

RESEARCH ARTICLE

Open Access

How has peritoneal dialysis changed over the last 30 years: experience of the Verona dialysis center

Gianluigi Zaza^{1*}, Carlo Rugiu¹, Alessandra Trubian¹, Simona Granata¹, Albino Poli² and Antonio Lupu¹

Abstract

Background: The last decade has witnessed considerable improvement in dialysis technology and changes in clinical management of patients in peritoneal dialysis (PD) with a significant impact on long term clinical outcomes. However, the identification of factors involved in this process is still not complete.

Methods: Therefore, to assess this objective, we retrospectively analyzed clinical records of 260 adult patients who started PD treatment from 1983 to 2012 in our renal unit. For the analysis, we divided them into three groups according to the time of starting dialysis: GROUP A (n: 62, 1983–1992), GROUP B (n: 66, 1993–2002) and GROUP C (n: 132, 2003 to 2012).

Results: Statistical analysis revealed that patients included in the GROUP C showed a reduction in mean patients' age ($p = 0.03$), smoking habit ($p = 0.001$), mean systolic blood pressure ($p < 0.0001$) and an increment in hemoglobin levels ($p < 0.0001$) and residual diuresis ($p = 0.016$) compared to the other two study groups. Additionally, patients included in GROUP C, mainly treated with automated peritoneal dialysis, showed a reduced risk of all-causes mortality and a decreased risk to develop acute myocardial infarction and cerebrovascular disease. Patients' age, diabetes mellitus and smoking habit were all positively associated with a significant increased risk of mortality in our PD patients, while serum albumin levels and residual diuresis were negatively correlated.

Conclusions: Therefore, the present study, revealed that in the last decade there has been a growth of our PD program with a concomitant modification of our patients' characteristics. These changes, together with the evident technical advances, have caused a significant improvement of patients' survival and a decrement of the rate of hospitalization. Moreover, it reveals that our pre-dialysis care, modifying the above-mentioned factors, has been a major cause of these clinical improvements.

Keywords: Continuous Ambulatory Peritoneal Dialysis, Automated peritoneal dialysis, Patient survival, Clinical outcomes

Background

Chronic kidney disease (CKD) is a progressive and irreversible deterioration of kidney function classified by the last international guidelines into five stages according to glomerular filtration rate [1]. In the last phase (end stage renal disease, ESRD) the kidney impairment is advanced and cellular/metabolic functions are significantly altered and unable to guarantee normal body homeostasis. Consequently, at this stage, renal replacement therapies (RRTs, peritoneal- or hemo-dialysis) or renal transplantation are needed to ensure patient's survival.

Although hemodialysis (HD) still represents the leading RRT, peritoneal dialysis (PD) procedure is utilized in more than 150,000 patients world-wide with a prevalence in Europe of 16% and in USA of 8% [2]. Recent data from the Italian Study Group of Peritoneal Dialysis have reported an incidence of this dialysis modality of approximately 20% with a prevalence of 15% [3,4].

PD seems to be a preferable choice for younger patients with high life expectancy and an elevated probability to undergo renal transplantation. In fact, this dialysis modality offers more flexibility allowing patients to continue working, a lesser cardiovascular impact and the maintenance of residual diuresis [5-7].

* Correspondence: Gianluigi.zaza@univr.it

¹Renal Unit, Department of Medicine, University-Hospital of Verona, Piazzale A. Stefani 1, 37126 Verona, VR, Italy

Full list of author information is available at the end of the article

However, peritoneal catheter and dialysis solutions (characterized by high concentration of glucose, glucose degradation products, low pH and high osmolality) used to remove waste products generated from normal metabolic processes, uremic toxins and to normalize body fluid and electrolytes [8] may still determine the systemic activation of a complex intracellular machinery leading to inflammation and oxidative stress [9-11]. These conditions may induce systemic non-infectious clinical complications including cardiovascular diseases.

Furthermore, recent studies have identified several demographic factors (e.g., age > 75 years at the start of dialysis, BMI < 18), clinical features (e.g., ischemic heart disease, anemia, heart failure and hemodynamic overload, cerebral vascular disease, peripheral vascular disease, diabetes) and metabolic causes (e.g., diabetes mellitus) associated with higher risk of mortality in PD patients compared with general population [12,13].

However, at the moment, only few reports have pointed out on the identification of risk factors for long-term clinical complications in PD patients living in Italy or other European countries [14-16]. It is unquestionable that geographically-related characteristics (e.g., diet, health care system), enhancing risk factors, may influence long term clinical outcomes.

Therefore, although monocentric, our study, performed on a large Italian cohort of PD patients (including 260 patients followed by our Renal/Dialysis Unit from 1983 to 2012), has been undertaken to identify changes across the last 30 years, to select clinical elements possibly predicting patients' survival and to recognize targets of intervention useful to minimize the onset and development of severe dialysis-associated clinical complications.

Methods

Patients

In this study, we retrospectively analyzed clinical records of 260 adult patients (older than 18 years) who started PD at the Renal Unit of the Hospital-University of Verona from 1983 to 2012.

Patients with fewer than 6 months' follow-up, who had been on HD or receiving a kidney graft before starting PD and had experienced at least one major cardiovascular event (myocardial infarction or cerebrovascular events, or who have undergone amputation) before dialysis were excluded.

Patients were, then, divided into 3 groups according to the time of starting PD:

- GROUP A (n: 62): patients starting PD from 1983 to 1992;
- GROUP B (n: 66): patients starting PD from 1993 to 2002;

- GROUP C (n: 132): patients starting PD from 2003 to 2012.

