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Research Article

The interrelationship between muscle endurance, self-perceived fatigue and pre-frailty in community-dwelling octogenarians.

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HIGHLIGHTS:

- Non-exhausted pre-frail elderly report higher fatigue levels than non-frail ones
- Combined muscle fatigability and self-perceived fatigue predicts pre-frailty
- Capacity-to-Perceived Vitality ratio (CPV) captures early signs of fatigue

ABSTRACT

Introduction: Low muscle endurance and high feelings of self-perceived fatigue could be an early characteristic of decline in intrinsic capacity, which comes to full expression as physical frailty in a later stage. This study aimed to investigate if the combination of muscle endurance and self-perceived fatigue is related to pre-frailty in well-functioning older adults aged 80 and over.

Methods: Four-hundred and five community-dwelling older adults aged 80 and over (214 robust and 191 pre-frail) were assessed for muscle endurance (Grip Work corrected for body weight (GW_bw)), self-perceived fatigue (MFI-20) and frailty state (Fried Frailty Index, FFI). A Capacity to Perceived Vitality (CPV) ratio was calculated by dividing GW_bw by the MFI-20 scores. ANCOVA analysis (corrected for age and gender) was used to compare robust and pre-frail older adults, and binary logistic regressions were applied to analyze the relationship between CPV and pre-frailty status.

Results: Pre-frail older adults who scored negative on the exhaustion item of the FFI still showed significantly lower GW_bw (p<0.001), CPV ratios (p<0.001) and higher self-perceived fatigue (p<0.05) compared to the robust ones. The risk for pre-frailty related significantly to higher age, being male and lower CPV ratios. In females, every unit increase in CPV ratio decreased the risk for pre-frailty by 78% (OR 0.22; 95% CI: 0.11-0.44), for males this effect was less strong (34%, OR 0.66; 95% CI: 0.47-0.93).

Conclusions: Pre-frail community-dwelling persons aged 80 years and over without clinical signs of exhaustion on the FFI still experience significantly higher fatigue levels (lower muscle endurance, higher self-perceived fatigue and lower CPV levels) compared to robust ones. CPV ratio could therefore be a good tool to identify subclinical fatigue in the context of physical (pre-)frailty.

Keywords: Fatigue; Tiredness; Muscle Endurance; Pre-frailty; Community-dwelling; Elderly; Vitality

1. INTRODUCTION

Frailty is a sign of losses in reserve capacity and can be conceptually defined as "a clinical state in which there is an increase in an individual's vulnerability to develop negative health-related events (including disability, hospitalizations, institutionalizations, and death) when exposed to endogenous or exogenous stressors" (Vella-azzopardi and others 2016). As a consequence, the frail person is at increased risk for different negative health outcomes such as disability, hospitalization and death (Vermeiren and others 2016). A widely accepted approach for the assessment of frailty consists of five components as described by Fried and others (2001): exhaustion, unintentional weight loss, low physical activity, slow walking and low grip strength. Fatigue, muscle atrophy (defined as sarcopenia) and muscle weakness (defined as dynapenia) are key elements in this phenotype. In the pathophysiology of frailty, sarcopenia, fatigue and inflammation have been recognized to play a predominant role (Bauer and Sieber 2008). Altogether, fatigue characterizes the depletion of physiological reserve capacity leading to a higher risk for negative health outcomes (Knoop and others 2021). However, it is unclear when fatigue plays a role in the development of frailty. Research on the occurrence of fatigue in the early stages of frailty is crucial, as frailty is believed to be reversible at this stage.

Generally speaking, fatigue can be divided into self-perceived feeling of fatigue and resistance to physical tiredness which includes a fatigue assessment such as muscle fatigability. A recent systematic review showed that fatigue has a prominent role in the operationalization of frailty and is included in most frailty scales (Knoop and others 2019). The different frailty instruments described in the literature cover a great diversity in fatigue constructs reflecting different underlying pathophysiological mechanisms by which fatigue relates to frailty. The fact that fatigue is a prominent element in the concept of frailty might be explained by the involvement of a chronic low-grade inflammatory profile (CLIP), considered as an important parameter in the pathogenesis of both fatigue and frailty (Hubbard and Woodhouse 2010; Krabbe and others 2004). Higher levels of circulating inflammatory markers such as C-reactive protein (CRP) are seen in older persons who are identified with physical frailty and sarcopenia (Marzetti and others 2019). Inflammation promotes sickness behaviour with fatigue as one of the symptoms (Dantzer and Kelley 2007) and is related to physical limitations and frailty (Cao Dinh and others 2019; Walston 2002).

Muscle endurance and self-perceived fatigue provide complementary information in well-functioning older people (Bautmans and others 2010; Bautmans and others 2008). Recent research has shown that fatigue is one of the early characteristics of frailty as signs of fatigue are already present approximately nine years prior to the occurrence of frailty (Stenholm and others 2019). However, it is unclear whether the combination of low muscle endurance and high self-perceived fatigue reflects an increased risk for the loss of reserve capacity, which comes to full expression as physical frailty. It is conceivable that good muscle endurance and low self-perceived fatigue outweigh the deficits in reserve capacity, thus conserving the robust and independent state. It is clinically relevant to identify early predictors for physical (pre-)frailty to start preventive interventions. Once an advanced frailty state is attained, it is very difficult to reverse to a robust state. Therefore, this study aimed to investigate

if the combination of muscle endurance and self-perceived fatigue is related to pre-frailty in well-functioning older adults aged 80 and over.

