our

knowledge, stable production in green microalgae has not been achieved yet. A “green” alternative to bacteria could indeed be represented by microalgae, which are well known as highly efficient CO₂-fixing organisms and can be cultivated in a circular economy approach, exploiting nutrients recovered from wastewaters [33,34]. Moreover, the possibility to efficiently manipulate metabolism in microalgae allows redirecting endogenous pathways toward the production of different molecules of interest and high-valuable compounds [35–38]. Recently the model organism for green algae, *Chlamydomonas reinhardtii*, was engineered to accumulate ketocarotenoids, as canthaxanthin and astaxanthin, suitable for human nutrition or livestock feed [36]. However, carotenoids are hydrophobic molecules requiring an organic solvent or amphiphilic environment to be maintained in solution.

In this work we investigate the expression of OCP2 from *Fischerella thermalis* (FtOCP2) in a microalgal host. OCP2 was chosen among the different OCP subunits due to its previously reported spontaneous ability to revert to OCP^O in the dark even in the absence of FRP [22]. *F. thermalis* is a thermophile cyanobacterium growing at temperature up to ~60 °C presenting in its genome genes for OCP2, HCP2, HCP4 and CTDH subunits [22,39]. FtOCP2 sequence was preferred to OCP2 subunits from other cyanobacteria being *F. thermalis* a thermophilic microorganism, suggesting a possible higher stability of the proteins encoded by its genome compared to the case of other mesophilic cyanobacteria.

Here, we report the expression and purification of a functional and photoactive FtOCP2 as a carotenoid carrier in *C. reinhardtii*, in order to improve carotenoids availability in aqueous solution upon extraction.

2. Materials and methods

2.1. Algal strains and culture conditions

C. reinhardtii UVM4 (UV-mediated mutant 4) strain [40] was used for all transformation experiments. Algal cells were cultivated in mixotrophic or photoautotrophic conditions, in Tris-acetate-phosphate (TAP) or High Salts (HS) minimal medium respectively [41,42], as described in the text. Liquid cultures were maintained in shake flasks, whereas agar-solidified medium in plates, at 25 °C and 100–150 μmol photons m⁻² s⁻¹ of continuous white light, unless otherwise stated. Algal growth and growth tests were conducted with different systems: shaking flasks, stirring flasks or Multi-Cultivator MC-1000 (Photon Systems Instruments, Czech Republic). Temperature was controlled to 25 °C while light intensities were varied as indicated in the text.

2.2. Construction of transformation vectors, cloning, transformation and mutant screening

The amino acid sequence of *F. thermalis* Orange Carotenoid-binding Protein 2 (hereafter FtOCP) (WP_009459388), was synthetically redesigned by codon optimization and intron spreading, to enhance transgene expression, as recently described [43]. The synthesized nucleotide sequence (Thermo Scientific) was cloned into pOpt2_PsaD_mVenus_Stag_Paro vector [44] to obtain a protein product flanked by the *C. reinhardtii* photosystem I subunit D (PsaD) chloroplast targeting peptide and a C-terminal mVenus encoding for Yellow Fluorescent Protein (YFP), fused with a Strep-tag® II flag, a short peptide (8 amino acids, WSHPQFEK), which binds with high selectivity to Strep-Tactin®, an engineered streptavidin. The CrBKT (*C. reinhardtii* β-carotene ketolase) gene (AY860820.1), synthetically redesigned as previously explained [36], was cloned both in pOpt2_PsaD_mVenus_Paro and pOpt2_PsaD_AadA vectors, in the latter fused to AadA (aminoglycoside-3'-adenyltransferase) gene (pOpt2_PsaD_CrBKT_AadA), conferring resistance to spectinomycin [45].

Stable nuclear transformation was carried out by glass beads agitation, as previously described [46] using 10 μg of linearized plasmid DNA. Selection of transformants was performed on TAP agar plates supplied with antibiotics (12 μg/mL for paromomycin and 200 μg/mL

for spectinomycin), for 5–7 days at $200 \mu\text{mol m}^{-2} \text{s}^{-1}$ light intensity. Antibiotic resistant colonies were then selected for target construct expression. 50 expressing colonies for both *CrBKT_YFP* and *CrBKT_AadA* were pre-screened visually for red coloration before further quantification. *FtOCP_YFP* antibiotic resistant colonies, instead, were picked to fresh plates and inoculated in 96-well microtiter plates with TAP medium, cultivating at $200 \mu\text{mol m}^{-2} \text{s}^{-1}$ light intensity until sufficiently dense. YFP fluorescence was measured in an Infinite PRO 200 plate reader (TECAN, Switzerland) with excitation at 509 nm and emission at 520–560, normalizing it to 720 nm absorbance (cell scattering), to determine the highest expressing strains. Nomenclature of strains generated in this work is reported in Table S1.

2.3. Protein extraction and SDS-PAGE

Total cells or purified proteins were separated by SDS-PAGE as described in [47]. Separated proteins were stained using Coomassie Brilliant Blue solution or were analyzed by immunodetection using anti-GFP (Green Fluorescent Protein) antibody (Sigma-Aldrich) and using an anti-rabbit Immunoglobulin G-Alkaline phosphatase conjugated secondary antibody (Sigma-Aldrich). Protein concentration of isolated proteins was performed measuring absorption at 280 nm by NanoDrop 2000 spectrophotometer (Thermo Scientific, Wilmington, DE, USA).