For all patients, we collected main demographic and clinical characteristics (Table 1). Estimated glomerular filtration rate (eGFR) has been calculated according to Cockcroft-Gault equation.

The study was carried out according to Declaration of Helsinki principles and approved by the Institutional Ethic Review Board of the University of Verona, Italy. All patients signed an informed consent.

Statistical analysis

All data were expressed as frequencies, percentages and mean values or median according to the variables' distribution. Parametric (*t*-test, ANOVA) and non-parametric (Mann-Whitney, Kruskal Wallis test) test and chi-square test/fisher test were used to assess differences in clinical and demographic features. Incidence rates of adverse events were calculated using the Poisson distribution and survival analysis was performed using Kaplan-Meier. Cox regression model was used to estimate the risk of acute myocardial infarction, cerebrovascular disease and vasculopathy. Tests have been corrected for baseline confounding factors as required. All tests were considered significant when $p < 0.05$. All statistical analysis were performed using SPSS, version 11.5.

Results

Demographic and clinical characteristics overtime

As showed in Table 1, we found significant differences in several demographic and clinical variables among the three study periods.

Statistical analysis revealed a significant and progressive reduction of the mean patients' age ($p = 0.03$), smoking habit ($p = 0.001$), mean systolic blood pressure ($p < 0.0001$) across the study periods. On the contrary, hemoglobin levels ($p < 0.0001$) and residual diuresis ($p = 0.016$) were increasing overtime.

Choice of PD methods

From 1983 to 2012, we found a clear trend toward increased number of PD patients undergoing automated peritoneal dialysis (APD) ($p < 0.0001$). This could reflect a change of the demographic characteristics of our PD population (younger and still able to work) (Figure 1A).

Waiting list for renal transplantation

During time, we found a significant increased number of PD patients on active renal transplant waiting list ($p < 0.0001$). This reflects a change of our center policy (mainly in the last decade) to use PD preferentially in younger patients without major comorbidities (e.g., cardio-vascular diseases, malignancies) (Figure 1B).

Table 1 Trends in demographic and clinical characteristics among the three study periods

| | GROUP A 1983-1992 (n = 62) | GROUP B 1993-2002 (n = 66) | GROUP C 2003-2012 (n = 132) | p value |
|--|-------------------------------|-------------------------------|--------------------------------|---------|
| Follow-up months mean (SD) | 29.3 (25.1) | 36.6 (27.4) | 33.7 (26.9) | 0.298 |
| *Age years mean (SD) | 63.5 (9.0) | 59.7 (12.7) | 59.0 (12.0) | 0.038 |
| Gender | | | | |
| Males n. (%) | 36 (58.1) | 46 (69.7) | 94 (71.2) | 0.174 |
| Females n. (%) | 26 (41.9) | 20 (30.3) | 38 (28.8) | |
| Smoking habits | | | | |
| No smokers n. (%) | 51 (82.2) | 39 (59.1) | 77 (58.3) | 0.001 |
| Ex smokers n. (%) | 7 (11.3) | 11 (16.7) | 38 (28.8) | |
| Smokers n. (%) | 4 (6.5) | 16 (24.2) | 17 (12.9) | |
| *Diabetes | | | | |
| No diabetes n. (%) | 44 (71.0) | 52 (78.8) | 102 (77.3) | 0.532 |
| Diabetes n. (%) | 18 (29.0) | 14 (21.2) | 30 (22.7) | |
| *Residual diuresis ml/die mean (SD) | 777.8 (560.0) | 924.2 (460.4) | 997.7 (469.2) | 0.016 |
| *Systolic blood pressure mmHg mean (SD) | 159.6 (25.2) | 154.0 (19.9) | 140.5 (15.4) | <0.0001 |
| *Albumin g/dl mean (SD) | 3.4 (0.49) | 3.2 (0.5) | 3.9 (2.8) | 0.06 |
| *Hb g/dl mean (SD) | 8.7 (1.5) | 9.6 (7.5) | 11.7 (1.36) | <0.0001 |
| *eGFR mean (SD) | 6.1 (1.9) | 6.4 (2.4) | 6.9 (2.6) | 0.076 |

eGFR: estimated Glomerular Filtration Rate. (*) At the start of PD treatment.

All-causes mortality risk

Multivariate analysis (Figure 2) showed that patients' age (hazard ratio (HR): 1.1, $p = 0.001$), diabetes mellitus (HR: 3.9, $p < 0.001$) and smoking habit (HR: 1.3, $p < 0.001$) were all positively associated with an increased risk of all-causes mortality in our PD patients' population, while serum albumin levels (HR: 0.6, $p = 0.001$) and residual diuresis (HR: 0.9, $p = 0.04$) were negatively correlated.

Gender, systolic blood pressure, hemoglobin levels and eGFR were not associated with patients' survival.

Kaplan-Meier revealed a better survival rate in PD patients of GROUP C compared to those included in the other two study-groups (Figure 3). Similar results were obtained evaluating transplantation and change in dialysis modality as competing outcomes and risks (Additional file 1: Table A and B, Figure S1). In particular, the statistical analysis confirmed a significantly higher risk of death in GROUP A versus both GROUP B and C ($p < 0.0001$) and an higher probability to undergo renal transplantation in last study period (GROUP C) compared to the other two periods (GROUP B and A) ($p < 0.0001$). On the contrary the risk to switch from PD to HD was similar among the three groups.

