

Rate of force development and rapid muscle activation characteristics of knee extensors in very old men

Giorgio Varesco, Loïc Espeit, Léonard Féasson, Thomas Lapole, Vianney

Rozand

▶ To cite this version:

Giorgio Varesco, Loïc Espeit, Léonard Féasson, Thomas Lapole, Vianney Rozand. Rate of force development and rapid muscle activation characteristics of knee extensors in very old men. Experimental Gerontology, 2019, 124, pp.110640. 10.1016/j.exger.2019.110640. hal-02409448

HAL Id: hal-02409448 https://hal.science/hal-02409448

Submitted on 25 Oct 2021

HAL is a multi-disciplinary open access archive for the deposit and dissemination of scientific research documents, whether they are published or not. The documents may come from teaching and research institutions in France or abroad, or from public or private research centers. L'archive ouverte pluridisciplinaire **HAL**, est destinée au dépôt et à la diffusion de documents scientifiques de niveau recherche, publiés ou non, émanant des établissements d'enseignement et de recherche français ou étrangers, des laboratoires publics ou privés.



Distributed under a Creative Commons Attribution - NonCommercial 4.0 International License

Rate of force development and rapid muscle activation characteristics of knee extensors in very old men.

- 3 Giorgio Varesco^a, Loïc Espeit^a, Léonard Feasson^{a,b}, Thomas Lapole^a & Vianney Rozand^{a,*}
- 4 ^aUniv Lyon, UJM Saint-Etienne, Laboratoire Interuniversitaire de Biologie de la Motricité,
- 5 EA 7424, F-42023, Saint-Etienne, France
- 6 ^bUnité de Myologie, Service de Physiologie Clinique et de l'Exercice, Centre Référent
- 7 Maladies Neuromusculaires Rares Euro-NmD, CHU de Saint-Etienne, F-42055 Saint
- 8 Etienne, France.
- 9
- 10 *Corresponding author at : Laboratoire Interuniversitaire de Biologie de la Motricité, Campus
- 11 Santé Innovations IRMIS, 10 rue de la Marandière, 42270 Saint Priest en Jarez, France.
- 12

Email adresses: giorgio.varesco@univ-st-etienne.fr (Giorgio Varesco), l.espeit@univ-stetienne.fr (Loïc Espeit), leonard.feasson@univ-st-etienne.fr (Léonard Feasson),
Thomas.lapole@univ-st-etienne.fr (Thomas Lapole), vianney.rozand@univ-st-etienne.fr
(Vianney Rozand).

17

18 Abbreviations

19

KE	knee extensors
F _{max}	maximal force
RFD	rate of force development
RFD _{max}	maximal rate of force development
tRFD _{max}	time to maximal rate of force development
RMS	Root Mean Square
RER	rate of EMG rise
MVC	maximal voluntary contraction
VL	vastus lateralis
CV	coefficient of variation

20

21 Highlights

- Rapid force development and activation capacity are reduced in the eight decade of
 age.
- Rate of force development at early (0-50 ms) but not late (100-200 ms) time intervals
 were lower in very old adults when normalized to the maximal force.
- Rate of force development capacity decline of knee extensors is mainly affected by
 age on the early phase of the contraction (first 50 ms from the force onset).

28 Abstract

The age-related decrease in neuromuscular performance is usually accentuated in very old 29 age (>80 yr) as evidenced by a marked reduction in maximal force production. However, 30 little is known on the ability to rapidly produce force, which limits daily activities and 31 32 increase the risk of falling. We aimed to assess rapid force production characteristics and rate of muscle activation of the knee extensors in 15 very old $(82\pm1 \text{ yr}) vs$ young $(24\pm4 \text{ yr})$ men. 33 Maximal force (F_{max}) and maximal rate of force development (RFD_{max}) were determined 34 during separated specific isometric contractions. EMG from the vastus lateralis was analyzed 35 to assess the rate of EMG rise (RER). Finally, RFD and RER were examined at time intervals 36 of 0-50, 50-100, 100-200, 0-200 ms and 0-30, 0-50, 0-75 ms, respectively. We reported lower 37 F_{max} (414±91 N vs. 661±139 N) and absolute RFD_{max} (8720±2056 N*s⁻¹ vs. 5700±2474 N*s⁻¹) 38 in the very old men compared to young men (P < 0.01). When normalized to F_{max}, RFD_{max} 39 40 was similar between groups. Normalized RFD at 0-50 and 0-200 ms were lower (-34% and -46%, P=0.04) for the very old men, while no difference was observed at 50-100 ms and 100-41 200 ms. RER values were higher (~346%, P<0.01) for the young men at every time interval. 42 43 These data suggest that the decline in RFD_{max} is associated to the decrease in F_{max} .

- 44 Impairments in RFD were accentuated in the first phase of the contraction, which has been
- 45 associated in the literature to impairments in neural factors associated with aging.