2.4. Pigment extraction and analysis

Pigments were extracted from 10^6 intact cells using 80% acetone, buffered with Na_2CO_3 . Pigment extracts were centrifuged at 15,000g and the supernatant loaded in HPLC at the same chlorophyll content (20 μl loaded at 0.01 $\mu\text{g}/\mu\text{l}$ concentration in acetone 80%) determined by absorption spectroscopy as previously reported [49]. In the case of isolated proteins, 25 μl of samples with the same absorption at 500 nm ($\text{OD} = 0.8$) were diluted to 200 μl in acetone buffered with Na_2CO_3 to obtain a final acetone concentration of 80%. Pigment extracts were then centrifuged at 15,000g before loading into HPLC. HPLC analysis was performed as described in [36] with the difference that here a Jasco LC-4000 Extrema HPLC system (Jasco, Italy) equipped with a C18 column (Synergi 4u Hydro-RP 80A, Phenomenex, USA) and 350–750 nm diode array detector was used. In particular, 15-min gradient of ethyl acetate (0 to 100%) in acetonitrile-water-triethylamine (9:1:0.01, vol/vol/vol) at a flow rate of 1.5 mL/min was applied [48]. Ketocarotenoids peaks were identified by comparing retention times and spectra to commercially available standards (CaroteNature GmbH, Switzerland) as reported in [36].

2.5. Growth analysis

Cell density of wild type and mutant strains was measured using Countess II FL Automated Cell Counter (Thermo Fisher Scientific, USA), whereas total dry biomass was evaluated by overnight lyophilization of washed cell pellets followed by gravimetric determination.

2.6. Confocal microscopy

The expression and subcellular localization of *FtOCP_YFP* were examined by confocal microscopy. Images were recorded on a Leica TCS-SP5 inverted confocal microscope (Leica Microsystem, Germany), equipped 63 \times oil immersion objectives and 2 Hybrid spectral detectors. mVenus (YFP) and chlorophyll were excited at 514 nm, whereas fluorescence emission was detected at 522–572 nm and 680–720 nm, for YFP and chlorophyll *a* respectively.

2.7. Protein extraction and purification

FtOCP_YFP protein was purified both from *ocp1* and *ocp1-bkt1* lines, exploiting the presence of the Strep-tag® II short peptide at the C-

terminal of the recombinant protein. 3 L of saturated culture of each line ($9 \cdot 10^{10}$ total cells) were pelleted and lysate in Buffer W (100 mM Tris/HCl, 150 mM NaCl, 1 mM EDTA at pH 8) and broken through a Cell Disruptor (Constant Systems Ltd, UK) at 16 kpsi. Crude cell lysate was pelleted at 1000 g to remove cell debris. Soluble recombinant *FtOCP_YFP* was purified by affinity chromatography, flowing the obtained supernatant through a Strep-Tactin® immobilized resin, following manufacture's protocol and eluting with 50 mM Biotin. Eluted solutions were concentrated through membrane spin columns (GE Healthcare, USA) to 1 mL each. Calculation of the fraction of holoprotein compared to the total *FtOCP* protein content was performed determining the molar ratio between YFP (fused to *FtOCP*) and carotenoids bound. Molar concentration of YFP and carotenoids were calculated from absorption spectra of isolated OCP samples, pigments extracted from recombinant OCP and isolated YFP, considering the extinction coefficients of carotenoids ($114,374 \text{ M}^{-1} \text{ cm}^{-1}$) and YFP ($83,000 \text{ M}^{-1} \text{ cm}^{-1}$), respectively at 496 nm and 514 nm, and the absorption ratios at 514/496 nm and 496/514 nm respectively for extracted carotenoids and for YFP. Nomenclature of isolated proteins is reported in Table S2.

2.8. Spectroscopic analysis and measurement of recombinant *Fischerella thermalis* Orange Carotenoid-binding Protein 2 photoconversion kinetics

Absorption spectra and kinetics were collected with Evolution 220 UV-Vis spectrophotometer (Thermo Scientific, USA). Photoconversion of dark-adapted *FtOCP_YFP* orange-inactive form to the red-active form was tested after 5 min of blue light illumination ($\lambda_{\text{max}} = 470 \text{ nm}$) at 900 $\mu\text{mol photons m}^{-2} \text{s}^{-1}$. The photoconversion and dark recovery kinetics were monitored from the absorption changes at 425–475 nm and 525–575 nm where the orange-inactive form (OCP^{O}) and the red-active form (OCP^{R}) are respectively mainly contributing.

2.9. Experimental replication and statistical treatment

All the experiments herein reported were performed at least three times. Errors are reported as standard deviations.

3. Results

3.1. Heterologous expression of *Fischerella thermalis* Orange Carotenoid-binding Protein 2

Codon optimization and incorporation of the first intron of *C. reinhardtii* Ribulose biphosphate carboxylase small chain 2 is a well-established technique to optimize nuclear transgene expression in this host organism [43]. This strategy was applied to generate an optimized synthetic gene, encoding for the 319 aa OCP2 peptide sequence from *F. thermalis* (Fig. S1). Optimized gene sequence was cloned into a pOpt2 vector: Photosystem I subunit D (PsaD) transit peptide was used as chloroplast targeting peptide, as previously reported [50], while mVenus was fused at the C-terminus of *FtOCP*, to allow screening of transgene expressing colonies. A Strep-tag® II affinity tag (WSHPQFEK*) was also fused at the C-terminus to allow protein purification. Hsp70A/Rbc2 hybrid promoter was used to drive gene expression as previously reported [43]. pOpt2_PsaD_*FtOCP_YFP* vector (Fig. 1a) was used to transform a *C. reinhardtii* UVM4 strain, a mutant strain previously generated by UV mutagenesis and screened for efficient expression of nuclear transgenes [40]. 96 transformed colonies grown on selective plates were screened for YFP fluorescence and 4 of them, showing the highest YFP fluorescence, were selected to be inoculated. The accumulation of *FtOCP_YFP* was confirmed in the transgenic cultures, herein called *ocp1-4*, by western blot analysis against YFP moiety, showing a 64 kDa protein band which is not present in their background UVM4 strain (Fig. 1d). As reported in Fig. 1b, YFP fluorescence correlates with protein accumulation. The localization of accumulated protein into the chloroplast was confirmed by confocal microscopy, observing the