2. METHODOLOGY

2.1 Study design:

Baseline data were used from the BUTTERFLY study (BrUssels sTudy on The Early pRedictors of Frailty) (Cao Dinh and others 2019; Vermeiren and others 2019), involving a cohort of community-dwelling older adults aged 80 and over, organized by the Vrije Universiteit Brussel (Belgium). This study was approved by the ethical committee of UZ Brussel (B.U.N. 143201421976), all participants signed informed consent.

2.3. Setting and participants

Four-hundred-ninety-four community-dwelling adults aged 80 years and over were recruited to participate in the BUTTERFLY study. The participants were recruited through advertisements via websites of the University Hospital in Jette in Belgium, general practitioners, pharmacies and health insurance companies. Participants were eligible for the study if they were aged 80 and over, could walk independently, lived independently, if they were mentally fit (MMSE>23/30), and not frail according to the Groningen Frailty Indicator (GFI<4/15) (Steverink and others 2001), Rockwood Frailty Index (RFI<0.25/10) (Collerton and others 2012), and/or the adapted Fried Frailty Index (FFI<3/4) (Sirola and others 2011); (exhaustion, weight loss, gait speed, and grip strength). Participants were excluded if they underwent surgery or any radiotherapy or chemotherapy during the past six months. Participants with CRP >10 mg/L were excluded, as this refers to an acute inflammatory state and not to a chronic low-grade inflammatory profile (Sproston and Ashworth 2018).

2.4. Measurements

2.4.1 Frailty score

As reported previously (Cao Dinh and others 2019), the adapted version of the physical phenotype of frailty as proposed by Sirola and others (2011) was used to determine frailty by a combination of 4 components: unintentional weight loss, exhaustion, weakness and low gait speed. Weight loss was evaluated by the self-reported question: "In the last six months, have you lost more than 4.5 kg unintentionally?" which was answered by yes (1) or no (0) (Fried and others 2001). Exhaustion was measured similarly to the original Fried phenotype, questioning two statements from the CES-D Depression Scale (Orme and others 1986): "I felt that everything I did was an effort" and "I could not get going". The participants were asked: "How often in the last week did you feel this way?" and were scored (0) for rarely or none of the time, (1) for some or a little of the time, (2) for a moderate amount of time, or (3) for most of the time. When participants scored a 2 or 3 on either of the two statements, they received a point on the frailty scale for exhaustion. Gait speed was measured by timing the walked distance of 4.5 m and was stratified for gender and height, as proposed by Fried and others (2001). Participants were scored a point for slow walking if their walking time was ≥7 seconds in men ≤173 cm and women ≤159 cm, and if their time was ≥6 seconds in men >173 cm and women >159 cm. Grip strength was assessed using the Martin Vigorimeter; cut-offs were 42 kPa for women and 71 kPa for men (Cao Dinh and others

2019). Participants showing a grip strength below these cut-offs received a point for this item. In analogy to the original version (Fried and others 2001), the following scoring system was put forward to assign the level of frailty: a score of 0/4 signifies robustness, 1-2/4 points means pre-frailty, and with a score of 3 or 4/4 one is considered frail.

Maximal grip strength (Gsmax), fatigue resistance (FR), Grip Work (GW)

Participants performed a maximal handgrip strength (GSmax) and a Fatigue resistance (FR) performance test using a Martin Vigorimeter (KLS Martin Group, Tuttlingen Germany). The Martin Vigorimeter is provided with 3 different sizes of compressible rubber bulbs. For this study the large bulb was used as described previously (Bautmans and others 2007; Bautmans and Mets 2005). Briefly, the participants were asked to squeeze the bulb (with the dominant hand) three times as hard as possible, the highest score of the three attempts was registered as GSmax. Afterwards, the FR was measured by asking the participants to squeeze in the bulb as hard as possible and to maintain this maximal effort as long as possible, under standardized verbal stimulation by the investigator. The time in seconds during which the GSmax dropped to 50% of its maximum was recorded as FR. These two parameters were used to calculate Grip Work (GW) by multiplying FR in seconds by 75% of the GSmax (GW= 0.75 * GSmax * FR) (Bautmans and others 2007). GW was also corrected for body mass (GW/body mass in kg) since overweight and obese participants will have to engage more strength and sustain that higher effort over time to execute their daily tasks (Bautmans and others 2011).

2.4.2 Self-perceived fatigue

The Multidimensional Fatigue Inventory (MFI-20) was used to assess the level of self-perceived fatigue. The MFI-20 is a self-reported fatigue questionnaire consisting of twenty items that cover five domains of fatigue: (1) general fatigue, (2) physical fatigue, (3) reduction in activity, (4) reduction in motivation, (5) cognitive fatigue (Smets and others 1995). If more than half of the items were missing, items were substituted by the mean of the non-missing items. Items are scored on a five-point Likert scale, a higher score (20-100) indicates higher fatigue and vice versa. In addition, the Mobility Tiredness scale- designed to measure mobility-related fatigue in older adults — was used. It is questionnaire-based and counts the number of items where the participant reported tiredness after performing six activities; indoor transfers, walk indoors, go outdoors, walk outdoors in nice weather and walk outdoors in poor weather, and climb stairs. Low score refers to low fatigue and vice versa (Avlund and others 1993).

2.4.3 Inflammation

Non-fasting blood samples were taken by venepunction on the day of assessment, before performing the physical tests. Circulating level of C-reactive protein (CRP) was obtained by nephelometry (Behring, Marburg, Germany). When measurements of CRP fell below the LOD (Lower limits of detection <0.5) (n=53), we imputed values from a uniform distribution between 0 and the LOD.