Interestingly, analyzing the trajectories of some of the above-mentioned clinical variables over-time (5 years), hemoglobin levels resulted constantly higher in GROUP C compared to B and A (Additional file 1: Table C).

This could mainly be due to a complete correction of anemia (with a more rational use of ESAs) in the last 10 years.

Risk of cardiovascular diseases

As shown in Figure 4, there was a gradual reduction in the risk of developing acute myocardial infarction and cerebrovascular complications overtime.

We did not find any statistically significant difference in the risk of vascular disease.

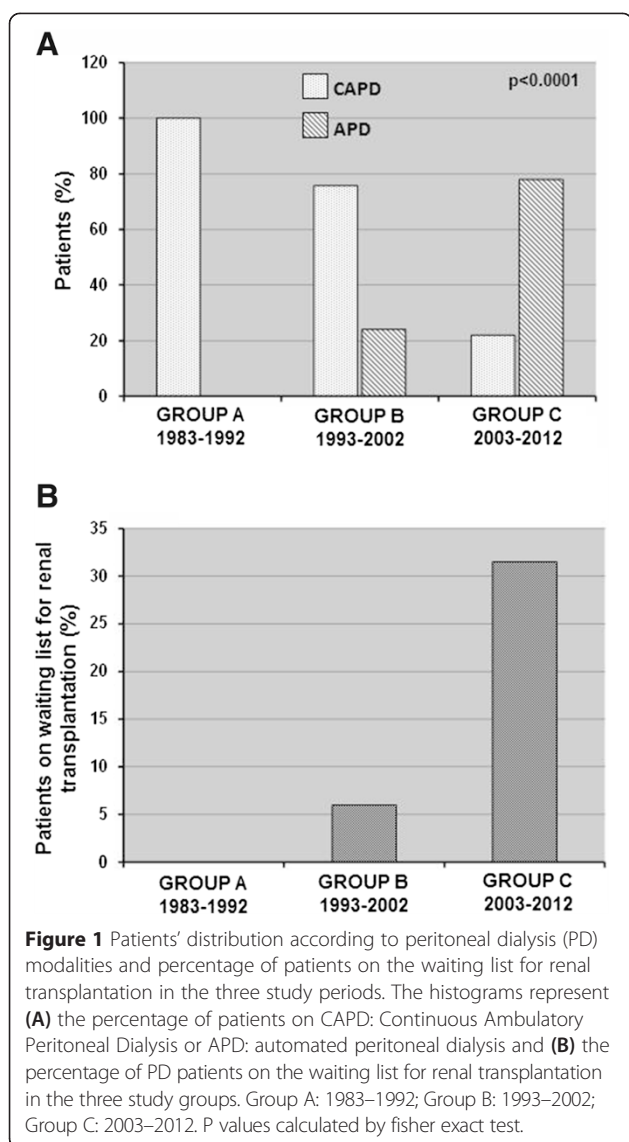
Days of hospitalization and complications

Interestingly, patients included in GROUP C showed a lower period of overall hospitalization for all causes and peritonitis compared to Group B and A (Figure 5).

The major causative agents of peritonitis were *Staphylococcus Aureus* and *Epidermidis* followed by *Pseudomonas Aeruginosa* (Table 2). This condition was similar in the 3 study groups.

Technique and catheter survival after one episode of peritonitis were considerably higher in the last study period (GROUP C: 2003–2012) compared to the other two periods.

Overall technique survival was similar in the 3 groups. No catheter has been surgically removed and re-inserted after a peritonitis episode (according to Hospital policy).



Discussion

Peritoneal dialysis (PD), although considered a valuable alternative renal replacement therapeutic option to hemodialysis (HD) in a great number of CKD patients (mainly young), is still associated with the development of severe long-term clinical complications (e.g., cardiovascular diseases) leading to significant reduction of patients' survival [17].

Therefore, in the last decade, researchers and clinicians worldwide had work together to avoid or minimize these complications by introducing more "biocompatible" fluids (with more physiologic pH and reduced glucose degradation products) and by ameliorating the selection of eligible patients for this dialysis procedure [18,19].

Additionally, several strategies have been undertaken to minimize, particularly during the pre-dialysis follow-up period, all corrigible factors known to be associated

with worst clinical outcomes of PD patients [20]. However, the complete identification of these elements represents a major target in nephrology.

To this purpose, we retrospectively analyzed our medical records regarding a large CKD patients cohort (n:260) starting PD treatment from 1983 to 2012.

Interestingly, several factors were significantly associated with an increased risk of mortality in our PD patients. Multivariate analysis showed that patients' age, diabetes mellitus and smoking habit were all positively associated with an increased risk of all-causes mortality in our PD patients' population.