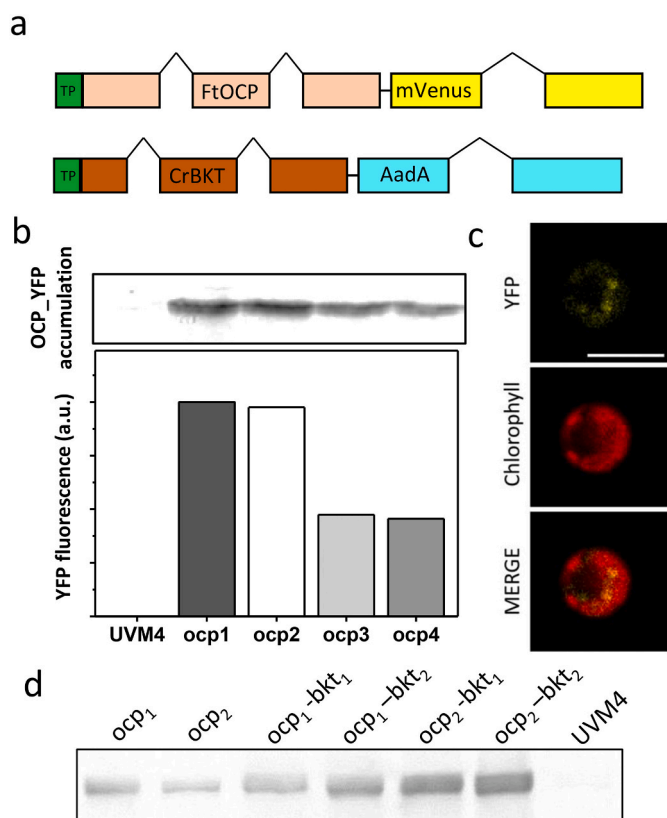


Fig. 1. FtOCP expression in *Chlamydomonas reinhardtii*: transformation, accumulation and localization a) schematic overview of vectors used for transformation of *C. reinhardtii* with *Fischerella thermalis* (Ft) OCP (Orange Carotenoid-binding Protein) and *C. reinhardtii* (Cr) BKT (β -carotene ketolase). Both vectors include Heat shock protein 70/Ribulose biphosphate carboxylase small chain 2 (Hsp70A/Rbcs2) hybrid promoter. FtOCP vector contains mVenus gene encoding for the YFP (Yellow fluorescence protein) fused to FtOCP, while CrBKT vector contains antibiotic resistance to spectinomycin (AadA, aminoglycoside-3'-adenyltransferase gene). PsaD (Photosystem I Subunit D) transit peptide (TP) was used to localize both BKT and OCP into the chloroplast. FtOCP and BKT sequences were designed in silico using codon optimization and inserting rbc2 intron 1. b) YFP fluorescence quantification, expressed as arbitrary unit (a.u.), and western blot analysis showing OCP_YFP accumulation in transformant strains *ocp*₁₋₄ compared to the background strain UVM4 (UV-mediated mutant 4). c) OCP_YFP cellular localization performed by confocal laser-scanning microscopy. Scale bar (white) represent 5 μ m. d) Western blot analysis showing OCP_YFP accumulation in transformant *ocp*₁ and *ocp*₂ lines as well as in *ocp*₁-bkt₁, *ocp*₁-bkt₂, *ocp*₂-bkt₁ and *ocp*₂-bkt₂ strains.

overlap between YFP fluorescence and chlorophyll autofluorescence (Fig. 1c). *ocp*₁ and *ocp*₂ were selected for next steps considering their higher protein accumulation, confirmed both by YFP fluorescence and western blot analysis.

3.2. Overexpression of *Chlamydomonas reinhardtii* β -carotene ketolase for ketocarotenoids accumulation

We have recently reported the possibility to selectively accumulate ketocarotenoids in *C. reinhardtii* by overexpressing an optimized endogenous β -carotene ketolase gene (*CrBKT*) [36]. In this mutant line, up to 70% of total carotenoids are converted to ketocarotenoids. The same strategy was here used to induce astaxanthin production in FtOCP_YFP transformed lines, considering the ability of OCP to bind ketocarotenoids [15,28]. In particular, the optimized CrBKT, deprived of the last 115 aa and targeted to chloroplast by PsaD transit peptide, was used as previously reported, fused to AadA gene, conferring resistance to spectinomycin [45]. This strategy allows to easily screen for