2.4.4 Anthropometry, body composition & Physical activity

Height and weight were measured (to the nearest exhaustio0.01m and 0.1kg respectively). Body fat was estimated using a fan beam whole-body DXA device (Hologic 4500 QDR upgraded to Discovery [Bedford, Massachusetts, USA]). Use of DXA to measure body composition is widely accepted and used in clinical practice (Smith-Ryan and others 2017; Vermeiren and others 2019). The Dutch version of the Yale Physical Activity Scale (YPAS) (Dipietro and others 1993) was used to measure the level of physical activity. This questionnaire is composed of two sections – 1) amount of physical activity/exercise performed during a typical week in the past month and 2) activities performed in the past month. The total energy expenditure summary index (kcal/wk) was calculated.

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2.5. Statistical analysis.

Statistical analyses were performed using Statistical Package of the Social Sciences (SPSS) version 26 (IBM, Amonk, New York, USA) and the statistical software RStudio version 1.1.463 running on R version 3.5.3 (R Foundation for Statistical Computing, Vienna, Austria). Significance was set a priori at two-sided p < 0.05. Average values are presented with mean \pm standard deviation (SD) or median \pm interquartile range (IQR, P75-P25) depending on measurement level and normality of distribution. Firstly, the prevalence of positive frailty components of the FFI are presented as percentages (expressing the number of positive cases divided by the number of the total study population). Secondly, differences in age between robust and pre-frail older adults were investigated by the independent samples T-test. Thirdly, differences between robust and pre-frail older adults (divided into categories based on the positive items of the FFI: (a) pre-frail based on low grip strength, (b) pre-frail based on exhaustion, (c) pre-frail based on weight loss, (d) pre-frail based on slow walking speed) were explored by one-way Analysis of Covariance (ANCOVA). We hypothesized that the relations between muscle endurance, self-perceived fatigue and frailty status might be influenced by sex and age (Collard and others 2012), and thus we corrected for these variables. Next, partial Pearson correlations (corrected for age and sex) were computed to determine the association between frailty score and fatigue (GSmax, FR, GW corrected for body mass, self-perceived fatigue, CPV ratio and CRP). Then, binary logistic regression analysis using forward selection was conducted to assess whether the variables (age, BMI, CRP, body fat, GW and MFI-20 as independent factors) discriminated between robustness and pre-frailty, and odds ratios were calculated. Non-significant parameters were not used in the final models. The presented p values in table 4 are based on the Wald test. To test the hypothesis whether persons with low muscle fatigability and high feelings of self-perceived fatigue are more likely to be pre-frail, we investigated the interaction between muscle fatigability and self-perceived fatigue (GW_{corrected for body weight}* 1/MFI-20) on the occurrence of pre-frailty. Since a high score on the GW is considered to be good, as well as a low score on the MFI-20, a classic interaction computation as the product of these parameters would neutralize the linear increase in interaction score with combined worsening scores on muscle fatigability and self-perceived fatigue. Therefore, we recomputed the MFI-20 scores as 1/MFI-20 for testing its interaction with GW. All these interactions significantly predicted pre-frailty. To better interpret this interaction, a "Capacity to Perceived Vitality" ratio (CPV) ratio was computed as GW_{corrected for body weight} / MFI-20, resulting in high 'combined' fatigue levels when the ratio was low, and low 'combined' fatigue levels when the ratio is high. In total 5 different ratios were computed, CPV-total (GWcorrected for body weight /MFI-20 total fatigue), CPV-general (GWcorrected for body weight /MFI-20 general fatigue), CPV-physical (GWcorrected for body weight /MFI-20 physical fatigue), CPV-redact (GWcorrected for body weight /MFI-20 reduced activity), CPV-redmot (GWcorrected for body weight /MFI-20 reduced motivation) and CPV-mental (GWcorrected for body weight /MFI-20 mental fatigue), resulting in in high 'combined' fatigue levels when the ratio was low, and low 'combined' fatigue levels when the ratio is high. Since the exhaustion component is part of the frailty index, we performed the same analysis on a subgroup, in which persons who scored positive on the CES-D item of the frailty index were excluded. Thus, model 1 includes all participants and model 2 excluded participants who scored positive on exhaustion (FFI). To avoid multicollinearity problems (r>0.80) we have implemented the CPV ratio's in independent regression models. To have a clear visualisation of the independent variables and their relation with pre-frailty, different effect plots were generated. Since we found an interaction between CPV ratios and gender (supplementary table 1), we performed separate logistic regression analyses for males and females for a better clinical interpretation of the results. Lastly, ROC curves were plotted to evaluate and visualize the fit of the logistic regression in distinguishing between robustness and pre-frailty.

3. RESULTS

This study included 405 participants (mean age 83 ± 3 years, flowchart shown in Fig. 1) among which 214 robust (91 males and 123 females) and 191 pre-frail (136 males and 55 females) older adults. Pre-frail participants were significantly older than their robust counterparts (respectively 84 ± 3 and 82 ± 2 years, p<0.001). Low grip strength is the most prevalent positive frailty criterion in the pre-frail older adults, followed by exhaustion and low gait speed (see table 1.). Twenty-five participants scored positive on more than one frailty criterion, of whom 14 on exhaustion combined with low grip strength (see table 1).

The participants' characteristics are shown in table 2. Robust participants had significantly higher GW (p<0.001), less fatigue on the MOB-T (p<0.05) and MFI-20 (total, general, physical) (p<0.05) and better CPV ratios (p<0.001). Pre-frail older adults who scored negative on the exhaustion item of the FFI index still showed lower levels of GW (p<0.001) and CPV ratios (p<0.001), and higher levels on the MFI-20-total (p<0.05), MFI-20 physical fatigue (p<0.05), MFI-20 general fatigue (p<0.05) and MOB-T (p<0.05) compared to their robust counterparts. When looking at pre-frail adults who scored positive on the exhaustion item of the FFI, it can be seen that they also scored lower on all muscle fatigue and CPV levels, and higher on self-perceived fatigue (except of MFI-20 mental fatigue).

