Fitness for Entering a Simple Exercise Program and Mortality: A Study Corollary to the Exercise Introduction to Enhance Performance in Dialysis (Excite) Trial

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Abstract

Background/Aims: In this corollary analysis of the EXCITE study, we looked at possible differences in baseline risk factors and mortality between subjects excluded from the trial because non-eligible (n=216) or because eligible but refusing to participate (n=116).

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Introduction

Precise definition of the population investigated in clinical trials and in well-planned observational studies is fundamental to understand the potential applicability of findings in these studies in clinical practice. The CONSORT (Consolidated Standards of Reporting Trials) document recommends a specific diagram for describing the flow of participants through the main phases of clinical trials [1], from eligibility to actual enrollment and follow-up. By the same token, a similar approach has been recommended for observational studies by STROBE, another document produced under the aegis of the EQUATOR (Enhancing the QUAlity and Transparency Of health Research) initiative. The importance of accurate reporting of the selection process of subjects enrolled into a trial is of obvious relevance for the generalizability of findings in the same trial. However, outcome data in subjects screened but not enrolled in clinical trials (as not eligible or as refusing to provide consent to participate) have received very little attention and we have been unable to identify even a single study focusing on this issue. The problem appears of particular relevance in trials testing exercise programs, i.e. trials which selects individuals with an inherently lower risk profile, i.e. with a degree of fitness allowing a physical exercise program. We hereby report a study corollary to the EXerCise Introduction To Enhance performance in dialysis (EXCITE) study, i.e. a multicentric, randomized controlled clinical trial on the effectiveness of exercise for improving physical performance and the quality of life and for reducing adverse clinical outcomes (mortality, cardiovascular events and hospitalizations) in dialysis patients (NCT01255969). We specifically designed this study to investigate differences in baseline risk factors and their influence on the overall survival in non-eligible patients and in eligible patients who declined the invitation to participate into the trial.

Patients and Methods

The study protocol was in conformity to the ethical guidelines of our institution and informed consent was obtained from each participant.

Patients

This is a study corollary to the EXerCise Introduction To Enhance performance in dialysis (EXCITE) study. More detailed information about this trial are available in ClinicalTrials.gov (NCT01255969). In this study we investigated the differences in baseline risk factors and their influence on the incidence rate of mortality in non-eligible patients (n=216; 90% on HD) and eligible patients who did not give the informed consent.
consent to take part into the EXCITE trial (n=116; 84% on HD). Deambulation ability was categorized as independent, assisted or total inability to deambulate (bedridden or wheel-chaired). The main demographic, clinical and biochemical characteristics of patients included in this study are given in Table 1.

**Laboratory measurements**

Blood sampling was performed during a mid-week day (short dialysis interval). Serum cholesterol, albumin, calcium, phosphate, C-Reactive Protein (CRP) and haemoglobin measurements were made using standard methods in the routine clinical laboratory.

**Follow up**

After the initial assessment, the median follow-up was 2.9 years in non-eligible patients (interquartile range 1.2 to 4.0 years) and 4.0 years in eligible patients unwilling to participate into the trial (interquartile range 2.3 to 4.0 years). During follow-up, death was accurately recorded. Each death was reviewed and assigned an underlying cause by a panel of 5 physicians. As a part of the review process, all available medical information about each death were collected. This information always included study and hospitalisation records. In the case of an out-of-hospital death family members were interviewed by telephone to better ascertain the circumstances surrounding death.

**Statistical analysis**

Data were expressed as mean ± standard deviation (normally distributed data), median and inter-quartile range (non-normally distributed data) or as per cent frequency (categorical data), and the comparisons among groups were made by One Way ANOVA, Kruskal-Wallis Test and Chi Square Test, as appropriate.

In each study cohort, the independent predictors of mortality were identified by Kaplan-Meier survival analysis and by univariate and multivariate Cox regression models. Tested variables included age, gender, smoking, diabetes, blood pressure, total cholesterol, triglycerides, haemoglobin, albumin, calcium, phosphate, C-Reactive Protein, history of neoplasm, myocardial infarction, stroke, TIA, angina episodes, arrhythmia, peripheral vascular disease, and heart failure. We built-up multiple Cox regression models specific to each study cohort, by introducing into these models all variables which were related to all-cause mortality with \( P < 0.10 \). To assess whether the observed difference in the death risk among the two study cohorts (i.e. non-eligible patients and eligible non-randomized patients) could be explained by differences in baseline risk factors (Table 1), we performed a multiple Cox regression analysis in the whole study population (n=648) including the group (cohort) variable and a series of risk factors which were related