46

- 47 Keywords: aging, rate of force development, rate of EMG rise, neuromuscular performance,
- 48 knee extensors.
- 49

50 **1. Introduction**

The ability to quickly produce a high amount of force is strongly related to the completion of 51 daily activities and can be further associated with the risk of falling in older adults [1,2,3]. A 52 53 useful tool to assess the capacity to rapidly increase force is the rate of force development (RFD), considered to be more sensitive than maximal isometric force (F_{max}) to detect changes 54 in neuromuscular function in clinical and old populations [4,5]. Another advantage of RFD is 55 the possibility to differentiate various phases of the force-time curve to better detect 56 performance changes after a training protocol or in clinical vs healthy populations [4,6,7]. 57 58 This analysis can be further implemented by studying the rise in the EMG signal. The rate of EMG rise (RER) has been used to evaluate the rate of muscle activation concurrent to the rise 59 in force in order to account for the neural factors that contribute to RFD [8,9,10]. 60

61 Healthy older adults present lower peak RFD than young adults for various upper and lower limb muscle groups, including the knee extensors [11,12], plantar flexors [10], dorsiflexors 62 [2] and elbow extensors [13]. Conflicting results were reported when RFD was normalized to 63 F_{max} : while some studies showed greater values in young vs older adults [2,10,12,14], others 64 did not [1,15]. The discrepancy in the results could be due to different time intervals 65 examined and different techniques used to normalize RFD by F_{max} (e.g. some author 66 calculated RFD on the force-time curve, while others used %MVC on the x-axis and 67 measured the RFD value at the point when the force reached 10%, 30% or 50% of the MVC 68 69 [10]). However, it remains unclear whether the age-related decrease in RFD capacity is only due to the decrease in F_{max}. For instance, RFD may be influenced more than F_{max} by changes 70 at the motor units level (i.e. loss of fast-type motor units [16]), or neural impairments (i.e. 71 72 motor unit discharge rate and doublets [2]).

Regarding the methodology, it has to be noted that a large part of the available literature
measured RFD during maximal isometric voluntary contractions (MVC), asking the

75 participants to exert isometric contractions as strong and as fast as possible (e.g.[10,11,12]). 76 Instead, recent recommendations suggest to perform specific impulsive isometric contractions as fast as possible to investigate RFD separately from F_{max} [4]. The rigor in the instructions is 77 78 essential, as Sahaly et al. [17] showed a significant improvement in peak RFD when participants were instructed to push "fast" rather than "hard and fast". Consequently, RFD 79 analyzed during MVC could be suboptimal and may not represent the maximal capacity to 80 81 rapidly increase force. Furthermore, considering that older adults exhibit greater variability in force measurements compared to young adults [18], measuring RFD during an MVC may 82 83 lead to inconsistent results and a misinterpretation of the age-related changes in RFD.

In this scenario, also the age group of participants is determinant: several cross-sectional and 84 longitudinal studies focusing on the advanced stages of human life documented a sudden 85 86 acceleration in the decline of neuromuscular function starting from the ninth decade [19,20]. 87 However, RFD has been mainly studied in individuals aged between 60 and 80 years and only one study focused on the RFD performance in octogenarians [13]. In their study, 88 89 Bellumori, Jaric, & Knight [13] analyzed the absolute peak RFD from MVC of the elbow extensors and index finger abductors in very old adults and observed a ~50% reduction in 90 RFD compared to young adults. Considering that the age-related impairments in 91 neuromuscular performance are typically greater in the lower limb compared to upper limb 92 muscles [19,21] with functional repercussions on balance and autonomy [22], investigating 93 94 the capacity to rapidly increase force of the lower limb muscles is particularly relevant in very old adults. Identifying the mechanisms of the possible impairment in RFD may help to 95 design specific interventions to preserve the neuromuscular function in very old age. 96

97 In the present study, we aimed to investigate the age-related changes in RFD 98 characteristics and RER of knee extensors (KE) between healthy young (<30 years) and 99 active very old men (>80 years). We hypothesized that 1) both absolute and relative (to F_{max}) 100 RFD characteristics of the KE would be considerably reduced in the octogenarians compared
101 to the young adults during specific impulsive contractions and 2) RER of the *vastus lateralis*102 would be lower in very old adults.

103 **2. Methods**

104 *2.1.Participants*

Twelve recreationally active males of 24±4 years (range 20 - 33 yr, 182±6 cm, 76±8 kg) and 105 fifteen active very old men of 82±1 years (range 80 - 83 yr, 168±6 cm, 71±9 kg) volunteered 106 to participate in the experiment and gave their written consent. Very old participants were 107 108 selected from the PROgnostic indicator OF cardiovascular and cerebrovascular events (PROOF) cohort, which is a prospective longitudinal cohort of old adults started in 2001 [23]. 109 Individual physical activity of the participants from this cohort is regularly assessed using a 110 111 wrist wearable accelerometer (wGT3X-BT, ActiGraph, Pensacola, USA) and analyzed using ActiLife software (v6.13.4, ActiGraph, Pensacola, USA). Considering those data, all the very 112 old men were considered as physically active (8840±1972 steps.day⁻¹, [24]). All participants 113 were free from musculoskeletal or neurological disorders. Before the experimental session, 114 participants were asked to avoid strenuous exercises for at least 24 hours. All the 115 experimental procedures were approved by the Comité de Protection des Personnes Sud-Est 1 116 (France), and the Agence Nationale Sanitaire et du Médicament (ANSM-France) and was 117 performed in accordance with the Declaration of Helsinki (ClinicalTrials.gov, Identifier: 118 119 NCT02675192).