transformed colonies. The gene of interest and resistance marker in fact are expressed as a fused protein, thus the survival of colonies on selective medium ensure the presence of CrBKT. The ability of this fused protein to successfully convert precursors into ketocarotenoids was tested in UVM4 and compared to previously reported CrBKT_YFP. After the transformation with these two expression vectors, an orange phenotype was clearly visible in both cases (Fig. S2). All the colonies transformed with PsaD_BKT_AadA construct showed an altered coloring, even if with different intensity grades, due to the position effect of the genomic insertion of the cassette. Differently, in the case of PsaD_BKT_YFP transformation, only 13% of total colonies showed a red phenotype (Fig. S2). Ketocarotenoids production was then evaluated in the case of 50 colonies screened for having orange/red phenotype, obtained using PsaD_BKT_AadA or PsaD_BKT_YFP constructs, showing comparable accumulation of BKT enzyme products (Fig. S2). pOpt2-PsaD_BKT_AadA vector was thus used to transform *ocp*₁ and *ocp*₂ lines. Again, as observed for the previous UVM4 transformation on selection plates, most of the surviving colonies were characterized by a clear red/orange phenotype (Fig. 2a). Transformed strains generated using *ocp*₁ or *ocp*₂ background, renamed as *ocp*₁-bkt₁₋₂, and *ocp*₂-bkt₁₋₂, were selected and analyzed for ketocarotenoids accumulation (Fig. 2b). Ketocarotenoids accumulation in selected lines leads to a visible color change in liquid culture (Fig. S3) and the absorption spectra of extracted

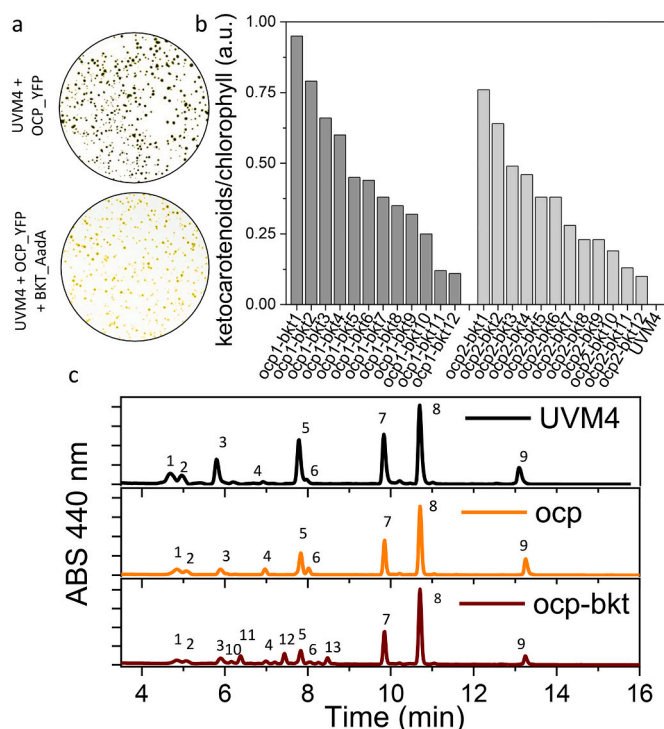


Fig. 2. Ketocarotenoids accumulation in transformed lines a) orange color phenotype of transformed lines in UVM4 (UV-mediated mutant 4) background expressing FtOCP (Orange Carotenoid-binding Protein from *Fischerella thermalis*) fused to YFP (Yellow Fluorescent Protein) alone or in addition to CrBKT (β -carotene ketolase from *Chlamydomonas reinhardtii*) fused to spectinomycin resistance (AadA, aminoglycoside-3'-adenyltransferase gene), indicated respectively as UVM4 + OCP_YFP or UVM4 + OCP_YFP + BKT_AadA. b) Ketocarotenoids accumulation expressed as arbitrary units (a.u.) for transformed lines expression both FtOCP and CrBKT (*ocp*₁-bkt₁₋₂ and *ocp*₂-bkt₁₋₂) compared to the background strain UVM4. c) Representative HPLC of pigments from UVM4 (black) *ocp* (orange) and *ocp*-bkt (red) strains. Chromatograms are shown reported the absorbance (ABS) of the different peaks at 440 nm. 1: neoxanthin; 2: lorenzoanthin; 3: violaxanthin; 4: antheraxanthin; 5: lutein; 6: zeaxanthin; 7: chlorophyll b; 8: chlorophyll a; 9 β -carotene; 10,11, 12: astaxanthin; 13: canthaxanthin. The data reported are representative of three independent experiments.

pigments revealed the presence of red shifted carotenoids, which are not present in UVM4, *ocp1* and *ocp2*. Pigments accumulated by top ketocarotenoids producing line, *ocp1-bkt1*, were further analyzed by high-performance liquid chromatography (HPLC) and compared to the background UVM4 and to the OCP only expressing lines (Fig. 2c; Table S3). HPLC analysis demonstrated similar pigment composition in UVM4 and *ocp1* lines, suggesting that the sole FtOCP expression could not alter the pigment profile in *C. reinhardtii* (Fig. 2c; Table S3). On the contrary, *ocp1-bkt1* lines showed the main representative peaks of astaxanthin (peaks 10, 11 and 12) and canthaxanthin (peak 13), as expected due to CrBKT enzyme activity [36] (Fig. 2c; Table S3). The three astaxanthin peaks identified in *ocp-bkt* lines are respectively related to the astaxanthin isomers as recently reported for the *C. reinhardtii* strains overexpressing the BKT enzyme [36].