### Table 1. Main demographic, clinical and biochemical characteristics of patients

|                          | Non eligible N=216 | Eligible non randomized N=116 | P     |
|--------------------------|--------------------|-------------------------------|-------|
| Age (years)              | 70±14              | 67±14                         | 0.043 |
| Male sex n. (%)          | 119 (56)           | 75 (65)                       | 0.111 |
| Smokers n. (%)           | 61 (29)            | 33 (49)                       | 0.049 |
| Diabetics n. (%)         | 64 (30)            | 24 (21)                       | 0.063 |
| History of neoplasia n. (%) | 44 (21)         | 29 (25)                       | 0.355 |
| Myocardial Infarction n. (%) | 51 (24)           | 18 (16)                       | 0.077 |
| Stroke n. (%)            | 29 (14)            | 11 (10)                       | 0.286 |
| Transient Ischemic Attack n. (%) | 44 (21)          | 11 (10)                       | 0.010 |
| Anginal episodes n. (%)  | 35 (16)            | 8 (7)                         | 0.015 |
| Arrhythmia n. (%)        | 53 (25)            | 13 (11)                       | 0.003 |
| Peripheral vascular disease n. (%) | 45 (21)       | 4 (4)                          | <0.001 |
| Heart failure n. (%)     | 145 (73)           | 96 (61)                       | 0.02  |
| NYHA class n. (%):       |                    |                               |       |
| 1                        | 54 (26)            | 50 (45)                       | <0.001 |
| 2                        | 29 (14)            | 7 (6)                         |       |
| 3-4                      | 70 (34)            | 10 (9)                        |       |
| Deambulation:            |                    |                               |       |
| Independent              | 79 (37)            | 107 (94)                      | <0.001 |
| Assisted                 | 77 (36)            | 7 (6)                         |       |
| Total inability (bedridden/ wheelchair) | 60 (28)         | 0 (0)                         |       |
| Systolic BP (mmHg)       | 127±19             | 133±19                        | 0.077 |
| Diastolic BP (mmHg)      | 69±11              | 71±10                         | 0.239 |
| Hypertension n. (%)      | 144 (67)           | 91 (78)                       | 0.028 |
| Total cholesterol (mg/dL) | 164±41            | 171±50                        | 0.198 |
| Triglycerides (mg/dL)    | 153±73             | 175±106                       | 0.05  |
| Haemoglobin (g/dL)       | 11±2               | 11±1                          | 0.571 |
| Albumin (g/dL)           | 4±3±8.4            | 3±9±0.4                       | 0.351 |
| Calcium (mg/dL)          | 7.9±2.0            | 8.3±2.0                       | 0.156 |
| Phosphate (mg/dL)        | 4.4±1.4            | 5.0±1.8                       | 0.002 |
| CRP (mg/L)               | 1.3 (0.4-3.6)      | 0.7 (0.4-2.9)                 | 0.020 |

Data are expressed as mean ± SD, median and inter-quartile range or as percent frequency, as appropriate.
(with $P \leq 0.10$) to mortality in each study cohort (Table 2) and/or differed (with $P \leq 0.10$) among the two cohorts (Table 1). By this strategy we constructed models of adequate statistical power (i.e. at least 10 deaths for each variable into the models). In the survival analysis, data were expressed as hazard ratio, 95% confidence interval and $P$ value.

The explained variation in mortality attributable to single variables was calculated by the method proposed by Hosmer and Lemeshow [2]. Statistical analysis was performed by using a standard statistical package (SPSS for Windows, Version 20, Chicago, Illinois, USA).

**Results**

The source population of the EXCITE study included 648 patients with ESKD. Among these, 216 patients were excluded because they did not meet inclusion criteria (33%), 116 patients were eligible but did not give the informed consent (18%) and the remaining 316 patients were randomized (49%). Most non-eligible patients (64%) did not deambulate autonomously and/or needed to be assisted in everyday life. Eligible patients non-participating into the study were significantly younger and displayed a lower proportion of patients with angina, arrhythmia, heart failure and TIA as compared to non-eligible patients. Circulating levels of phosphate and the prevalence of smokers and hypertensive patients were higher in eligible non-randomized patients than in non-eligible patients (Table 1). Circulating levels of C-Reactive Protein (CRP) and the prevalence of peripheral vascular disease were higher in non-eligible patients than in eligible patients who did not give the informed consent (Table 1).