120 *2.2.Mechanical recording and electromyography*

Participants were seated upright on a custom-built chair with hip angle set at 120° (180° corresponds to full extension) and knee angles set at 90° of flexion (Figure 1A). Knee extensors force of the right leg was assessed using a calibrated force transducer (Omega Engineering Inc., Stamford, CT, USA). Extraneous movements of the upper body were

minimized by three belts across the thorax and waist. The force transducer was firmly 125 attached above the malleoli. Force was sampled at 2 kHz and recorded without analogical or 126 digital filter using a Power-Lab data acquisition system (16/30-ML880/P, ADInstruments, 127 Bella Vista, Australia). Passive resting force was subtracted from the signal so that the 128 baseline was set at 0 N. Electromyographic (EMG) activity was recorded from the vastus 129 *lateralis* (VL) muscle of the right leg using pairs of self-adhesive Ag/AgCl surface electrodes 130 (recording diameter = 10 mm, Kendall MediTrace foam electrode, MA) placed in bipolar 131 configuration over the muscle belly with an interelectrode (center to center) distance of 30 132 133 mm. Electrodes were placed following the SENIAM project recommendations. A reference electrode was placed on the right patella. Prior to electrode placement, the skin was shaved, 134 abraded and cleaned with alcohol swab to lower the impedance (<5 k Ω , 87v, Fluke Corp., 135 136 Everett, WA, USA). EMG and force signals were digitized simultaneously at the same sampling rate (2 kHz) by PowerLab System (16/30- ML880/P, AD Instruments, Bella Vista, 137 Australia). Force and EMG data were analyzed offline using Labchart 8 software 138 (ADInstruments). EMG signals were amplified (gain = 500) using an octal bio-amplifier 139 (ML138, AD Instruments) with a bandwidth frequency range of 10-500 Hz obtained through 140 a band-pass filter. Prior to analysis, data were rectified and smoothed using a moving root-141 mean-square (RMS) with a time constant interval of 50 ms [8]. 142

143 *2.3.Experimental protocol*

All the participants were clearly informed on the modality of the test, and an experimenter provided practical demonstration if required. Participants performed a standardized warm-up consisting in 10 isometric contractions of the knee extensors at increasing intensities until reaching 90% of their perceived F_{max} . Warm-up was followed by familiarization with the experimental procedures: participants were instructed to extend their knee (in isometric condition) "as fast as possible" in order to assess the RFD variables [4]. Familiarization 150 consisted in 5 to 10 non-successive impulsive contractions, allowing participant to feel comfortable with the procedures. Successively, 5-min of rest were allowed before asking 151 participants to perform two 5-s MVC separated by at least 60 s of rest. If the difference 152 between the two MVCs was greater than 5%, participants performed a third MVC. In order to 153 perform MVC, participants were asked to contract "as hard as possible", strong verbal 154 encouragements and visual feedback were provided. After a 3-min rest, participants 155 performed a series of eight isometric contractions (0.5-1.5 s; Figure 1B) separated by 30 s of 156 rest. Strong verbal encouragement was provided. In case of countermovement, i.e. quick 157 158 flexion prior to the knee extension, determined from a force drop of 2 N below the baseline, or if the force attained was less than 70% of the F_{max} [7,25], the trial was repeated again in 159 order to obtain eight reliable contractions. 160

161 *2.4.Data analysis*

All data were analyzed offline using Labchart 8 Software (ADInstruments). The F_{max} and 162 associated RMS-EMG signal (EMG_{max}) were defined as the average of the highest 0.5-s 163 window among the best MVC. Repeatability of MVC was tested using a bivariate correlation. 164 The maximal rate of force development (RFD_{max}) was determined as the maximal value of 165 the force-time derivative curve for each RFD contraction. Absolute RFD_{max} was used to 166 identify the best five impulsive contractions retained and then averaged for further analysis, 167 while the others were discarded (Figure 1B). To assess the influence of within-subject 168 169 variability among young and very old adults, the coefficient of variation (CV) was calculated for each participant and compared between groups. 170

Time to RFD_{max} (tRFD_{max}) and force level at 25, 50, 100 and 200 ms from the onset of the force (see below) were identified. Then, the mean RFD was calculated at three time intervals of 0-50 ms (RFD0-50), 50-100 ms (RFD50-100), 100-200 ms (RFD100-200) and 0-200 ms (RFD 0-200) from the linear slope of the force-time curve [8] (Figure 1C). The onset of the force development was defined automatically as the point at which the level of force exceeded three standard deviations over the average resting baseline [25]. This method has been chosen because it provides a valid and reliable method to determine the onset of the contraction when low-noise dynamometers (~1 N) are used [4]. All RFD parameters are presented in both absolute values and expressed as percentage of F_{max} .

180 The rate of EMG rise (RER) was calculated as the linear slope of the RMS EMG-time curve

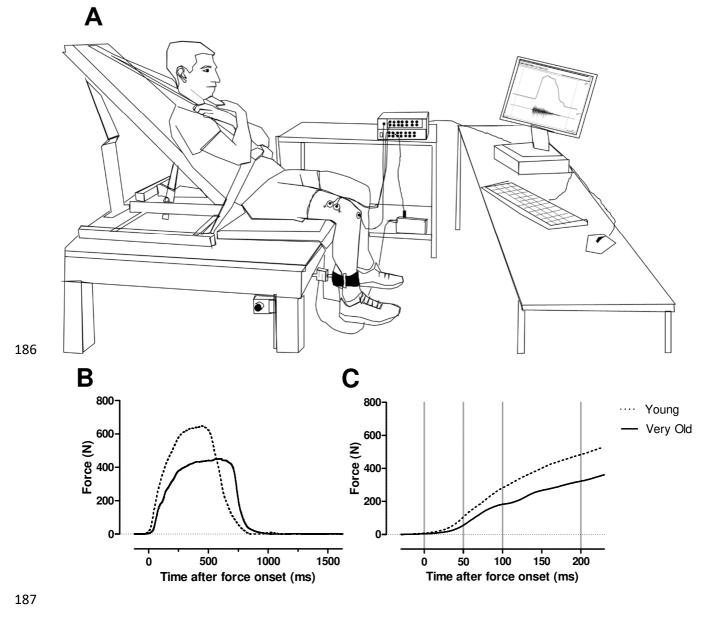
181 from two different onsets: the first onset was set 70 ms prior to the force onset as suggested

by Aagaard et al. [8], and the second onset was visually identified from the rise in the RMS

183 EMG-time curve. RER was analyzed at time intervals of 0-30 ms (RER30), 0-50 ms (RER50)

and 0-75ms (RER75). All EMG signals were normalized to EMG_{max} [10]. Electromechanical

delay was measured from the RMS EMG onset visually identified to the force signal onset.