Table 3 shows that in the participants who scored negatively on exhaustion, better GW was related to a better score on the FFI. No significant relations between MFI-20 and the FFI score were found in this group. All CPV ratios were negatively related to the FFI, considering lower ratios for a higher FFI score. Considering all participants, higher levels of fatigue based on the MFI-20 (total and subscales) and MOB-T scores were related to higher frailty scores. In contrast, lower CPV ratios were significantly related to higher FFI scores.

A binary logistic regression was performed per CPV ratio (sub)scale with pre-frailty as dependent variable (table 4 and supplementary table 1). The regression analyses indicated that in all models age and the interaction

between sex and CPV ratios were significantly associated with pre-frailty. CRP, physical activity and percentage body fat, did not significantly contribute and thus were not retained in the final models. The risk for being prefrail increases with age and decreases with a higher CPV ratio. A significant interaction between CPV ratios and sex was found (supplementary table 1). Figure 2 shows this interaction and indicates that with increasing CPV ratio, the risk for pre-frailty decreases more for females compared to males. For a better clinical interpretation of this interaction, we performed the same analyses for males and females separately (table 4). In our study, males have a higher risk to be pre-frail compared to females. With increasing CVP ratio, the risk for pre-frailty decreases significantly. These results were found for all CPV ratios and the visualisations of these effects are shown in supplementary Figures 1a-2f. A separate logistic regression was performed per gender; in females every unit increase in fatigue ratio showed to decrease the risk for pre-frailty with 77.6% (CPV total model 1 OR 0.224; 95% CI: 0.114-0.440). In males these results were also present but less strong as for females (34,4%; CPV total model 1 OR 0.656; 95% CI: 0.465-0.926). The odds ratio for the total group using the CPV-total scale was slightly higher when excluding persons who scored negative on the CES-D item for males (model 1 OR 0.656; 95% CI: 0.465-0.926: model 2 OR 0.711; 95% CI: 0.501-1.008), for females we found the opposite (model 1 OR 0.224; 95% CI: 0.114-0.440: model 2 OR 0.173; 95% CI: 0.076-0.391) . ROC curves were plotted in order to evaluate and visualize the fit of the logistic regression for all total models (supplementary Figures 3a- 3L). The highest AUC was observed for CPV-physical ratio (AUC: 0.782, see supplementary table 1).

4. **DISCUSSION**

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In this study, we investigated whether the combination of muscle endurance and self-perceived fatigue is related to frailty in well-functioning older adults aged 80 and over. The main finding of this study was that also in participants who are pre-frail but did not show clinical signs of fatigue on the exhaustion component in the FFI, still experienced significantly higher fatigue levels (lower muscle endurance, higher self-perceived fatigue and lower CPV levels) compared to their robust counterparts. Logistic regression analysis showed that age, gender, CPV ratio, and the interaction between CPV ratio and gender were significantly associated with pre-frailty, in community dwelling octogenarians, with overall predictive accuracy of 77%. As a significant interaction was found between muscle fatigue and self-perceived fatigue, we computed a CPV ratio since this value is clinically more informative to interpret than the statistical interaction. Females were less likely to be pre-frail and with increasing CVP ratio the risk for pre-frailty decreased significantly, for males this effect was also found but less strong. Muscle endurance correlated significantly but moderately with self-perceived fatigue, suggesting that both provide complementary information regarding the clinical expression of fatigue. Therefore, the interplay between these two variables expressed as the CPV ratio might be a good fatigue "index" in frailty assessment. All CPV ratios showed a stronger relationship with the FFI than muscle endurance and self-perceived fatigue separately; and logistic regression showed that CPV ratio can discriminate between robust and pre-frailty status. Results showed that age and the CPV ratio were significantly associated to the frailty status, even when the prefrail participants did not score positively on the fatigue item in the FFI.

In this study we found that low muscle endurance is related to higher frailty scores, this is in line with earlier research where it was shown that muscle endurance and frailty share the same biomedical determinants (i.e. aging, disease, inflammation, physical inactivity, malnutrition, hormonal deficiencies, subjective fatigue and neuromuscular function and structure) (Theou and others 2008). Also, both muscle endurance and selfperceived fatigue are part of the "vicious cycle of frailty" as proposed by Fried and others (2004). The group of Westerblad showed that muscle endurance decreases before the onset of muscle weakness in a mouse model of premature aging (Yamada and others 2012). This implies that low muscle endurance is probably an important early marker for frailty (Kent-Braun and others 2002). Thereby, it has been shown that muscle endurance can help to discriminate older women with different degrees of frailty (De Dobbeleer and others 2018). Interestingly, we found significant differences in self-perceived fatigue and muscle endurance between robust and pre-frail participants, also within participants who scored negatively for the exhaustion item in the FFI. This could imply that persons who do not score positively on exhaustion in the FFI show already early signs of fatigue. However, this was not detected by the CES-D questions, one of the key elements in the FFI (Orme and others 1986). The CES-D items in the FFI were included by Fried and others (2001) assuming that they indicate exhaustion, comprising low energy and poor endurance. These CES-D items are associated with the VO2max and are able to predict cardiovascular disease (Fried and others 2001). Multiple adapted FFI versions used other fatigue instruments to capture the exhaustion item including generic questions such as feeling tired or feeling worn out, and questions from the 36-item Short Form Health questionnaire (Knoop and others 2019). Despite the large array of operationalization, all exhaustion items belong to the same ICF category (Azzopardi and others 2016). However, it is questionable if these tools are capable to detect early stages of fatigue. In our study, we found that participants show already signs of losses in reserve capacity expressed by a low CPV ratio before they score positive on the CES-D exhaustion items. Therefore, it can be hypothesized that including a measurement of muscle endurance in combination with self-perceived fatigue can be a valuable asset in early physical frailty identification. This will help developing preventive and therapeutic interventions.