**Predictors of mortality in eligible non-randomized patients and in ineligible patients**

During the follow-up period, the number of deaths was 119 in non-eligible patients (55%) and 41 in eligible non-randomized (35%) patients. In non-eligible patients, age,

### Table 2. Univariate Cox regression analyses in the two study cohorts. Categories grading ambulation impairment are listed in Table 1.

| Variable                                    | Unit of increase | Non-eligible patients* | Eligible non-randomized patients** |
|---------------------------------------------|------------------|------------------------|------------------------------------|
| Age                                         | 1 year           | 1.04 (1.02-1.06), $p < 0.001$ | 1.04 (1.01-1.06), $p = 0.01$       |
| Gender                                      | 0=F; 1=M         | 1.17 (0.81-1.68), $p = 0.41$ | 0.90 (0.48-1.70), $p = 0.74$       |
| Smoking                                     | 0=no; 1=yes      | 1.41 (0.95-2.11), $p = 0.09$ | 1.32 (0.60-2.94), $p = 0.49$       |
| Diabetes                                    | 0=no; 1=yes      | 1.13 (0.77-1.66), $p = 0.54$ | 1.04 (0.50-2.19), $p = 0.91$       |
| Systolic BP                                  | 1 mmHg           | 0.99 (0.99-1.01), $p = 0.47$ | 1.01 (0.98-1.03), $p = 0.56$       |
| Diastolic BP                                 | 1 mmHg           | 0.99 (0.97-1.00), $p = 0.11$ | 0.97 (0.93-1.01), $p = 0.12$       |
| Total cholesterol                           | 1 mg/dl          | 1.00 (0.99-1.00), $p = 0.97$ | 1.00 (1.00-1.01), $p = 0.35$       |
| Triglycerides                                | 1 mg/dl          | 0.99 (0.99-1.00), $p = 0.01$ | 1.00 (0.99-1.00), $p = 0.74$       |
| Haemoglobin                                  | 1 g/dl           | 0.98 (0.90-1.06), $p = 0.58$ | 1.02 (0.82-1.27), $p = 0.87$       |
| Albumin                                     | 1 g/dl           | 0.99 (0.94-1.04), $p = 0.71$ | 0.93 (0.41-2.13), $p = 0.86$       |
| Calcium                                     | 1 mg/dl          | 0.99 (0.90-1.08), $p = 0.79$ | 1.02 (0.87-1.20), $p = 0.84$       |
| Phosphate                                    | 1 mg/dl          | 0.88 (0.77-1.00), $p = 0.05$ | 1.00 (0.84-1.20), $p = 0.99$       |
| CRP                                         | 1 mg/L           | 1.01 (1.00-1.01), $p = 0.03$ | 0.99 (0.94-1.04), $p = 0.57$       |
| History of myocardial infarction             | 0=no; 1=yes      | 1.25 (0.82-1.88), $p = 0.29$ | 0.50 (0.18-4.11), $p = 0.19$       |
| History of stroke                           | 0=no; 1=yes      | 1.15 (0.68-1.95), $p = 0.60$ | 1.34 (0.52-3.41), $p = 0.54$       |
| History of TIA                               | 0=no; 1=yes      | 1.23 (0.79-1.89), $p = 0.36$ | 1.28 (0.50-3.26), $p = 0.61$       |
| History of angina                            | 0=no; 1=yes      | 1.31 (0.82-2.07), $p = 0.26$ | 1.00 (0.31-3.23), $p = 0.99$       |
| History of arrhythmia                        | 0=no; 1=yes      | 1.39 (0.93-2.06), $p = 0.11$ | 2.24 (1.03-4.85), $p = 0.04$       |
| History of heart failure                     | 0=no; 1=yes      | 1.46 (1.25-1.72), $p < 0.001$ | 1.05 (0.74-1.49), $p = 0.78$       |
| History of peripheral vascular disease       | 0=no; 1=yes      | 1.53 (1.00-2.32), $p = 0.05$ | 1.51 (0.37-6.13), $p = 0.57$       |
| History of neoplasia                         | 0=no; 1=yes      | 1.37 (0.89-2.11), $p = 0.15$ | 1.98 (1.04-3.77), $p = 0.04$       |