188

189 *(2-column fitting image)*

Figure 1. Example of the force recording during the isometric impulsive contraction for tworepresentative participants. A: Schematic of the experimental setup.

192 B: Rapid force development of the knee extensors between young (dotted line) and very old

193 men (solid line). C: Rapid force development at different time intervals: 0-50 ms, 50-100 ms,

194 100-200 ms and 0-200 ms.

195

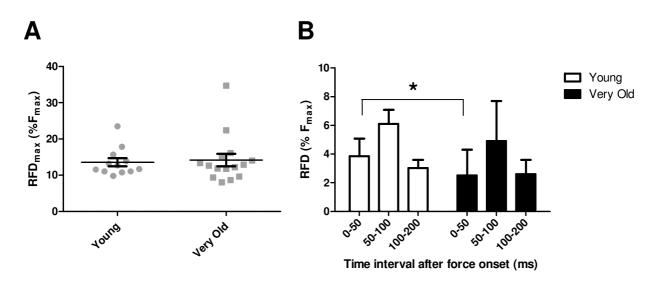
196 *2.5.Statistical analysis*

Data are reported as mean \pm SD in the text, the table and the figures. Assumption of 197 normality (Shapiro-Wilk test) was tested for all variables and outliers were determined using 198 a step of 1.5 x Interquartile range from the Turkey's Hinges. Intraclass correlation coefficient 199 (ICC, Two-way mixed model), standard error of the measurement (SEM), average CV and 200 repeated-measure analysis of variance (ANOVA) were used to assess reliability of RFD 201 measures [26] on both groups pooled together and separated. Independent samples T-tests 202 were used to compare F_{max}, EMG_{max}, and CV between groups. Mann-Withney U Tests were 203 204 used to compare time to RFD_{max}, absolute and relative RFD_{max}. Two-way ANOVAs were performed with age as between-subjects factor and time as within-subjects factor for all the 205 remaining force and EMG-derived variables. Sphericity (Mauchly test) were tested for all 206 207 dependent variables. If the assumption of sphericity was violated, the correction of Greenhouse-Geisser was applied. When significant main effects were observed, Bonferroni's 208 test was used for post-hoc analysis. Significance was set as $P \le .05$ for all tests. Statistical 209 analysis was conducted using IBM SPSS software (Version 23.0, SPSS, Inc, Chicago, IL). 210

3. Results

Due to technical issues, the EMG signals for two very old men were unusable and thus 212 discarded. Thus, EMG parameters are presented for 13 very old men. Absolute values for 213 Fmax, EMGmax, RFD, RFDmax and tRFDmax are presented in Table 1. Reliability of RFD 214 measurement was excellent either when both groups were pooled together (ICC=0.98, 215 CI=0.97-0.99; SEM=893.87 N*s⁻¹; CV=11±6%; ANOVA: $F_{(4,104)}=0.84$, P=.50) and when the 216 young (ICC=0.96, CI=0.91-0.98; SEM=998.78 N*s⁻¹; CV=12±6%; ANOVA: $F_{(4,44)}=2.10$, 217 P=.10) and very old men (ICC=0.98, CI=0.96-0.99; SEM=810.80 N*s⁻¹; CV=9±6%; 218 ANOVA: $F_{(4,56)}=0.27$, P=.90) were analyzed separately. F_{max} was consistent between 219 measures (r = 0.98, P<0.001) and significantly lower (37%) for the very old men compared to 220

the young men (P < 0.001). The associated EMG_{max} was also lower for the older adults (27%, *P*=0.42). Absolute RFD_{max} was greater in the young men (35%, P < 0.001), however the agerelated difference was not observed when normalized to the maximal force (Figure 2A, *P*=0.78). Normalized RFD was not influenced by the between-subject variability, as CV was similar between young (9±6%) and very old men (12±6%, P=0.21).



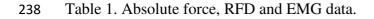
226

227 (2-column fitting image)

Figure 2. Normalized RFD performance of young *vs* very old adults. A: Differences in the RFD_{max} normalized to the F_{max} . B: The ratio of the RFD and F_{max} at three time intervals of 0-50 ms, 50-100 ms and 100-200 ms after the force onset between young men (white) and very old men (black). *=P<0.05.

It can be noticed on Figure 2A that two outliers from the very old group may have influenced the mean. Nevertheless, the statistical analysis was also performed without these two participants and the results remained unchanged. Finally, no age-related differences were observed for the tRFD_{max} (P=0.25).

Of those two outliers, one very old man showed higher F_{max} (554 N) and RFD (12409 N*s⁻¹) than some young men.