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In contrast to other studies (Cao Dinh and others 2019; Walston 2002), we did not find a significant relationship between inflammation and frailty status. In addition, no significant relationship was found between CRP and muscle endurance. These results are contradictory to other articles that showed a relationship with inflammation in community-dwelling older persons (Bautmans and others 2007; Beyer and others 2012a) and in hospitalized geriatric patients (Arnold and others 2017; Beyer and others 2012b). However, literature in healthy community-dwelling older adults is scarcer, which could explain the lack of similar findings. In our study, we included relatively healthy, high performing participants who were not frail and who performed sufficient physical activity, which might decrease the risk for inflammaging. As CRP was used to exclude participants with ongoing inflammatory pathology; not primarily to quantify low-grade inflammation we did not use high sensitivity (hs)CRP. Therefore, we might have missed the influence of low-grade inflammation in our statistical analysis.

We hypothesized that fatigue is a valid parameter for intrinsic capacity, a concept introduced by the World Health Organization (WHO). Intrinsic capacity expresses all physical and mental capacities of an individual (Cesari and others 2018). Intrinsic capacity and the loss of reserve capacity as seen in frailty (Baerd JR 2019) are highly interrelated. Where intrinsic capacity is more presenting the reserves of an individual, frailty focuses more on the deficit accumulation with ageing (Fried and others 2001). Reduced intrinsic capacity will have implications on the level of physical functioning. The WHO suggests that changes in intrinsic capacity are likely to start at midlife, before persons will experience problems in daily functioning. It is hypothesized that some parameters will influence the depletion of reserve capacity leading to enlarged risk for frailty. The combination of high selfperceived fatigue and low muscle endurance could be recognized as deficits modifying intrinsic capacity and partly determine the development of frailty. However, the exact role of these two parameters in this concept has not been investigated before. The results of this study show that muscle fatigue and self-perceived fatigue are a good indicator for early stages of pre-frailty. Males were more likely to be pre-frail in this cohort, especially those with a low CPV ratio. This is in contrast with earlier research where females were more likely to be prefrail (Collard and others 2012; Gordon and others 2017). However, this is the first report on persons aged 80 and over. Since in our sample there were more males than females, this could have influenced the results. Females with better CPV ratios are less likely to be pre-frail; in males the same results were found but the effects were less strong. This suggests that the level of fatigue might be more important for females in the pathophysiology for frailty, considering that other characteristics are involved in males and females.

As far as we know, this is the first study suggesting that the CPV ratio could be an additional characteristic in the early identification of physical frailty. However, our study is based on cross-sectional data, and the results should be confirmed prospectively. The ROC curves found in this study show acceptable to good AUC value (range 0.755 – 0.782)(Mandrekar 2010), suggesting a decent prediction of pe-frailty. For clinical practice, the results showed that robust older adults showed on average a CPV total score >1, while pre-frail older adults had on average a score <1. Every unit increase of CPV score decreased the risk for pre-frailty with 34% for males and 78% for females.

This study has certain strengths and limitations. The strength of this study relies in the inclusion of a relatively large and unique cohort considering the age and independence level of the participants, providing unique data on potentially early determinants of frailty. It cannot be excluded that the lack of statistical differences and relationships between frailty status and inflammation is due to the fact that on the one hand we recruited participants without ongoing inflammatory pathology, and on the other hand that the CRP-assay lacked sensitivity. The finding that males were more likely to be pre-frail might be due to an uneven distribution of males and females in our study sample. However, the relationship between fatigue and frailty status was similar when analysing female and male separately. The total accuracy rate of the models leaves room for improvement and implies that other variables should be included in the early detection of physical frailty. However, we have included percentage body fat, physical activity and inflammation, but these parameters did not contribute significantly. On the other hand, it might be possible that the CPV captures a phenotype that is not completely

detected by the FFI. Finally, all findings in this study are based on a cross-sectional analysis, and prospective confirmation is advocated, as pre-frailty is an unstable condition and reverse causation cannot be excluded.

5. CONCLUSION

Our study showed that older pre-frail persons without clinical signs of exhaustion experience significantly higher fatigue levels (i.e. lower muscle endurance, higher self-perceived fatigue and lower CPV levels) compared to their robust counterparts. The "Capacity to Perceived Vitality" (CPV) ratio, representing the ratio between muscle endurance and self-perceived fatigue, can be considered as a good candidate as early marker of (pre-frailty. For clinical practice, every unit increase CPV ratio decreased the risk for pre-frailty with 34% for males and 78% for females. CPV ratio could therefore be a good tool to identify subclinical fatigue in the context of physical (pre-)frailty.

6. STATEMENTS

6.3. Acknowledgement

N/A

7.1 Statements of ethics

All participants gave their written informed consent and the study protocol was approved by ethical committee of the Universitair Ziekenhuis Brussel (B.U.N. 143201421976).

7.2 Conflict of interest

The authors have no relevant conflicts of interest to declare.