The impact of age on survival is still debated. In fact, as a continuous home-based therapy, PD offers several potential advantages for older people, and it remains an important modality of renal replacement therapy, but patients with advanced age have an high risk to develop clinical complications and undergo peritonitis [21-24]

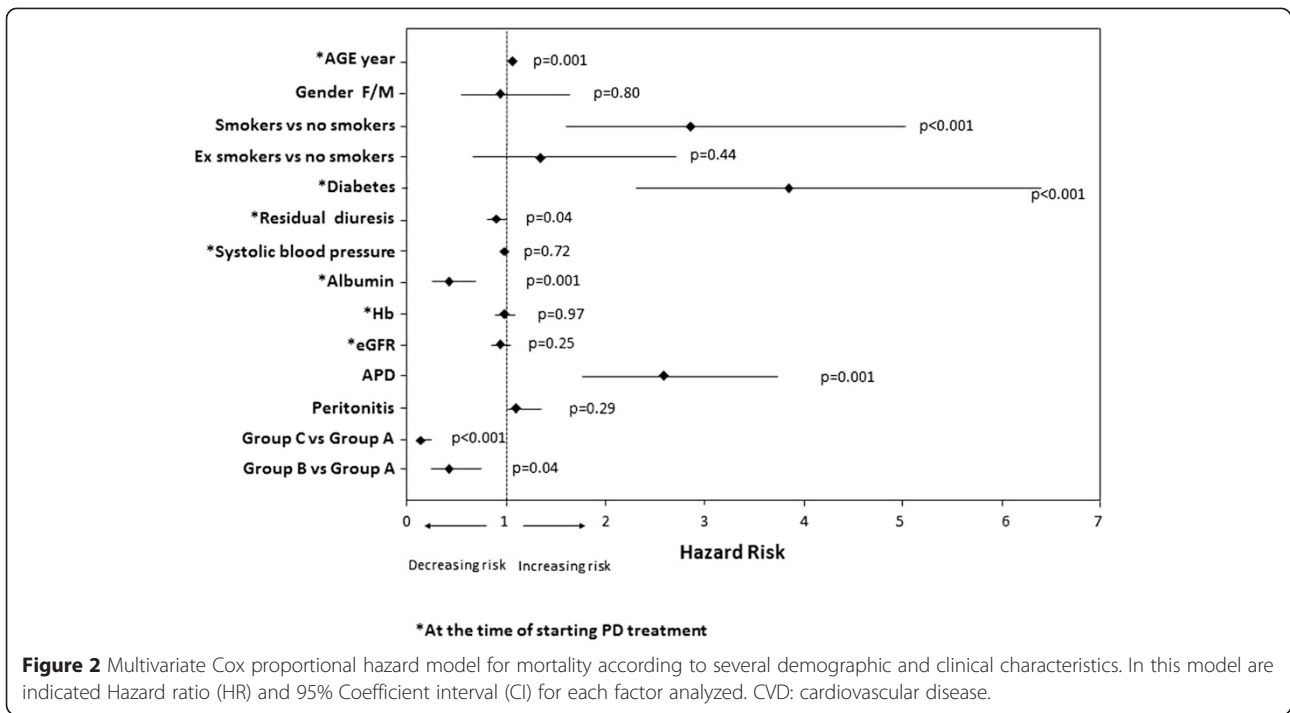
However, in a recent paper, Nessim et al. did not find any relationship between peritonitis and older age in the subgroup of patients who initiated dialysis in more recent years (2001 – 2005) [25] probably because of the recent advances in PD connection methods and exit-site care [26].

Also diabetes has been previously associated with worst outcomes in PD patients. Duong et al. have recently reported that a poor glycemic control ($A1c \geq 8\%$ or serum glucose ≥ 300 mg/dl) appears to be associated with a decreased survival in PD patients. Authors suggested also that a better glycemic control could slow down the progression of microvascular disease and loss of residual renal function [27].

Therefore, to improve the management of diabetic patients undergoing PD represents a great challenge in nephrology.

To this purpose, icodextrin use has been encouraged in this large patients' population. The benefit of this colloid osmotic agent, derived from maltodextrin, has been recently evaluated by Paniagua et al. in a prospective, randomized controlled trial in 60 diabetic patients undergoing PD [28]. These authors demonstrated that icodextrin, as compared with conventional glucose solution, reduces blood glucose concentration, a finding that was accompanied by a concomitant improvement in HbA1c and a reduction in insulin dosage. In addition, icodextrin-treated PD patients necessitated of a lower food intake.

Our study, then, reported that PD patients were highly vulnerable to the adverse consequences of smoking. This is in line with a paper published by Braatvedt et al. describing, in a large New Zealand database (more than 1000 patients), an higher age-adjusted mortality rate in PD patients with a history of current or former smoking compared to non-smokers [29].