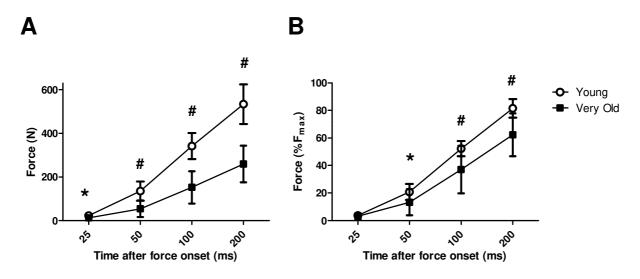
	Young Men	Range (min. – max.)	Very Old Men	Range (min. – max.)
F _{max} (N)	660.9±139.1	455.0 - 911.0	414.3±90.8 #	268.0 - 554.0
EMG _{max} (mV)	0.62±0.22	0.24 - 1.09	0.45±0.16 *	0.26 - 0.80
$RFD_{max} (N*s^{-1})$	8719.7±2055.9	6278.0 - 11625.0	5700.5±2473.9 #	$3262.0 - 12409.0^{a}$
$tRFD_{max}(s)$	70.3±36.1	40.9 - 168.5	85.8±33.1	45.0 - 127.0
RFD 0-50 ms (N*s ⁻¹)	2494.7±877.7	$1543.0 - 4684.0^{a}$	1034.4±738.4 #	222.0 - 2470.0
RFD 50-100 ms (N*s ⁻¹)	3992.7±929.8	2885.0 - 6033.0	2040.2±1258.7 #	568.0 - 4035.0
RFD 100-200 ms (N*s ⁻¹)	1986.6±513	$1283.0 - 3071.0^{a}$	1101.6±544.2 #	379.0 - 1925.0
RFD 0-200 ms (N*s ⁻¹)	2725.2±410.8	2097.7 - 3441.85	1470.5±503.3 #	618.38 - 2241.95

239

240 F_{max} = maximal force; EMG_{max}: maximal EMG; RFD= rate of force development; 241 RFD_{max}=maximal rate of force development; tRFD_{max}= time to the maximal rate of force 242 development. ^a= from an outlier; *= P < 0.05; #= P < 0.01.

243

From the onset of the contraction, there were significant age-related differences in the 244 absolute force level at every time point and were 23.1±8.5 N vs 12.6±10.5 N (25-ms, 245 P=0.01), 135.2±43.8 N vs 53.8±38.4 N (50-ms, P<0.001), 340.7±59.6 N vs 152.4±73.5 N 246 (100-ms, P<0.001) and 533.3±91.1 N vs 258.9±84.2 N (200-ms, P<0.001) for young and 247 very old men, respectively (Figure 3A). When normalized to F_{max}, the differences between 248 groups were less pronounced. Normalized force values were 3.6±1.3% vs 3.1± 2.3% (25-ms, 249 250 P=0.49), 20.7±5.8% vs 13.3±9.3% (50-ms, P=0.02), 52.1±5.5% vs 36.9±17.3% (100-ms, P=0.007) and 81.5±6.7% vs 62.2±15.6% (200-ms, P=0.001) for young vs very old men, 251 respectively (Figure 3B). 252



253

254 (2-column fitting image)

Figure 3. Differences between young men (empty circles) and very old men (black squares) at four time points after the force onset. A: Difference in absolute force level. B: Difference in force level normalized to the F_{max} . *=P<0.05; #=P<0.01.

258

Average RFD were greater for the young than the very old men at every time interval (Table 1). When normalized to the F_{max} , average RFD were $3.8\pm1.2\%$ (0-50 ms), $6.1\pm0.9\%$ (50-100 ms) and $3.0\pm0.5\%$ (100-200 ms) for young men and $2.5\pm1.7\%$ (0-50 ms), $4.8\pm2.7\%$ (50-100 ms) and $2.5\pm1.0\%$ (100-200 ms) for very old men (Figure 2B). Significance was found only at the 0-50 ms time interval (*P*=0.04), while no differences were observed at 50-100 ms (*P*=0.16) and 100-200 ms (*P*=0.20). Finally, significant difference was found at the 0-200 ms time interval between young ($4.2\pm0.7\%$) and very old men ($3.5\pm0.9\%$; *P*=0.04).

RER of the VL, expressed as %EMGmax, exhibited different results depending on the onset considered (Figure 4). Data obtained with the RER onset 70 ms prior to the force onset showed significantly lower RER during the 0-30 ms time interval in the young men (P=0.04), while no age-related differences were observed during the 0-50 ms (P=0.26) and 0-75 ms time intervals (P=0.36; Figure 4A). The opposite trend was observed when RER onset was visually set at the point of RMS EMG rise, RER was greater for the young than the very old 272 men for every time interval (RER30: P=0.003; RER50: P<0.001; RER75: P=0.007; Figure 273 4B). Electromechanical delay was longer in the very old men (76.8±27.5 ms) than in the 274 young men (37.4±0.9 ms; P<0.001).

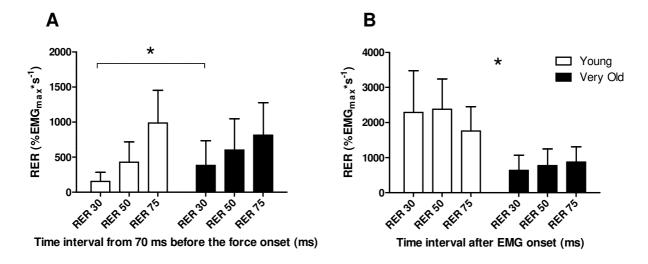




Figure 4. Comparison between young (white) and very old (black) men in the RER normalized to the maximal EMG at three time intervals of 0-30 ms, 0-50 ms and 0-75ms. A: onset was automatically set 70 ms prior to the force onset. B: onset was visually set at the rise in RMS EMG-time curve. *=P<0.05.

280

281 **4. Discussion**

The aim of the present study was to assess the age-related impairments in rapid force 282 production capacity and rate of muscle activation of the KE. We reported reduced F_{max} and 283 absolute RFD_{max} in active octogenarians compared to young men. For every time interval of 284 285 the force-time curve, absolute force and RFD values were considerably lower in very old men, as initially hypothesized. Contrary to our hypothesis, no age-related difference in 286 RFD_{max} was observed when normalized to F_{max}. However, normalized RFD values in the 287 early phase of the contractions (0-50 ms) were lower in very old compared to young men. 288 This age-related difference disappeared in the successive phases of the contraction. 289

Regarding RER, when the onset was defined manually on the RMS EMG-time curve, RERwas significantly lower for very old men at every time intervals.