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7.3 Author Contributions

V.K. contribute to the data collection, data analysis, and writing of the manuscript. A.C., A.D., R.V.A. and S.V. contribute to data collection and reviewed the manuscript. S.v.L. contribute to the statistical analyses. A.S. and B.J. contribute to supervision and reviewed the manuscript. I.B. conceived the study, was responsible for funding acquisition, supervision, reviewing and editing of the manuscript. On behalf of the Gerontopole Brussels Study Group

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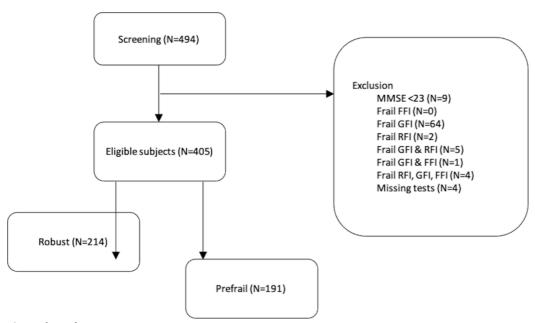


Fig 1. Flow chart. *MMSE: Mini-mental state examination, FFI: Fried Frailty Index, GFI: Groningen Frailty Index, RFI: Rockwood Fraily Index*

Interaction effect 'Gender x CPV-total-BW'

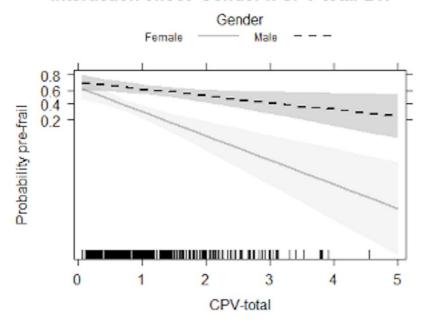


Fig. 2. Plot of the interaction between gender and CPV-total ratio on the predictive value of pre-frailty.This figure shows the interaction factor between gender and CPV-total that was found in the logistic regression analysis. The risk for pre-frailty is lower for females compared to males. With an increased CPV ratio decreased the chance to be pre-frail for females.

Table 1. Prevalence of Fried phenotype criteria

Variables	No (other) frailty		Combined with Low		Combined with			Combined with Low				
	markers	S		grip st	rength		Exhaus	stion		gait speed		
	Total	Male	Female	Total	Male	Female	Total	Male	Female	Total	Male	Female
No frailty markers (robust)	214	91	123									
	52.8%	22.5%	30.4%									
Positive on Low grip strength	150	118	32									
	37.0%	29.1%	7.9%									
Positive on Exhaustion	9	2	7	14	8	6						
	2.2%	0.5%	1.7%	3.5%	2.0%	1.5%						
Positive on Low gait speed	6	1	5	3	3	0	2	2	0			
	1.5%	0.2%	1.3%	0.7%	0.7%	0%	0.5%	0.5%	0%			
Positive Unintentional	1	0	1	6	2	4	0	0	0	0	0	0
weight loss	0.2%	0%	0.2%	1.5%	0.5%	1%	0%	0%	0%	0%	0%	0%

%: Percentage; N: Number of participants; Percentages indicate the number of positive cases in the specific frailty item and in combination with other frailty items. Different combinations were present; positive on low grip strength combined with exhaustion, positive on low grip strength combined with low gait speed and positive on low grip strength combined with unintentional weight loss, positive on exhaustion combined with low gait speed and positive on exhaustion combined with unintentional weight loss.

Table 2. Participants' characteristics according to pre-frailty profile

Variables	Robust	Pre-frail				
	Robust	Pre-Frail	Positive on Low grip	Positive on Low gait	Positive on	Positive Unintentional
	n= 214	n= 191	strength	speed	Exhaustion CES-D	weight loss
			n= 173	n= 11	n= 25	n= 7
	Males n=91	Males n=136	Males n=131	Males n=6	Males n=12	Males n=2
. ,	Females n=123	Females n=55	Females n=42	Females n=5	Females n=13	Females n=5
Age (years)	82.30±2.13	83.79±3.15 [†]	83.76±3.08 [†]	83.84±3.81 [†]	84.23±3.03*	82.38±3.22
Height (m)	1.6±0.1*	1.7±0.1*	1.7±0.1 [†]	1.7±0.1*	1.6±0.1*	1.6±0.1*
Weight (kg)	71.4±12.8	73.6±11.2	73.9±11.1	72.4±12.8	74.6±14.8*	65.2±12.8
BMI	26.2±3.3	26.6±3.9	26.8±3.4	23.8±8.4*	27.2±3.5	24.5±2.7
Body fat (%)	34.6±6.3	33.0±7.0	32.5±6.8	33.9±8.7	36.3±6.9	34.3±5.0
MMSE	28.0±1.6	27.7±1.8	27.5±1.8	27.5±1.6	27.8±1.9	27.7±1.7
Physical activity (Kcal) ⁵	6496.0±3747.0	5942.8±3846.8	6093.1±3967.3	5256.6±3120.3	4745.5±2273.5	6455.8±4164.1
Grip strength (Kpa)	65.1±16.4	52.5±14.1 [†]	51.9±13.8 ⁺	60.3±15.3 ⁺	50.8±14.9 [†]	47.9±19.3 [†]
CRP	2.4±5.2	3.6±6.9	3.6±7.1	1.7±1.8	6.1±11.8*	3.8±4.1
Fatigue resistance ¹	70.6±34.7	67.7±38.4	68.1±38.8	60.7±42.4	64.8±35.0	61.2±32.6
Grip work ²	3376.4±1707.5	2714.2±1752.7 [†]	2685.9±1683.1 [†]	2966.3±2683.0 [†]	2596.3±2125.1 [†]	1976.2±824.8 [†]
Grip work corrected for body weight ²	48.0±24.2	37.4±23.9 [†]	37.0±23.3 [†]	41.3±34.2 [†]	35.9±27.1 [†]	30.9±13.7 [†]
MOB-T ¹	0 (0-1)	0 (0-2)*	0 (0-2)	2 (0-3)*	2 (0-4) †	0 (0-1)*
MFI-20 total fatigue score (20-100) ²	40 (31-49)	45 (32-56)*	44 (32-55)	52 (45-61)*	60 (43-68) [†]	31 (26-58)
MFI-20 General fatigue (4-20) ²	8 (5-10)	9 (6-12)*	8 (5-11)*	10 (9-13)*	13 (10-15) [†]	9 (5-9)*
MFI-20 physical fatigue (4-20) ²	7 (5-10)	9 (5-12)*	8 (5-11)*	11 (9-13) [†]	12 (10-16) [†]	6 (5-12)*
MFI-20 reduced activity (4-20) ²	8 (6-11)	10 (6-13)	10 (6-13)	11 (9-13)	13 (10-15) [†]	8 (4-13)
MFI-20 reduced motivation (4-20) ²	8 (5-11)	9 (6-11)	9 (6-11)	11 (8-12)	11 (8-14)*	7 (6-10)
MFI-20 mental fatigue (4-20) ²	7 (5-11)	8 (5-11)	8 (5-11)	7 (5-10)	7 (5-12)	5 (4-12)
CPV- total ⁴	1.3±0,8	1.0±0.8 [†]	1.0±0.8 [†]	0.8±0.5 [†]	0.7±0.7 [†]	0.9±0.4 [†]
CPV- general ⁴	7.0±4.5	5.4±4.5 [†]	5.3±4.5 [†]	3.8±2.6 [†]	3.3±2.5 [†]	4.4±2.2 [†]
CPV- physical ⁴	7.4±5.0	5.6±4.9 [†]	5.6±4.9 [†]	3.5±2.5 [†]	3.9±4.9⁺	4.4±2.0 [†]
CPV- redact ⁴	6.7±4.7	4.8±4.2 [†]	4.8±4.2*	3.8±2.6 [†]	3.4±3.1 [†]	4.7±2.5 [†]
CPV- redmot ⁴	6.9±4.5	5.1±4.3 [†]	5.1±4.3 [†]	4.0±2.7 [†]	4.2±4.8 [†]	4.4±2.1 [†]
CPV- mental ⁴	7.2±4.8	5.4±4.3 [†]	5.4±4.3 [†]	5.7±4.6 [†]	5.4±5.5 [†]	5.0±2.2 [†]