3.3. Ketocarotenoids accumulation does not interfere with Orange Carotenoid-binding Protein accumulation

Possible effects of ketocarotenoids presence on FtOCP_YFP protein accumulation was evaluated by western blot on transformed lines with different amount of ketocarotenoids (*ocp1-bkt1*, *ocp1-bkt2*, *ocp2-bkt1* and *ocp2-bkt2*). As shown in Fig. 1d, all tested lines showed similar FtOCP_YFP protein accumulation, thus excluding negative or positive effects of CrBKT expression of recombinant FtOCP protein.

3.4. Orange Carotenoid-binding Protein and ketocarotenoids accumulation has no effects on growth

Effects of FtOCP_YFP and CrBKT on cell growth were also evaluated in both autotrophy (HS) and mixotrophy (TAP) conditions, using two different light regimes: 100 $\mu\text{mol photons m}^{-2} \text{s}^{-1}$ for control light and

500 $\mu\text{mol photons m}^{-2} \text{s}^{-1}$ for high light. Algae were grown in 80 mL airlift photobioreactors systems until reaching the stationary phase. As reported in Fig. S4, *ocp1* and *ocp1-bkt1* lines show no significant differences in biomass accumulation compared to WT in conditions of autotrophy or mixotrophy at both light intensities herein tested. These results demonstrate that OCP accumulation, neither in absence nor in presence of ketocarotenoids influenced *C. reinhardtii* growth.

3.5. Orange Carotenoid-binding Protein purification from *Chlamydomonas reinhardtii* cells

Purification of recombinant FtOCP proteins was then performed from *ocp1* and *ocp1-bkt1* lines grown in a semi-batch system in mixotrophy, at 300 $\mu\text{mol photons m}^{-2} \text{s}^{-1}$. Cells for both genotypes were harvested and lysed to release soluble proteins. OCP proteins were then purified by affinity chromatography. After elution from Strep-tag® II affinity column, recovered fractions were characterized by a yellow-orange color (Fig. 3a). FtOCP presence was confirmed by western blot analysis in eluted fractions purified from either *ocp1* or *ocp1-bkt1* (Fig. 3b) and absorption spectra of the eluted samples were collected, showing a main peak at 512–513 nm (Fig. 3c), which is ascribable to the YFP fused to FtOCP, as it can be seen from the comparison with the isolated YFP absorption spectrum in Fig. 3c. The spectra of purified proteins from *ocp1* (OCP) and *ocp1-bkt1* (OCP-BKT) show significant differences: OCP presents a blue shifted absorption with peaks at 436 nm and 476 nm while OCP-BKT shows a red-shifted tail in the 550–600 nm region, suggesting different carotenoid binding properties among the two samples. By subtraction of YFP signal it was possible to calculate the absorption spectra of OCP and OCP-BKT deprived of the contribution given by the fluorescence protein (Fig. S5). Fluorescence emission spectra at room temperature were also recorded (Fig. 3d): both OCP and