Although investigating the capacities of the KE is functionally relevant in old and very old 292 293 age [19,27], the literature regarding RFD performance with aging for this muscle group is limited. Thompson and colleagues [1] observed ~28% lower F_{max} and absolute RFD_{max} 294 measured during isometric KE MVC in old (67 yr) compared to young adults (25 yr), while 295 Mau Moeller et al. [12] reported even greater age-related impairments (F_{max}, ~40%; RFD_{max}, 296 ~65%) between young (25 yr) and old (70 yr) men during the same task. In agreement with 297 298 the previous studies, we reported a $\sim 36\%$ lower absolute F_{max} and RFD_{max} in very old men compared to young men. During the rapid isometric contractions, the absolute force level of 299 very old men was significantly lower than the young men at every time point, and this 300 301 difference persisted when normalized to F_{max} . These results confirm the marked impairment 302 in the capacity to rapidly increase the level of force with very old age that accompany the decline in maximal force production capacity [20]. 303

304 Interestingly, no age-related differences were observed in RFD_{max} when normalized to F_{max}, which could indicate similar factors are responsible for the decline of those two parameters. 305 Maximal force and rapid force development are both affected by sarcopenia and the 306 consequent muscle remodeling process with aging [28]. Only one very old man was more 307 performant in both F_{max} and RFD_{max} than some young men. This is in agreement with 308 309 previous investigations showing no age-related change in normalized RFD in plantar flexors [15], dorsiflexors [15], leg extensors [1] and leg flexors [1]. However, other studies reported 310 a reduction in the relative RFD_{max} of plantar flexors [10], dorsiflexors [2] and leg extensors 311 312 [12,14]. In contrast with the previous investigations, in the present study F_{max} and RFD were obtained during different contractions and giving specific instructions, according to the most 313 recent recommendations [4,5], obtaining good within-subject reliability of the RFD measures 314

315 in both groups. Consequently, different muscles involved, data analysis and protocols adopted may partially explain the discrepancies in the literature. The majority of the studies 316 analyzed RFD during MVC, while we used specific impulsive isometric contractions. As we 317 318 asked our participants to push "as hard as possible" during MVC rather than "hard and fast", we could not compare RFD during MVC and the impulsive contractions. Other than 319 methodological factors, the high level of physical activity of our participants may influenced 320 age-related differences in RFD. In the present study, the very old participants were healthy 321 community-dwelling volunteers, and classified as "active" according to the number of 322 323 steps/day measured (threshold defined at 7000 steps/day between active and sedentary very old men [29]). These healthy individuals may not represent the overall population over 80 324 years, but rather those with greater lifespan physical activity behaviors and/or with biological 325 326 superiority [30]. The natural selection bias of healthy participants over 80 years (alive, 327 autonomous, without motor or neural impairments, etc...) could also explain why the agerelated declines in performance observed in previous cross-sectional and longitudinal studies 328 329 were less important than what expected in the ninth decade from linear extrapolation models [31,32]. Moreover, physical activity has important implications in the age-related changes in 330 RFD. For example, Aagaard et al. [3] showed that chronic exposure to strength training 331 preserves RFD_{max} at a similar level in young and old individuals, and Caserotti et al. [33] 332 reported similar performances after 12-weeks of heavy-resistance training in old (60-65 333 334 years) and very old women (80-89 years). Further investigations should compare normalized RFD of very old men with different levels of physical activity to clarify the relative 335 contribution of maximal force decline, aging and physical activity level in RFD performance. 336 337 The normalized 0-200 ms time interval was used to provide a more representative indicator of the age-related RFD changes, and was significantly lower in the very old men. This reflects 338 an overall impairment in the capacity of rapid force generation with aging that has been 339

evidenced at every time point on the force-time curve. To better illustrate RFD differences
between groups, different time epochs were analyzed and normalized to F_{max}.

The average normalized RFD values were significantly lower only in the early phase of the 342 343 contraction (0-50 ms), but not in the successive time intervals (50-100 ms and 100-200 ms), contrary to the absolute RFD values and the normalized force values. The non-significant 344 difference in the middle time epochs regarding RFD data could be explained by the higher 345 between-subject variability in very old men. Despite higher variability, the difference at 0-50 346 ms was great enough to reach statistical significance. In other words, the force increment in 347 348 this short time period (50 ms) was greater for the young men, which determined a difference in the relative RFD between groups. This could suggest that different neural strategies could 349 be adopted during aging to compensate the progressive loss in RFD capacity, still evident in 350 351 the 0-50 ms time interval. This could be partially supported by our results on time to reach RFD_{max}, which was similar between groups. Similar results were obtained by Thompson et al. 352 [10] and Mau-Moeller et al. [12]. Thompson et al. [10] found a difference between young and 353 354 old adults in the plantar flexors at time intervals of 0-50 ms and 100-200 ms. However, these authors didn't report the time interval 50-100 ms. Thus, it seems that the early phase of the 355 contraction is determinant for the age-related impairment in absolute RFD capacity of the leg 356 extensors. Previous investigations associated the first phase of the contraction (0-50 ms) to 357 the rate of muscle activation. For instance, a recent study of Gerstner et al. [34] on the plantar 358 359 flexors found similar rate of torque development capacity between young and old men (~69 yr) in the firsts 50 ms of contraction, while it was reduced in the late interval (100-200 ms). 360 These authors suggested that while factors like poorer muscle quality, the inability to sustain 361 362 a high rate of muscle activation and reduced level of physical activity could influence the late interval of contraction in old men, muscle activation could reflect changes in the early time 363 interval. To provide further insight, we quantified using the rate of EMG rise (RER). We set 364