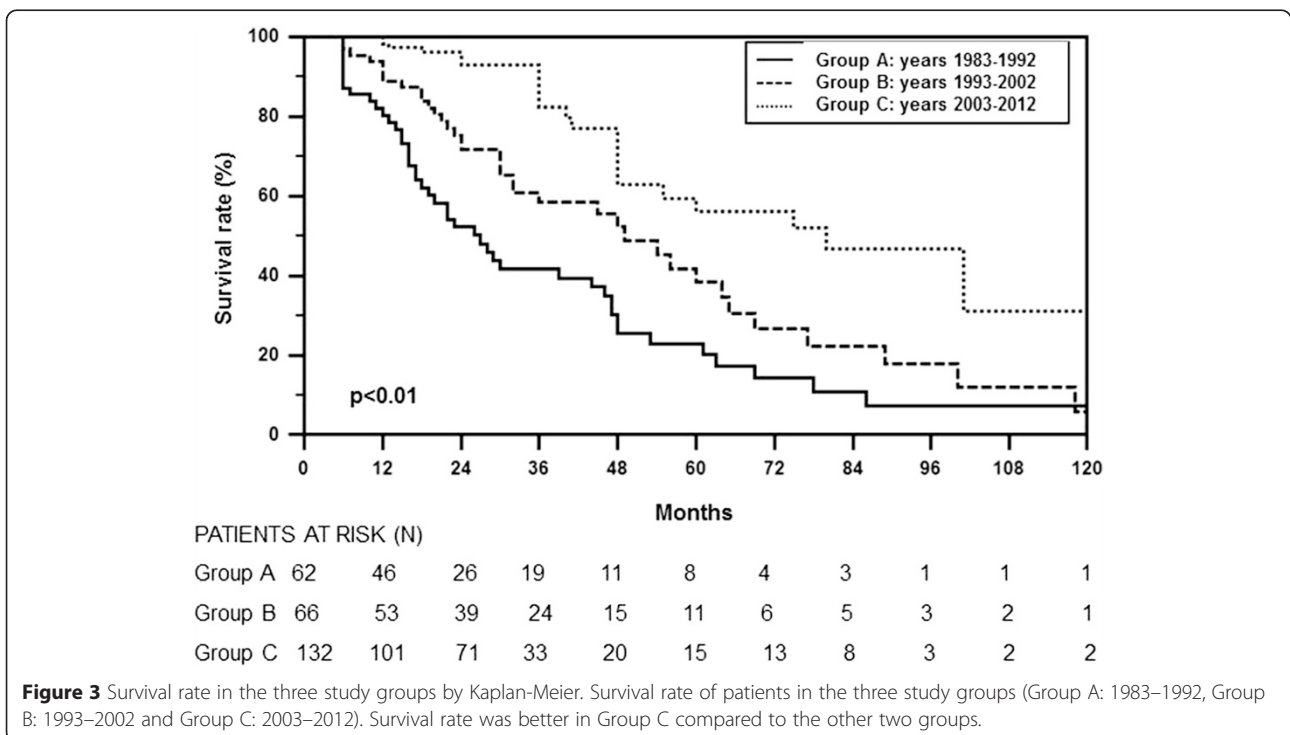


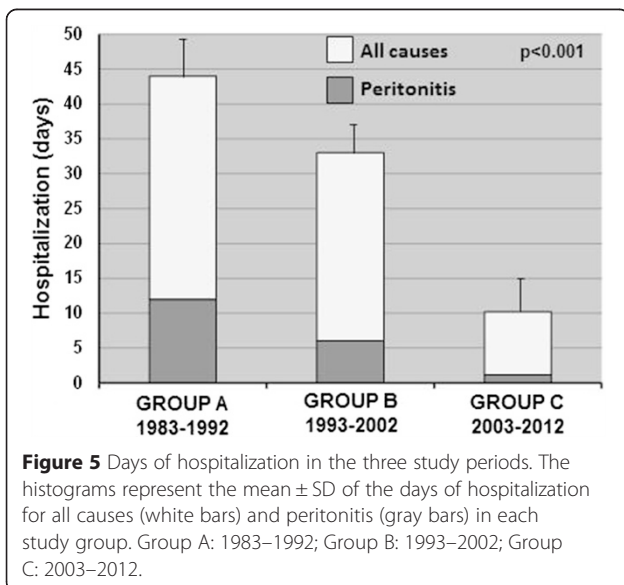
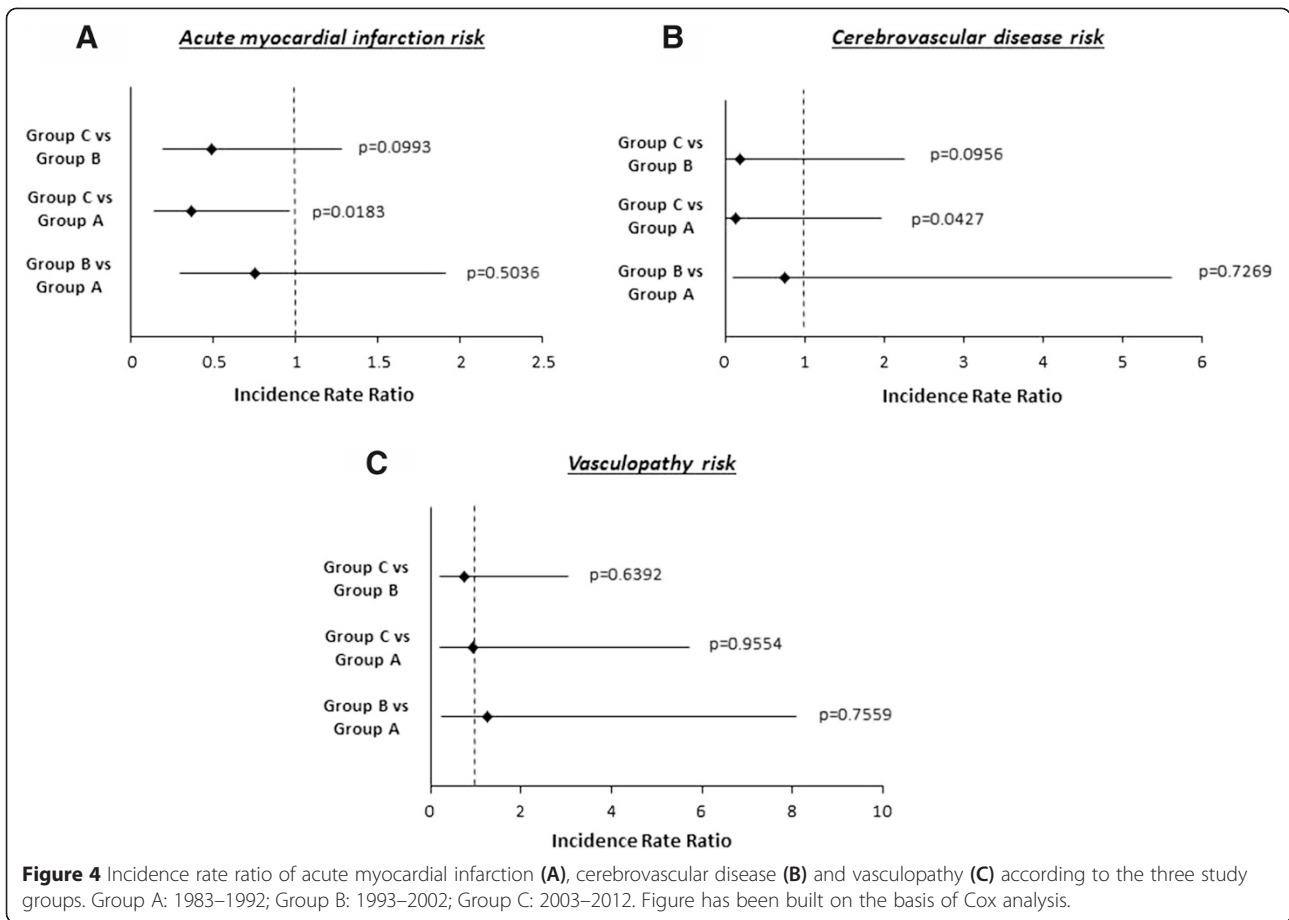
On the contrary, serum albumin levels and residual diuresis were negatively associated with the risk of mortality for all-causes in our PD patients' population.