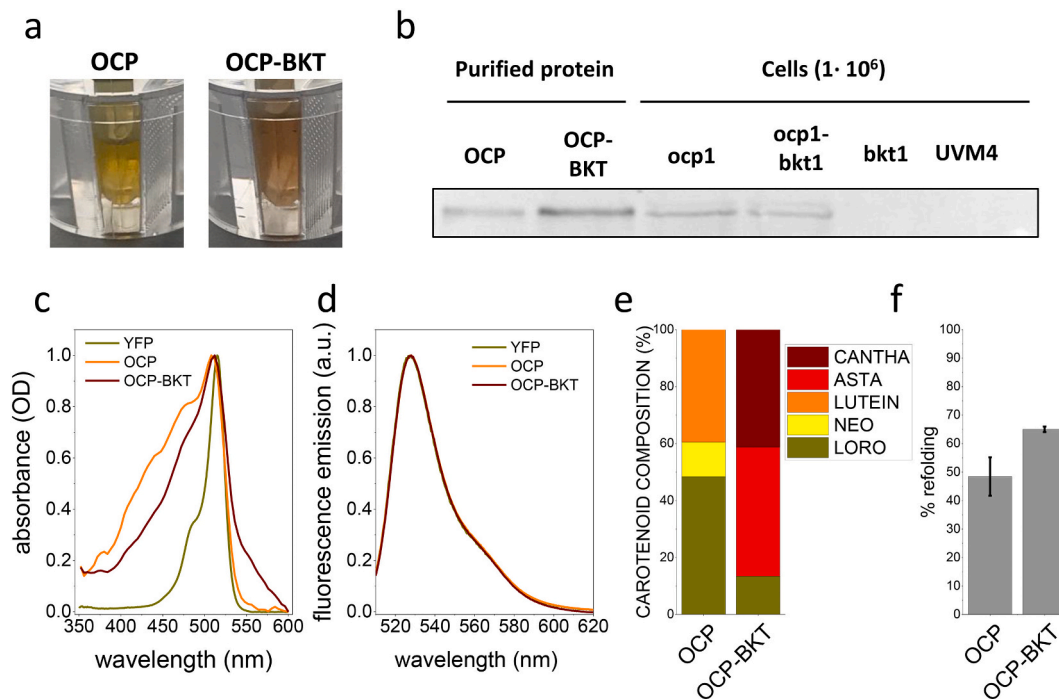


Fig. 3. Purification of OCP proteins a) red/orange color of purified OCP (Orange Carotenoid-binding Protein) proteins isolated from *Chlamydomonas reinhardtii* strains expressing recombinant OCP protein (OCP) or both BKT (β -carotene ketolase) and OCP (OCP-BKT); b) Western blot analysis performed by using specific antibody recognizing YFP (Yellow Fluorescent Protein) fused to OCP proteins in in isolated fractions after affinity chromatography (OCP and OCP-BKT) and in total protein extracts from the background strains UVM4 and from the transformant strains *ocp1*, *ocp1-bkt1* and *bkt1* expressing respectively OCP, OCP and BKT or BKT only; c) absorption spectra of isolated proteins reported as optical density (OD); d) fluorescence emission spectra of OCP proteins purified from *ocp1* (OCP), *ocp1-bkt1* (OCP-BKT) and YFP expressing lines reported as arbitrary units (a.u.); e) composition of carotenoids bound by OCP and OCP-BKT isolated proteins. Data are reported in percentage of total carotenoids. NEO: neoxanthin; CANTHA: canthaxanthin; LORO: loroxanthin; LUTEIN: lutein; ASTA: astaxanthin. f) percentage of refolded FtOCP holoproteins compared to total FtOCP isolated polypeptides. Error bars are reported as standard deviations ($n = 3$).

OCP-BKT fluorescence emission spectra perfectly overlap the YFP emission spectra, suggesting a correct folding of fused YFP. As reported in Fig. 3f the fraction of holoproteins compared to total isolated FtOCP protein content was ~50% and ~65% in the case of FtOCP isolated respectively from *ocp1* and *ocp1-bkt1* lines. The increased fraction of carotenoid binding protein in presence of ketocarotenoid (FtOCP isolated from *ocp1-bkt1*) is consistent with the preferential binding of ketocarotenoids rather than xanthophylls or carotenes to OCP subunits [15,22,27,31]. HPLC was used to evaluate pigments bound to the purified protein from either *ocp1* or *ocp1-bkt1* lines. FtOCP was shown to bind neoxanthin, lutein and zeaxanthin in the *ocp1* line, whereas FtOCP from *ocp1-bkt1* line was found to bind lutein, zeaxanthin and antheraxanthin (Fig. 3e). Interestingly, lutein was the xanthophyll molecule with the highest affinity for FtOCP, being the predominant carotenoid found in the holoproteins isolated from *ocp1*, while in presence of ketocarotenoids antheraxanthin was the ligand with the highest affinity (Fig. S6). Carotenoids bound by FtOCP are not soluble in water. Data reported in Fig. 4, indeed, show that after pigments extraction using ethanol and subsequent lyophilization, carotenoids bond to FtOCP cannot be solubilized in water but only in organic solvent. FtOCP protein, on the contrary, can be an efficient carrier for xanthophyll or ketocarotenoids in aqueous solution considering its stability in this buffer.

3.6. Purified Orange Carotenoid-binding Protein can switch from inactive to active form

The ability to be photoconverted into the active form was tested for purified FtOCP_YFP obtained from both *ocp1* and *ocp1-bkt1* lines. The conversion from OCP^O, inactive, to the active OCP^R has been reported to induce a red shift of the absorption spectrum of the protein [28]. Dark adapted purified proteins were thus illuminated using strong blue light. The conversion rate to the active form was monitored by measuring absorption spectra both at 425–475 nm and 525–575 nm (Fig. 5a, b). Indeed, these regions include respectively the absorption peaks of the inactive and the active form of OCP respectively. As shown in Fig. 5a and

c blue light illumination did not induce any significant change in the absorption spectrum of FtOCP_YFP purified from *ocp1*. Differently in the case of FtOCP_YFP purified from *ocp1-bkt1* blue light illumination caused a measurable reduction of absorption values in the 425–475 nm region and, at the same time, an increase in 525–575 nm region for (Fig. 5b, d, e). When blue light was switched off, an increase in the absorption at 425–475 nm and a simultaneous decrease in 525–575 nm region were recorded, indicating a spontaneously back-conversion from the active to inactive form (Fig. 5b, d, e). FtOCP_YFP purified from *ocp1*, does not bind ketocarotenoids as astaxanthin or canthaxanthin, and it is not able to photoswitch from the inactive to active form, differently from FtOCP_YFP purified from *ocp1-bkt1*. These results confirm the requirement of ketocarotenoids for the photoactivation of OCP proteins (Fig. 5f) [51].

4. Discussion

Carotenoids are antioxidant molecules which recently raised interest for their possible use as nutraceuticals for human health [52–55]. Ketocarotenoids as astaxanthin or canthaxanthin are, among carotenoid molecules, those with the highest antioxidant properties, with a high potential for their application in different industrial sectors. Microalgae are primary producers of carotenoids, including ketocarotenoids, being thus considered as potential platform for sustainable production of these molecules [52–60]. Differently from other potential heterotrophic sources of carotenoids, as bacteria or yeasts, microalgae have the important benefit of exploiting the photosynthetic process to sustain metabolism, converting light energy into chemical energy and leading to CO₂ fixation [61,62]. Carotenoids are essentially hydrophobic and require an amphiphilic carrier to be maintained in aqueous solution for industrial production and application. OCP is a small soluble protein able to bind one carotenoid molecule that could be suitable to this purpose, preventing carotenoids precipitation in aqueous environment (Fig. 4) [31]. OCP proteins in cyanobacteria are responsible for photo-protection, interacting with phycobilisomes and quenching the excess energy to prevent photodamage [63]. Several successful attempts to purify OCP have been performed in cyanobacteria [64] and its recombinant expression in bacteria has also been described [31]. To extend the range of possible platforms to produce carotenoid binding OCP proteins, here we report the overexpression and purification of FtOCP protein, a member of the OCP2 protein family [22], in *C. reinhardtii*. Using *C. reinhardtii* as a host for OCP expression permits to induce the formation of OCP holoproteins bound to different carotenoids, which are not available in cyanobacteria. Moreover, the powerful synthetic biology tools recently developed in this model organism allow the application of this strategy [43]. OCP protein was successfully accumulated in *Chlamydomonas* without any major effect on growth, both in mixotrophy and in autotrophy, even at high light intensity (Fig. S4).