two different RMS EMG onset for the RER analysis. With the EMG onset fixed 70 ms prior 365 to force onset [8], RER was significantly higher in very old compared to young men only in 366 the early phase of the contraction (0-30ms), while no age-related differences were observed at 367 368 0-50 ms and 0-75ms. However, the electromechanical delay was significantly different between groups, and only the electromechanical delay of the very old men was close to the 369 fixed value of 70 ms suggested by Aagaard et al. [8]. Thus, setting the onset visually on the 370 RMS EMG-time curve seems to be the best option when two different populations are 371 compared. In the present study, when the RMS EMG onset was set manually, RER was 372 significantly lower for very old men for all the time intervals considered. The longer 373 electromechanical delay in the very old group compared to the young, despite in small part 374 dependent from the electrode placement, indicate a deficit in the transmission and conversion 375 376 of the voluntary drive to force. The reduced capacity to rapidly activate the muscle suggests that impairments occur also in the voluntary drive, e.g. the generation of doublet discharge, 377 the decrease in discharge frequency and/or the inability to maintain high discharge 378 frequencies, leading to different or less efficient activation pattern with aging [2,35]. To 379 interpret those results, it has been shown that the age-related slowing of muscle contractile 380 kinetics, the progressive reduction in type II muscle fibers [19] and changes in the muscle 381 architecture [34] are muscle dependent and could influence the EMG signal. Limiting our 382 analysis to the VL and recognizing the limitations of surface EMG measurements [36], 383 384 further evidences are needed to clarify the specific role of neural vs contractile mechanisms involved in the loss of RFD across different muscles. 385

386

5. Conclusions

388 The present study demonstrated that rapid force development capacity is impaired in active 389 octogenarians. However, the age-related difference is minimized when RFD is normalized by the maximal force of each individual. The results suggest that muscle force and RFD have common mechanisms determining neuromuscular changes associated with healthy aging. The impairments in RFD are accentuated in the early phase of the contraction, which has been associated in the literature with the rate of muscle activation. Further studies are needed to establish whether similar levels of Fmax or RFDmax are related to similar motor function in very old age.

396

397 Conflicts of interest

398 No conflicts of interests, financial or otherwise are declared by the authors.

399 Funding

400 This research did not receive any specific grant from funding agencies in the public,401 commercial, or not-for-profit sectors.

402

403 Reference	5
----------------------	---

404

405	1.	Thompson	n BJ.	Ryan ED	, Sobolewski EJ,	Conchola EC.	Cramer JT. Age relat	ed

differences in maximal and rapid torque characteristics of the leg extensors and flexors

in young, middle-aged and old men. Exp Gerontol 2013; 48: 277–282.

2. Klass M, Baudry S, Duchateau J. Age-related decline in rate of torque development is

409 accompanied by lower maximal motor unit discharge frequency during fast

410 contractions. J Appl Physiol 2008; 104: 739–746.

411 3. Aagaard P, Magnusson PS, Larsson B, Kjaer M, Krustrup P. Mechanical muscle

412 function, morphology, and fiber type in lifelong trained elderly. Med Sci Sports Exerc
413 2007; 39: 1989–96.

414 4. Maffiuletti NA, Aagaard P, Blazevich AJ, Folland J, Tillin N, Duchateau J. Rate of

415	force development: physiological and methodological considerations. Eur J App
416	Physiol 2016; 116: 1091–1116.

- 417 5. Rodríguez-Rosell D, Pareja-Blanco F, Aagaard P, González-Badillo JJ. Physiological
 418 and methodological aspects of rate of force development assessment in human skeletal
 419 muscle. Clin Physiol Funct Imaging 2017; 1–20.
- 420 6. Andersen LL, Aagaard P. Influence of maximal muscle strength and intrinsic muscle
 421 contractile properties on contractile rate of force development. Eur J Appl Physiol
 422 2006; 96: 46–52.
- 423 7. Folland JP, Buckthorpe MW, Hannah R. Human capacity for explosive force
- 424 production: neural and contractile determinants. Scand J Med Sci Sports 2014; 24:
 425 894–906.
- 426 8. Aagaard P, Simonsen EB, Andersen JL, Magnusson P, Dyhre-Poulsen P. Increased
 427 rate of force development and neural drive of human skeletal muscle following
 428 resistance training. J Appl Physiol 2002; 93: 1318–1326.
- 429 9. Clark DJ, Patten C, Reid KF, Carabello RJ, Phillips EM, Fielding RA. Muscle
- 430 performance and physical function are associated with voluntary rate of neuromuscular
- 431 activation in older adults. Journals Gerontol Ser A Biol Sci Med Sci 2011; 66 A:
- 432 115–121.
- 433 10. Thompson BJ, Ryan ED, Herda TJ, Costa PB, Herda AA, Cramer JT. Age-related
- changes in the rate of muscle activation and rapid force characteristics. Age (Omaha)
 2014; 36: 839–849.
- Izquierdo M, Ibanez J, Gorostiaga E *et al*. Maximal strength and power characteristics
 in isometric and dynamic actions of the upper and lower extremities in middle-aged
 and older men. Acta Physiol Scand 1999; 167: 57–68.
- 439 12. Mau-Moeller A, Behrens M, Lindner T, Bader R, Bruhn S. Age-related changes in