ANCOVA corrected for age and gender, values expressed as mean ± SD

548 Bonferroni post-hoc test 549 m: meters; kg: kilogram; 550 Vitality ¹: 1 missing value

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m: meters; kg: kilogram; %: percentage; Kcal: calories; Kpa: Kilo pascal; MOB-T: Mobility Mobility Tiredness scale; MFI-20: Multidimensional Fatigue Inventory; CPV: Capacity to Perceived Vitality 1: 1 missing value; 2; 2 missing values; 4: 4 missing values; 5: 5 missing values

^{*}Significant different from robust (p<0.05) on all ANCOVA analysis corrected for age and sex (except for age)

[†]Significant different from robust (p<0.001) on all ANCOVA analysis corrected for age and sex (except for age)

Table 3. Relationships between muscle endurance, self-perceived fatigue, CRP, physical activity and robustness or prefrailty.

	All participants N=405 Participants who a fatigued on CES-D			
	Grip Work	Fried score	Grip Work	Fried score
Grip Work		-0.24 [†]		-0.22 [†]
Body fat (%)	-0.27 [†]	0.07	-0.28 [†]	0.05
Physical activity (Kcal)	0.12*	-0.03	0.13*	-0.01
CRP	-0,08	0.10	-0.08	0.07
MFI-20 total fatigue score (20-100)	-0.14*	0.18 [†]	-0.13*	0.05
MFI-20 General fatigue (4-20)	-0.11*	0.19 [†]	-0.12*	0.06
MFI-20 physical fatigue (4-20)	-0.21 [†]	0.22 [†]	-0.19 [†]	0.09
MFI-20 reduced activation (4-20)	-0.15*	0.14*	-0.15*	0.03
MFI-20 reduced motivation (4-20)	-0.09	0.11*	-0.08	0.01
MFI-20 mental fatigue (5-20)	0.01	0.02	0.02	0.00
CPV- total		-0.26 [†]		-0.23 [†]
CPV- general		-0.26 [†]		-0.21 [†]
CPV- physical		-0.26 [†]		-0.22 [†]
CPV- redact		-0.25 [†]		-0.21 [†]
CPV- redmot		-0.24 [†]		-0.22 [†]
CPV- mental		-0.22 [†]		-0.22 [†]

Values represent Partial correlation coefficients corrected for age and sex*p<0.05, †p<0.01

^{%:} percentage; Kcal: calories; GW: Grip Work; MFI-20: Multidimensional Fatigue Inventory; CPV: Capacity to Perceived Vitality

Table 4. Logistic regression with GW/MFI-total score discriminating prefrailty/robustness for males and females