OCP protein has been reported to bind mainly ketocarotenoids in cyanobacteria, with 3'-hydroxyechinenone being the main carotenoid bound [15,23,25,51]. Here we demonstrate that FtOCP can bind, in absence of ketocarotenoids, also lutein, zeaxanthin or lutein. In particular, lutein was the xanthophyll for which FtOCP showed the highest affinity (Fig. 3). These results were independent of the relative concentration of lutein among *C. reinhardtii* carotenoids, which is accumulated to a much lower level compared to other pigments, as lutein or β -carotene [65]. As mentioned above, the use of *C. reinhardtii* as host for OCP overexpression permit to exploit the synthetic biology tools available for this organism [36,37,43]: the expression of FtOCP in the *bkt* background allowed accumulating this protein in presence of ketocarotenoids, including astaxanthin (Figs. 2, 3). The results obtained from FtOCP expressed in *ocp1-bkt1* confirms that FtOCP preferentially binds ketocarotenoids (Fig. 3) and for the first time, to the best of our knowledge, demonstrate the possible direct binding of astaxanthin to a OCP2 subunit (Fig. 3). Indeed, while astaxanthin was reported to bind a OCP2 subunit (Fig. 3). Indeed, while astaxanthin was reported to bind a novel water-soluble carotenoid binding protein, named AstaP, found in

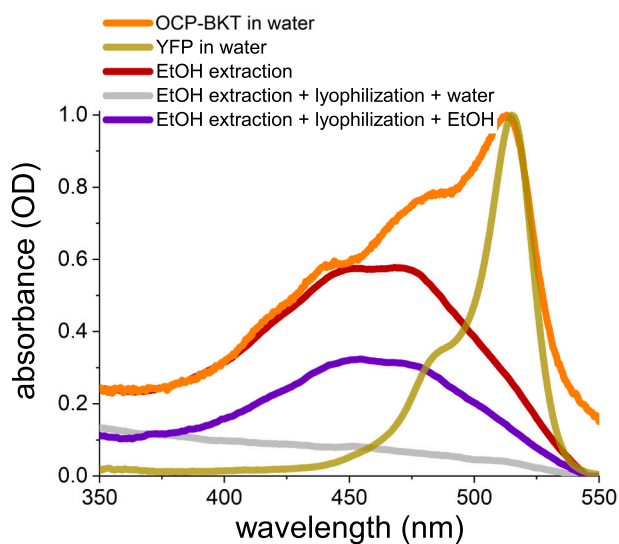


Fig. 4. OCP as a soluble carrier of carotenoids in water. Absorption spectra, reported as optical density (OD) of OCP-BKT (recombinant Orange Carotenoid-binding Protein, OCP, purified from *Chlamydomonas reinhardtii* strain expressing both OCP and BKT, β -carotene ketolase) in water (orange line), YFP (Yellow Fluorescent Protein) in water (gold line) and OCP-BKT after ethanol (EtOH) extraction (red line). Grey line and violet lines represent spectra of OCP-BKT after ethanol extraction which was then lyophilized and resuspended respectively in water (grey line) or in ethanol (violet line). The data reported are representative of three independent experiments.