- 440 neuromuscular function of the quadriceps muscle in physically active adults. J
- 441 Electromyogr Kinesiol 2013; 23: 640–648.
- 442 13. Bellumori M, Jaric S, Knight C a. Age-related decline in the rate of force development
 443 scaling factor. Motor Control 2013; 17: 370–81.
- **14**. Korhonen MT, Cristea A, Alén M *et al.* Aging, muscle fiber type, and contractile
- function in sprint-trained athletes. J Appl Physiol 2006; 101: 906–917.
- Thelen DG, Schultz AB, Alexander NB, Ashton-Miller JA. Effects of Age on Rapid
 Ankle Torque Development. J Gerontol 1996; 51A: 226–232.
- 448 16. Hepple RT, Rice CL. Innervation and neuromuscular control in ageing skeletal muscle.
 449 J Physiol 2016; 594: 1965–1978.
- 450 17. Sahaly R, Vandewalle H, Driss T, Monod H. Maximal voluntary force and rate of
- 451 force development in humans Importance of instruction. Eur J Appl Physiol 2001;
 452 85: 345–350.
- 453 18. Hunter SK, Pereira HM, Keenan KG. The aging neuromuscular system and motor
 454 performance. J Appl Physiol 2016; 121: 982–995.
- 455 19. Mitchell WK, Williams J, Atherton P, Larvin M, Lund J, Narici M. Sarcopenia,
- 456 dynapenia, and the impact of advancing age on human skeletal muscle size and
- 457 strength; a quantitative review. Front Physiol 2012; 3 JUL: 1–18.
- 458 20. Venturelli M, Reggiani C, Richardson RS, Schena F. Skeletal Muscle Function in the
- 459 Oldest-Old: The Role of Intrinsic and Extrinsic Factors. Exerc Sport Sci Rev 2018; 46:
 460 188–194.
- 461 **21**. Frontera WR, Hughes VA, Fielding RA, Fiatarone MA, Evans WJ, Roubenoff R.
- 462 Aging of skeletal muscle: a 12-yr longitudinal study. J Appl Physiol 2000; 88: 1321–
 463 1326.
- 464 22. McGregor RA, Cameron-Smith D, Poppitt SD. It is not just muscle mass: A review of

muscle quality, composition and metabolism during ageing as determinants of muscle 465 function and mobility in later life. Longev Heal 2014; 3: 1-8. 466 23. Barthélémy JC, Pichot V, Dauphinot V et al. Autonomic nervous system activity and 467 decline as prognostic indicators of cardiovascular and cerebrovascular events: The 468 'PROOF' study: Study design and population sample. Associations with sleep-related 469 breathing disorders: The 'SYNAPSE' study. Neuroepidemiology 2007; 29: 18-28. 470 Tudor-Locke C, Williams JE, Reis JP, Pluto D. Utility of Pedometers for Assessing 471 24. Physical Activity. Sport Med 2006; 32: 795-808. 472 473 25. De Ruiter CJ, Kooistra RD, Paalman MI, de Haan A. Initial phase of maximal voluntary and electrically stimulated knee extension torque development at different 474 knee angles. J Appl Physiol 2004; 97: 1693–1701. 475 476 **26**. Weir JP. Quantifying Test -Retest Reliability Using the Intraclass Correlation 477 Coefficient and the SEM. J Strength Cond Res 2005; 19: 231–240. 27. Laukkanen P, Heikkinen E, Kauppinen M. Muscle strength and mobility as predictors 478 479 of survival in 75-84-year-old people. Age Ageing 1995; 24: 468-73. Gueugneau M, Coudy-Gandilhon C, Théron L et al. Skeletal Muscle Lipid Content **28**. 480 and Oxidative Activity in Relation to Muscle Fiber Type in Aging and Metabolic 481 Syndrome. Journals Gerontol - Ser A Biol Sci Med Sci 2015; 70: 566–576. 482 29. Tudor-locke CE, Craig CL, Brown WJ et al. How Many Steps / day are Enough ? For 483 484 Older Adults and Special Population. Int J Behav Nutr Phys Act 2011 2011; 1–17. 30. Porter MM, Vandervoort AA, Lexell J. Aging of human muscle : structure , function 485 and adaptability. Scand J Med Sci Sport 1995; 5: 129-142. 486 487 31. Aniansson A, Hedberg M, Henning G-B, Grimby G. Muscle morphology, enzymatic activity, and muscle strength in elderly men: A follow-up study. Muscle Nerve 1986; 488 9: 585–591. 489

490	32.	Greig CA, Botella J, Young A. The quadriceps strength of healthy elderly people
491		remeasured after eight years. Muscle Nerve 1993; 16: 6–10.
492	33.	Caserotti P, Aagaard P, Buttrup Larsen J, Puggaard L. Explosive heavy-resistance
493		training in old and very old adults: Changes in rapid muscle force, strength and power.
494		Scand J Med Sci Sport 2008; 18: 773–782.
495	34.	Gerstner GR, Thompson BJ, Rosenberg JG, Sobolewski EJ, Scharville MJ, Ryan ED.
496		Neural and Muscular Contributions to the Age-Related Reductions in Rapid Strength.
497		Med Sci Sports Exerc 2017; 49: 1331–1339.
498	35.	Mitchell C, Cohen R, Dotan R, Gabriel D, Klentrou P, Falk B. Rate of Muscle
499		Activation in Power- and Endurance-Trained Boys. Int J Sport Physiol Perform 2011;
500		6: 94–105.
501	36.	Farina D, Merletti R, Enoka RM. Neural Control of Movement: The Extraction of
502		Neural Strategies from the Surface {EMG}. J Appl Physiol 2004; 96: 1486–1495.
503		