Males							
		Significant variables	В	S.E.	Sig.	Odds ratio	95% confidence interval
CPV- total	Model 1	Age	0.234	0.061	<0.001	1.264	1.122-1.423
	R ² : 0.138	CPV- total	-0.422	0.176	0.016	0.656	0.465-0.926
	Model 2	Age	0.222	0.061	<0.001	1.248	1.108-1.406
	R ² : 0.120	CPV- total	-0.342	0.178	0.055	0.711	0.501-1.008
CPV- general	Model 1	Age	0.233	0.061	< 0.001	1.262	1.120-1.421
	R ² : 0.130	CPV- general	-0.065	0.031	0.035	0.937	0.882-0.995
	Model 2	Age	0.221	0.061	< 0.001	1.247	1.107-1.405
	R ² : 0.113	CPV- general	-0.050	0.031	0.108	0.951	0.894-1.011
CPV- physical	Model 1	Age	0.231	0.061	<0.001	1.260	1.118-1.419
	R ² : 0.130	CPV- physical	-0.059	0.028	0.035	0.943	0.893-0.996
	Model 2	Age	0.219	0.061	<0.001	1.245	1.105-1.403
	R ² : 0.114	CPV- physical	-0.046	0.028	0.104	0.955	0.904-1.009
CPV- redact	Model 1	Age	0.239	0.062	<0.001	0.270	1.126-1.433
	R ² : 0.144	CPV- redact	-0.085	0.032	0.009	0.919	0.862-0.979
	Model 2	Age	0.227	0.062	<0.001	1.254	1.112-1.415
	R ² : 0.126	CPV- redact	-0.071	0.033	0.030	0.932	0.874-0.933
CPV- redmot	Model 1	Age	0.237	0.061	<0.001	1.267	1.124-1.428
	R ² : 0.140	CPV- redmot	-0.079	0.032	0.013	0.924	0.868-0.983
	Model 2	Age	0.224	0.061	<0.001	1.251	1.110-1.410
	R ² : 0.123	CPV- redmot	-0.065	0.032	0.042	0.937	0.880-0.998
CPV- mental	Model 1	Age	0.233	0.060	<0.001	1.263	1.122-1.422
	R ² : 0.138	CPV- mental	-0.071	0.030	0.018	0.931	0.878-0.988
	Model 2	Age	0.220	0.061	<0.001	1.246	1.106-1.404
	R ² : 0.122	CPV- mental	-0.061	0.030	0.045	0.941	0.886-0.999
Females							
CPV- total	Model 1	Age	0.245	0.073	0.001	1.278	1.107-1.474
	R ² : 0.279	CPV- total	-1.495	0.344	<0.001	0.224	0.114-0.440
	Model 2	Age	0.208	0.077	0.007	1.231	1.059-1.431
	R ² : 0.290	CPV- total	-1.757	0.417	<0.001	0.173	0.076-0.391

CPV- general	Model 1	Age	0.240	0.071	0.001	1.272	1.106-1.463
	R ² : 0.292	CPV- general	-0.294	0.068	<0.001	0.745	0.653-0.851
	Model 2	Age	0.210	0.075	0.005	1.234	1.064-1.431
	R ² : 0.271	CPV- general	-0.296	0.075	<0.001	0.744	0.642-0.862
CPV- physical	Model 1	Age	0.247	0.074	0.001	1.280	1.108-1.480
	R ² : 0.292	CPV- physical	-0.271	0.063	<0.001	0.763	0.674-0.863
	Model 2	Age	0.210	0.079	0.008	1.234	1.056-1.441
	R ² : 0.327	CPV- physical	-0.355	0.082	<0.001	0.701	0.597-0.823
CPV- redact	Model 1	Age	0.227	0.070	0.001	1.255	1.094-1.440
5. V . Cdd Ct	R ² : 0.241	CPV- redact	-0.234	0.060	<0.001	0.791	0.703-0.890
	Model 2	Age	0.195	0.074	0.008	1.215	1.051-1.404
	R ² : 0.227	CPV- redact	-0.248	0.070	<0.001	0.780	0.680-0.895
CPV- redmot	Model 1	Age	0.241	0.071	0.001	1.272	1.106-1.463
	R ² : 0.229	CPV- redmot	-0.220	0.059	<0.001	0.802	0.714-0.901
	Model 2	Age	0.205	0.075	0.006	1.228	1.059-1.424
	R ² : 0.255	CPV- redmot	-0.289	0.075	<0.001	0.749	0.646-0.868
CPV- mental	Model 1	Age	0.240	0.070	0.001	1.272	1.110-1.458
	R ² : 0.209	CPV- mental	-0.191	0.054	<0.001	0.826	0.743-0.917
	Model 2	Age	0.225	0.076	0.003	1.252	1.079-1.454
	R ² : 0.268	CPV- mental	-0.293	0.073	<0.001	0.746	0.647-0.860

Logistic regression analysis to predict pre-frailty in males and females separately; Model 1: All participants included n= 405; Model 2: Participants who are not fatigued on CES-D; R2: R square

LEGENDS TO THE SUPPLEMENTARY FIGURES

Supplementary Fig. 1. Plots of the interaction between gender and CPV ratios on the predictive value of prefrailty. Fig. 1a CPV-general ratio, Fig. 1b CPV-physical ratio, Fig. 1c CPV-redact ratio, Fig. 1d CPV-redmot ratio, Fig. 1e CPV-mental ratio

Supplementary Fig. 2. Plots of the interaction between gender and CPV-total ratio on the predictive value of pre-frailty in participants who were not exhausted on CES-D. Fig. 2a CPV-total ratio, Fig2b CPV-general ratio, Fig. 2c CPV-physical ratio, Fig. 2d CPV-redact ratio, Fig. 2e CPV-redmot ratio, Fig. 2f CPV-mental ratio

Supplementary Fig. 3. ROC Curves reflecting the discriminative value of CPV ratios for identifying pre-frailty In all participants Fig. 3a CPV-total ratio, Fig 3b CPV-general ratio, Fig. 3c CPV-physical ratio, Fig. 3d CPV-redact ratio, Fig. 3e CPV-redmot ratio, Fig.3f CPV-mental ratio; and in participants who were not exhausted on CES-D Fig. 3g CPV-total ratio, Fig. 3h CPV-general ratio, Fig. 3i CPV-physical ratio, Fig. 3j CPV-redact ratio, Fig.3k CPV-redmot ratio, Fig.3l CPV-mental ratio