In the past, serum albumin has been shown to predict all-cause mortality, and peritonitis risk in PD patients;

however, the data are significantly more limited than for HD patients and most of the time based on small research studies [30-33].

The biological basis for the association of hypoalbuminemia with mortality remains uncertain. A main reason





could be the magnitude of daily peritoneal protein losses (5–10 g/protein per day) in the dialysate effluent [34]. On the contrary, several researchers believe that since albumin is a negative acute phase reactant, this association in dialysis patients is secondary to the confounding influence of systemic inflammation [35]. A recent cross-sectional study of PD patients suggests that many of the patients with hypoalbuminemia are volume overloaded and hypervolemia may be an additional confounding influence [36].

Residual diuresis was the other clinical factor negatively associated with mortality in our PD patients confirming previous literature evidences reporting that a residual renal function (RRF) favoring the clearance of middle molecules, sodium removal and better control of volume status could have positive cardiovascular and systemic effects [37,38].

Moreover, as additional results of our study, we found a significant change of some of the above-mentioned mortality risk factors overtime. Mean patients' age and smoking habit were reduced, while residual diuresis raised during-time.

We suppose that all these positive modifications have been possible thanks to the implementation in our Renal

Table 2 Frequency of occurrence of peritonitis and overall/post-peritonitis technique survival in the three study periods

| | GROUP A (1983–1992) | GROUP B (1993–2002) | GROUP C (2003–2012) |
|--|---------------------|---------------------|---------------------|
| Culture negative peritonitis (CNP) | 35% | 25% | 17% |
| Culture positive peritonitis (CPP) | 65% | 75% | 83% |
| Staphylococcus aureus (% of total CPP) | 20% | 17% | 19% |
| Staphylococcus epidermidis (% of total CPP) | 40% | 40% | 36% |
| Pseudomonas (% of total CPP) | 20% | 18% | 16% |
| Other gram-negative organisms (including Klebsiella, Serratia and Enterobacter species) (% of total CPP) | 15% | 17% | 18% |
| Other organisms (% of total CPP) | 5% | 8% | 11% |
| Post-peritonitis technique survival (at 1 year after peritonitis) | 58% | 75% | 85% |
| Overall technique survival (at 5 years) | 63% | 73% | 79% |

Unit of a specific pre-dialysis out-patients follow-up strategy by a dedicated multi-disciplinary team (involving medical doctors, nurses, psychologists, nutritionists) that, personalizing medical assistance, ensures an adequate metabolic balance, a good correction of anemia (with the more rational use of ESAs) and an implementation of all possible strategies to maintain the residual diuresis (for example by limiting the use of nephrotoxic drugs and by an overfluid control). Additionally, the correct education of our PD patients has increased their therapeutic compliance/adherence, reduced/stopped smoking habit and improved dietary intake.

In the last decade, then, this multi-disciplinary pre-dialysis program, together with the reduction of our patients' age, has been responsible of the significant increment of PD patients undergoing APD modality ($p = 0.0001$). The APD, giving the possibility to continue to work and to maintain familial and social activities, can be considered a good "bridge" between CKD and renal transplantation (as showed in Figure 1A, at the moment, patients added to the waiting list for transplantation are more than 30%).

Other factors modified during the study periods were hemoglobin levels (increased) and systolic blood pressure (decreased). Both represent well known risk factors for cardiovascular mortality in the general and PD population [39,40].

This can partially justify the significant decreased risk of death for acute myocardial infarction and cerebrovascular complications in our PD patients overtime.

Conclusions

Therefore, the present study, although limited by the exclusion of other important factors that could influence clinical outcomes (including nutritional status, peritoneal transport characteristics) and absence of a control group (i.e., patients undergoing hemodialysis treatment) clearly underlines that in the last decade there has been

a significant increment in the number of patients undergoing PD and a profound change in their demographic and clinical characteristics.