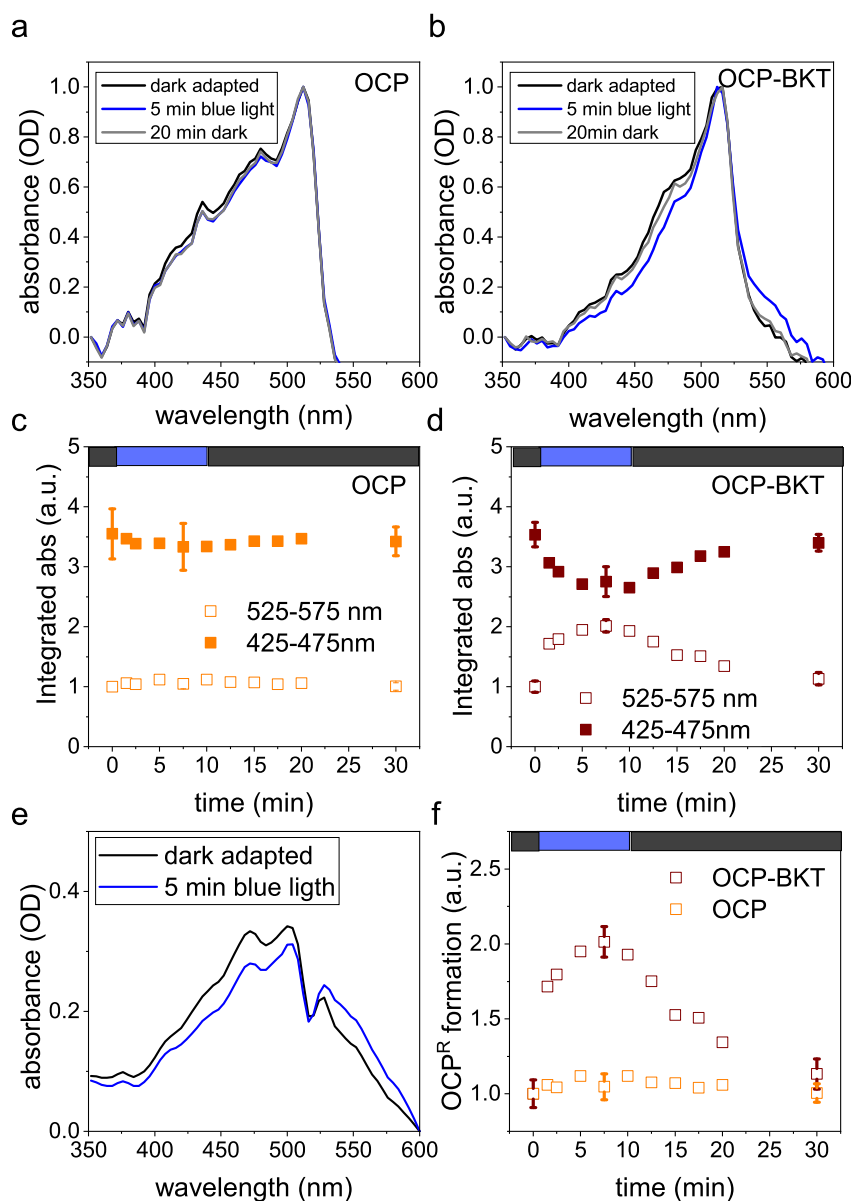


Fig. 5. Blue light excitation of OCP proteins a) b) Absorption spectra of recombinant OCP Orange Carotenoid-binding Protein) isolated from transformed *Chlamydomonas reinhardtii* strains expressing OCP only (ocp1 a, OCP) or both BKT (β -carotene ketolase) and OCP (ocp1-bkt1 b, OCP-BKT). Absorption spectra are reported as optical density (OD) for dark-adapted protein (black), after 5 min of blue light illumination (blue) and after 20 min of dark recovery (grey). c) d) Absorption spectra integrated in 525–575 nm range (active OCP^R) or 425–475 nm range (inactive OCP^D) reported as arbitrary units (a.u.). e) Absorption spectra of OCP-BKT after subtraction of YFP (Yellow Fluorescent Protein) signal for dark adapted sample and after 5 min of blue light illumination. f) Comparison of OCP^R formation (spectra integrated in 525–575 nm range) for OCP and OCP-BKT. Integrated absorption data reported in c), d) and f) were normalized to 1 in the case of 525–572 nm absorption of dark-adapted samples (time 0). Error bars are reported as standard deviation in c), d) and f) for dark adapted samples, after the blue light treatment and after dark recovery (n = 3).

an eukaryotic microalgal strain [66], the affinity of OCP1, OCP2 or OCPX for this ketocarotenoids was not investigated so far. As previously reported, only in the presence of ketocarotenoids, including astaxanthin, recombinant *FtOCP* behaved as a photoswitchable protein in presence of blue light (Fig. 5). Astaxanthin or canthaxanthin, or both, are thus able to allow OCP conformational change into the OCP^R form. Moreover, the ability to solubilize carotenoids in a water-based solution was confirmed also in the case of ketocarotenoid binding *FtOCP* (Fig. 4). Interestingly, neither accumulation of *FtOCP* nor ketocarotenoids biosynthesis influenced the biomass production rate of *C. reinhardtii* (Fig. S4). This result allows considering *C. reinhardtii* as an optimal host for the expression of OCP, which could act as a carrier of ketocarotenoids in aqueous solution. Metabolic engineering of *C. reinhardtii* for the production of astaxanthin has been recently reported as a promising step toward efficient and sustainable industrial production of ketocarotenoids for human nutrition, aquaculture or animal feed [36]. Overexpression of OCP in this strain allows further improvement of *C. reinhardtii* as ketocarotenoid production platform, opening the possibility to keep astaxanthin and canthaxanthin in aqueous solution upon extraction, for specific application requiring the absence of membranes or organic solvents.

However, proper optimization of OCP accumulation and its purification from engineered *C. reinhardtii* strains are required to make this process feasible at industrial scale. Possibly, other carotenoid binding proteins can be considered as soluble carrier for carotenoids/ketocarotenoids as for instance the C-terminal domain-like carotenoid proteins (CCPs): these subunits are homologs to the C-terminus of the OCP proteins and where recently characterized as a new class of soluble carotenoid binding proteins being able to bind canthaxanthin. Overexpression of CCPs subunits in *C. reinhardtii* might lead to a further carrier for ketocarotenoids [67]. It is also important to note that the ability to switch from the inactive to active form of *FtOCP* expressed in *bkt* background could allow the use of this holoprotein as a blue light-dependent molecular switch, to be used upon further engineering steps for regulating and studying gene expression or other cell activities [68].

CRediT authorship contribution statement

Matteo Pivato: Investigation, Data curation, Validation, Visualization, Writing - original draft; **Federico Perozeni:** Investigation, Data curation, Validation, Visualization, Writing - original draft; **Francesco**

Licausi: Conceptualization, Writing - review & editing; **Stefano Cazzaniga:** Investigation, Data curation, Writing - review & editing; **Matteo Ballottari:** Conceptualization, Funding acquisition, Supervision, Methodology, Validation, Visualization, Project administration, Writing - original draft, Writing - review & editing.

Declaration of competing interest

The authors declare that they have no known competing financial interests or personal relationships that could have appeared to influence the work reported in this paper.

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Appendix A. Supplementary data

Supplementary data to this article can be found online at <https://doi.org/10.1016/j.algal.2021.102255>.

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