* HR (95% CI) and P; ** HR (95% CI) and P
The incidence rate of mortality was substantially higher in non-eligible patients (incidence rate: 21.0 deaths/100 persons-year) than in eligible non-randomized patients (incidence rate: 10.9 deaths/100 persons-year) (Log rank test: $\chi^2=13.85$, $p<0.001$) (Fig. 1a). To assess whether the difference in baseline risk factors could explain the difference in the death risk of the two study cohorts, we performed two multivariate analyses including the group variable of eligible and non-eligible patients (cohorts) and a series of risk factors which were related to mortality (with $p<0.10$) in each study cohort (Table 2) and/or differed (with $p<0.10$) among the two cohorts (Table 1). To estimate the impact of the degree of deambulation impairment on mortality we tested two models, the first excluding deambulation ability and the second including this co-variate. In the first model the crude difference in the mortality risk among the two cohorts (HR 1.96; 95% CI 1.36 to 2.77; $P<0.001$) was reduced modestly (HR 1.60; 95% CI 1.10 to 2.35; $P=0.017$) (Fig. 1b and Table 4). However, additional adjustment for deambulation almost nullified the excess risk of death seen in non-eligible patients (HR 1.16; 95% CI 0.75 to 1.80; $P=0.513$) (Fig. 1c and Table 4). Of note, the explained variation in all-cause mortality was 33% for the model excluding and 38% for the model including the deambulation variable ($P<0.0005$). Thus, deambulation captures as much as the 13% in the explained variability in the risk of death.

**Discussion**

This study corollary to a multicenter randomized, clinical trial testing the effect of physical exercise in dialysis patients shows that the mortality risk is twice higher in patients who were excluded from the trial than in eligible patients who did not accept to participate. Importantly, the mortality rate in this group (10.9 deaths/100 persons-year) was lower than the average mortality rate in the ERA-EDTA registry [3] and in the Italian registry of dialysis [4]. As expected, the two populations which remained external to the trial, i.e. the

| Units of increase | HR (95% CI) and P |
|-------------------|------------------|
| **a) Number of deaths, n=119** |
| Age | 1 year | 1.04 (1.02-1.06), $p<0.001$ |
| Smoking | 0=no; 1=yes | 1.13 (0.74-1.72), $p=0.59$ |
| Triglycerides | 1 mg/dL | 1.00 (0.99-1.00), $p=0.04$ |
| Phosphate | 1 mg/dL | 0.95 (0.82-1.10), $p=0.95$ |
| CRP | 1 mg/L | 1.01 (1.00-1.02), $p=0.05$ |
| History of peripheral vascular disease | 0=no; 1=yes | 1.29 (0.84-1.99), $p=0.25$ |
| Deambulation | 1 unit | 0.72 (0.57-0.91), $p=0.01$ |
| History of heart failure | 0=no; 1=yes | 1.81 (1.03-3.16), $p=0.04$ |

| **b) Number of deaths, n=41** |
| Age | 1 year | 1.03 (1.00-1.06), $p=0.04$ |
| History of arrhythmia | 0=no; 1=yes | 1.91 (0.87-4.17), $p=0.11$ |
| History of neoplasia | 0=no; 1=yes | 1.62 (0.84-3.15), $p=0.15$ |
ineligible population and the population with sufficient fitness but unwilling to undergo the exercise program, showed several differences in baseline risk factors such as age, blood pressure, phosphate, CRP, smoking, diabetes, triglycerides, cardiovascular comorbidities and history of neoplasia. We hypothesized that these differences could explain the higher death risk of ineligible patients and tested this hypothesis by modeling death risk in an analysis where the two cohorts were nominally identified by a specific covariate. The proviso was that the inclusion of known risk factors differentiating the two cohorts should attenuate substantially or cancel out the excess death risk of unfit patients ineligible to the trial. However, in contrast to our working hypothesis, we found that adjustment for background risk factors only in limited part explained the between cohorts difference in mortality. This finding indicates that major risk factors fail to fully capture the higher probability of death in patients ineligible to clinical studies. Thus, the higher death rate in those patients must necessarily depend on unmeasured risk factors of paramount importance for human health. Most non-eligible patients (64%) did not deambulate autonomously or needed to be assisted, indicating that comorbidities limiting motor activity in everyday life mark a degree of severity that is unaccounted for by nominally defined, ungraded, major comorbidities like arrhythmia, heart failure, cerebro- and peripheral-vascular disease, ischemic heart disease and neoplasia. Indeed, when we introduced into the model deambulation impairment, the between cohorts difference in the risk of death almost disappeared (HR 1.16; 95% CI 0.75