Patients are younger, no smokers, with a residual diuresis, normal hemoglobin level and lower blood pressure. All these changes have definitely improved patients' survival (all-causes and cardiovascular diseases) and caused a fall of the hospitalization rate. Moreover, our pre-dialysis care, modifying most of the above reported risk factors, has been a major actor of the clinical improvement observed in our PD patients' population in the last 10 years.

Additional file

Additional file 1: Additional Statistics.

Abbreviations

CKD: Chronic kidney disease; RRT: Renal replacement therapies; HD: Hemodialysis; PD: Peritoneal dialysis; BMI: Body mass index; eGFR: Estimated glomerular filtration rate; APD: Automated peritoneal dialysis; HR: Hazard ratio; RRF: Residual renal function; ESRD: End stage renal disease.

Competing interest

The authors declare that they have no competing interests.

Authors' contributions

GZ and CR conceived the study and wrote the manuscript. AT and SG collected the clinical data and helped to write the manuscript. AP participated in the design of the study and performed the statistical analysis. AL helped in the manuscript writing and data analysis. All authors read and approved the final manuscript.

Acknowledgments

The authors thank Dr. Paola Tomei (Renal Unit, Department of Medicine, University-Hospital of Verona, Verona, Italy) for her collaboration in collecting the clinical data.

Author details

¹Renal Unit, Department of Medicine, University-Hospital of Verona, Piazzale A. Stefani 1, 37126 Verona, VR, Italy. ²Department of Public Health and Community Medicine, University of Verona, Verona, Italy.

Received: 1 October 2014 Accepted: 1 April 2015

Published online: 14 April 2015

References

- National Kidney Foundation. K/DOQI clinical practice guidelines for chronic kidney disease: evaluation, classification, and stratification. *Am J Kidney Dis.* 2002;392 Suppl 1:S1–266.
- Biesenbach G, Pohanka E. Dialysis in diabetic patients: hemodialysis and peritoneal dialysis. *Pros Cons Minerva Uro INefrol.* 2012;64(3):173–82.
- Cabiddu G, Marinangeli G. Censimento 2012 del Gruppo di Studio di Dialisi Peritoneale. <http://www.dialisiperitoneale.org/web/eventi/GDS/censimento.cfm>
- Marinangeli G, Cabiddu G, Neri L, Viglino G, Russo R, Teatini U, et al. Old and new perspectives on peritoneal dialysis in Italy emerging from the peritoneal dialysis study group census. *Perit Dial Int.* 2012;32(5):558–65.
- Van Biesen W, Verbeke F, Vanholder R. Cardiovascular disease in haemodialysis and peritoneal dialysis: arguments pro peritoneal dialysis. *Nephrol Dial Transplant.* 2007;22(1):53–8.
- Tam P. Peritoneal dialysis and preservation of residual renal function. *Perit Dial Int.* 2009;29 Suppl 2:S108–10.
- Theofilou P. Quality of life in patients undergoing hemodialysis or peritoneal dialysis treatment. *J Clin Med Res.* 2011;3(3):132–8.
- Haddy FJ, Meyer TW. Uremia. *N Engl J Med.* 2007;357:1316–25.
- Kalantar-Zadeh K, Kopple JD. Relative contribution of nutrition and inflammation to clinical outcome in dialysis patients. *Am J Kidney Dis.* 2001;38:1343–50.
- Noh H, Kim JS, Han KH, Lee GT, Song JS, Chung SH, et al. Oxidative stress during peritoneal dialysis: implications in functional and structural changes in the membrane. *Kidney Int.* 2006;69:2022–8.
- Breborrowicz A. Free radicals in peritoneal dialysis: agents of damage? *Perit Dial Int.* 1992;12:194–8.
- Wang AY. Cardiovascular risk factors in peritoneal dialysis patients revisited. *Perit Dial Int.* 2007;27 Suppl 2:S223–7.
- Dong J, Han QF, Zhu TY, Ren YP, Chen JH, Zhao HP, et al. The associations of uric acid, cardiovascular and all-cause mortality in peritoneal dialysis patients. *PLoS One.* 2014;9(1):e82342.
- Stinghen AE, Barretti P, Pecoito-Filho R. Factors contributing to the differences in peritonitis rates between centers and regions. *Perit Dial Int.* 2007;27 Suppl 2:S281–5.
- Mehrotra R, Story K, Guest S, Fedunyszyn M. Neighborhood location, rurality, geography, and outcomes of peritoneal dialysis patients in the United States. *Perit Dial Int.* 2012;32(3):322–31.
- Tonelli M, Hemmelgarn B, Culleton B, Klarenbach S, Gill JS, Wiebe N, et al. Mortality of Canadians treated by peritoneal dialysis in remote locations. *Kidney Int.* 2007;72:1023–8.
- Wang AY, Wang M, Lam CW, Chan IH, Lui SF, Sanderson JE. Heart failure in long-term peritoneal dialysis patients: a 4-year prospective analysis. *Clin J Am Soc Nephrol.* 2011;6(4):805–12.
- Lee HY, Park HC, Seo BJ, Lee HY, Park HC, Seo BJ, et al. Superior patient survival for continuous ambulatory peritoneal dialysis patients treated with a peritoneal dialysis fluid with neutral pH and low glucose degradation product concentration (balance). *Perit Dial Int.* 2005;25(3):248–55.
- Cho Y, Badve SV, Hawley CM, Wiggins K, Johnson DW. Biocompatible peritoneal dialysis fluids: clinical outcomes. *Int J Nephrol.* 2012;2012:812609.
- Khan SS, Xue JL, Kazmi WH, Gilbertson DT, Obrador GT, Pereira BJ, et al. Does predialysis nephrology care influence patient survival after initiation of dialysis? *Kidney Int.* 2005;67(3):1038–46.
- Foot C, Ninomiya T, Gallagher M, Perkovic V, Cass A, McDonald SP, et al. Survival of elderly dialysis patients is predicted by both patient and practice characteristics. *Nephrol Dial Transplant.* 2012;27(9):3581–7.
- Verdalles U, Abad S, Aragoncillo I, Villaverde M, Jofre R, Verde E, et al. Factors predicting mortality in elderly patients on dialysis. *Nephron Clin Pract.* 2010;115(1):c28–34.
- Brown EA. Peritoneal dialysis in elderly patients: clinical experience. *Perit Dial Int.* 2005;25 Suppl 3:S88–91.
- Lim WH, Dogra GK, McDonald SP, Brown FG, Johnson DW. Compared with younger peritoneal dialysis patients, elderly patients have similar peritonitis-free survival and lower risk of technique failure, but higher risk of peritonitis-related mortality. *Perit Dial Int.* 2011;31(6):663–71.
- Nessim SJ, Bargman JM, Austin PC, Story K, Jassal SV. Impact of age on peritonitis risk in peritoneal dialysis patients: an era effect. *Clin J Am Soc Nephrol.* 2009;4:135–41.
- Okayama M, Inoue T, Nodaira Y, Kimura Y, Nobe K, Seto T, et al. Aging is an important risk factor for peritoneal dialysis-associated peritonitis. *Adv Perit Dial.* 2012;28:50–4.
- Duong U, Mehrotra R, Molnar MZ, Noori N, Kovesdy CP, Nissenson AR, et al. Glycemic control and survival in peritoneal dialysis patients with diabetes mellitus. *Clin J Am Soc Nephrol.* 2011;6(5):1041–8.
- Paniagua R, Ventura MJ, Avila-Diaz M, Cisneros A, Vicenté-Martínez M, Furlong MC, et al. Icodextrin improves metabolic and fluid management in high and high-average transport diabetic patients. *Perit Dial Int.* 2009;29:422–32.
- Braatvedt GD, Rosie B, Bagg W, Collins J. Current and former smoking increases mortality in patients on peritoneal dialysis. *N Z Med J.* 2006;119(1234):U1977.
- Diaz-Buxo JA, Lowrie EG, Lew NL, Zhang SM, Zhu X, Lazarus JM. Associates of mortality among peritoneal dialysis patients with special reference to peritoneal transport rates and solute clearance. *Am J Kidney Dis.* 1999;33(3):523–34.
- Mehrotra R, Kopple JD. Protein and energy nutrition among adult patients treated with chronic peritoneal dialysis. *Adv Ren Replace Ther.* 2003;10(3):194–212.
- Wang Q, Bernardini J, Piraino B, Fried L. Albumin at the start of peritoneal dialysis predicts the development of peritonitis. *Am J Kidney Dis.* 2003;41(3):664–9.
- Mehrotra R, Duong U, Jiwakanon S, Kovesdy CP, Moran J, Kopple JD, et al. Serum albumin as a predictor of mortality in peritoneal dialysis: comparisons with hemodialysis. *Am J Kidney Dis.* 2011;58(3):418–28.
- Westra WM, Kopple JD, Krediet RT, Appell M, Mehrotra R. Dietary protein requirements and dialysate protein losses in chronic peritoneal dialysis patients. *Perit Dial Int.* 2007;27(2):192–5.
- de Mutser R, Grootendorst DC, Indemans F, Boeschoten EW, Krediet RT, Dekker FW. Association between serum albumin and mortality in dialysis patients is partly explained by inflammation, and not by malnutrition. *J Ren Nutr.* 2009;19(2):127–35.
- John B, Tan BK, Dainty S, Spanel P, Smith D, Davies SJ. Plasma volume, albumin, and fluid status in peritoneal dialysis patients. *Clin J Am Soc Nephrol.* 2010;5(8):1463–70.
- Ate S K, Nergizoğlu G, Keven K, Sen A, Kutlay S, Ertürk S, et al. Effect of fluid and sodium removal on mortality in peritoneal dialysis patients. *Kidney Int.* 2001;60:767–76.
- Davies SJ, Phillips L, Russell GI. Peritoneal solute transport predicts survival on CAPD independently of residual renal function. *Nephrol Dial Transplant.* 1998;13:962–8.
- Molnar MZ, Mehrotra R, Duong U, Kovesdy CP, Kalantar-Zadeh K. Association of hemoglobin and survival in peritoneal dialysis patients. *Clin J Am Soc Nephrol.* 2011;6(8):1973–81.
- Udayaraj UP, Steenkamp R, Caskey FJ, Rogers C, Nitsch D, Ansell D, et al. Blood pressure and mortality risk on peritoneal dialysis. *Am J Kidney Dis.* 2009;53:70–8.

Submit your next manuscript to BioMed Central and take full advantage of:

- Convenient online submission
- Thorough peer review
- No space constraints or color figure charges
- Immediate publication on acceptance
- Inclusion in PubMed, CAS, Scopus and Google Scholar
- Research which is freely available for redistribution

Submit your manuscript at
www.biomedcentral.com/submit