Fig. 1. Unadjusted and adjusted survival analyses in the two study groups (non-eligible and eligible non randomized patients).
Table 4. Multivariate Cox regression analysis in the whole study population

| Eligible non-randomized patients | Units of increase | Crude analysis** | Excluding Deambulation** | Including Deambulation** |
|----------------------------------|-------------------|-----------------|--------------------------|-------------------------|
| Non eligible patients            | 1                 | 1*              | 1*                       | 1*                      |
| Age                              | 2                 | 1.94 (1.36-2.77), p <0.001 | 1.60 (1.10-2.35), p=0.017 | 1.16 (0.75-1.80), p=0.513 |
| Systolic BP                      | 1 year            | 1.03 (1.02-1.05), p=0.001 | 1.03 (1.02-1.05), p=0.001 | 0.93 (0.88-1.01), p=0.27 |
| Triglycerides                    | 1 mmHg            | 1.00 (1.00-1.01), p=0.86 | 1.00 (1.00-1.01), p=0.77 | 0.99 (0.89-1.11), p=0.85 |
| Phosphate                        | 1 mg/dL           | 1.00 (1.00-1.00), p=0.13 | 1.00 (1.00-1.00), p=0.07  | 0.99 (0.89-1.11), p=0.85 |
| CRP                              | 1 mg/L            | 1.01 (1.00-1.02), p=0.08 | 1.01 (1.00-1.01), p=0.14 | 1.01 (1.00-1.01), p=0.14 |
| Smoking                          | 0=no; 1=yes       | 1.26 (0.87-1.83), p=0.21 | 1.27 (0.88-1.85), p=0.21 | 1.27 (0.88-1.85), p=0.21 |
| Diabetes                         | 0=no; 1=yes       | 1.10 (0.75-1.60), p=0.64 | 0.92 (0.92-1.37), p=0.69 | 0.92 (0.92-1.37), p=0.69 |
| History of myocardial infarction | 0=no; 1=yes       | 0.87 (0.57-1.34), p=0.53 | 0.87 (0.57-1.33), p=0.52 | 0.87 (0.57-1.33), p=0.52 |
| History of TIA                   | 0=no; 1=yes       | 1.07 (0.71-1.62), p=0.75 | 0.96 (0.63-1.47), p=0.86 | 0.96 (0.63-1.47), p=0.86 |
| History of angina                | 0=no; 1=yes       | 0.97 (0.59-1.58), p=0.90 | 1.03 (0.63-1.67), p=0.92 | 1.03 (0.63-1.67), p=0.92 |
| History of arrhythmia            | 0=no; 1=yes       | 1.17 (0.81-1.70), p=0.40 | 1.20 (0.83-1.74), p=0.34 | 1.20 (0.83-1.74), p=0.34 |
| History of peripheral vascular disease | 0=no; 1=yes     | 1.41 (0.92-2.16), p=0.12 | 1.38 (0.90-2.13), p=0.14 | 1.38 (0.90-2.13), p=0.14 |
| History of heart failure         | 0=no; 1=yes       | 1.46 (0.96-2.21), p=0.08 | 1.34 (0.88-2.04), p=0.18 | 1.34 (0.88-2.04), p=0.18 |
| Hypertension                     | 0=no; 1=yes       | 0.99 (0.67-1.46), p=0.97 | 1.04 (0.71-1.54), p=0.84 | 1.04 (0.71-1.54), p=0.84 |
| History of neoplasm              | 0=no; 1=yes       | 1.26 (0.86-1.84), p=0.24 | 1.29 (0.88-1.90), p=0.19 | 1.29 (0.88-1.90), p=0.19 |
| Deambulation                     | 1 category       | 0.66 (0.52-0.84), p=0.001 |                        |                         |

*Reference group; **HR (95% CI) and P

to 1.80; P=0.513). This observation is of importance because physical performance and inability to deambulate are very rarely taken into account when describing the risk profile of dialysis patients in major clinical trials and observational studies in this population. For example, neither in the 4D [5], nor in the AURORA study [6], nor in the more recent EVOLVE trial [7], physical disability was an exclusion criterion but in none of these trials information on physical disability was given. About 1/3 (33%) of the source cohort of the EXCITE study had a relevant degree of physical impairment and it is likely that a similar proportion existed in the source populations of other trials in dialysis patients. The variability in mortality rate explained by unfitness to a simple exercise program is substantial and ignoring severe limitation in physical functioning may have a non trivial effect for the interpretation of the risk profile of patients included in clinical studies. However, as discussed, information on ambulatory ability is almost always omitted in trials in the dialysis population. Findings in this study represent a call for systematic reporting information on physical activity and/or (motor) ability in clinical trials and observational studies in this population. Indeed -well beyond classical risk factors, background co-morbidities and risk factors peculiar to end-stage renal disease- deambulation impairment explains a relevant proportion of the variability in mortality in ESKD. Furthermore, our findings suggest that extending outcome reporting and data analyses to the whole source population of clinical trials may provide relevant information to better frame the implications of the selection process applied to restrict the trial population to patients with well-defined demographic and clinical characteristics.
Disclosure Statement

The authors of this manuscript state that they do not have any conflict of interests and nothing to disclose.

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