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Anticipatory Postural Adjustments and Spatial Organization of Motor Cortex: Evidence of Adaptive Compensations in Healthy Older Adults

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1 **Anticipatory Postural Adjustments and Spatial Organization of Motor Cortex – Evidence**
2 **of Adaptive Compensations in Healthy Older Adults**

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16

17 **Author contributions:**

18 JS and BF designed the study. JS performed data collection, processing and analyses. JS and
19 BF interpreted the data. JS drafted the manuscript and BF provided critical review of the
20 manuscript.

21

22 **Running head:**

23 APAs and M1 organization in older adults

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30 **ABSTRACT**

31 During anticipated postural perturbations induced by limb movement, the central nervous
32 system generates anticipatory postural adjustments (APAs) in the trunk and hip musculature to
33 minimize disturbances to equilibrium. Age-related changes in functional organization of the
34 nervous system may contribute to changes in APAs in healthy older adults. Here we examined if
35 altered APAs of trunk/hip musculature in older adults are accompanied by changes in the
36 representation of these muscles in motor cortex. 12 healthy older adults, 5 with a history of falls
37 and 7 non-fallers, were compared to 13 young adults. APAs were assessed during a
38 mediolateral arm raise task in standing. Temporal organization of postural adjustments was
39 quantified as latency of APAs in the contralateral external oblique, lumbar paraspinals and
40 gluteus medius relative to activation of the deltoid. Spatial organization was quantified as extent
41 of synergistic coactivation between muscles. Volume and location of the muscle representations
42 in motor cortex were mapped using transcranial magnetic stimulation. We found that older
43 adults demonstrated significantly delayed APAs in the gluteus medius muscle. Spatial
44 organization of the three muscles in motor cortex differed between groups, with the older adults
45 demonstrating more lateral external oblique representation than the other two muscles.
46 Separate comparisons of the faller and non-faller subgroups with young adults indicated that
47 non-fallers had the greatest delay in gluteus medius APAs and a reduced distance between the
48 representational areas of the lumbar paraspinals and gluteus medius. This study indicates that
49 altered spatial organization of motor cortex accompanies altered temporal organization of APA
50 synergies in older adults.

51 **KEYWORDS**

52 Motor cortex; transcranial magnetic stimulation; torso; functional organization; aging; postural
53 control

54 **NEW AND NOTEWORTHY**

55 Anticipatory postural adjustments (APAs) are a critical component of postural control. Here we
56 demonstrate that in healthy older adults with and without a history of falls, delayed APAs in the
57 hip musculature during mediolateral perturbations are accompanied by altered organization of
58 trunk/hip muscle representation in motor cortex. The largest adaptations are evident in older
59 adults with no history of falls.

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1. INTRODUCTION

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66 Falls are a significant cause of morbidity and mortality among older adults. Although falls have
67 multiple causes, changes in postural control in older adults contribute significantly to fall risk
68 (Rubenstein and Josephson 2006). It is becoming clear that adaptations in structure and
69 function occur at every level of the postural control system in association with aging (Papegaaij
70 et al. 2014a). In order to design effective exercise interventions to reduce the risk of falls, it is
71 critical to understand how nervous system adaptations may contribute to age-related changes in
72 postural control in healthy older adults.

73 Anticipatory postural adjustments (APAs) are an important component of postural control (Horak
74 2006). Anticipatory postural adjustments are synergies of feedforward muscle activation or
75 inhibition that occur before a predictable perturbation. Disordered APAs may result in postural
76 instability during self-initiated movements (Horak 2006; Kubicki et al. 2012). APA synergies can
77 be characterized in terms of the timing of muscle activation or inhibition relative to the
78 destabilizing event (temporal organization); in terms of the three-dimensional coordination of
79 activity in multiple muscles (spatial organization); and in terms of the magnitude of muscle
80 activation (amplitude scaling). The standing rapid arm flexion task is a simple paradigm that is
81 often used to quantify these characteristics of APAs. Anticipatory postural control of the trunk
82 and hip musculature during rapid arm raising in standing counteracts reactive forces from upper
83 limb motion and helps to maintain the mass of the head and trunk within the base of support.
84 During rapid arm flexion, APAs occur in the abdominals, paraspinals and hip extensors in
85 healthy young adults (Hodges et al. 1999; Massé-Alarie et al. 2012). In older adults, APAs in the
86 hip extensors are delayed relative to the onset of the agonist (deltoid) muscle compared to
87 young adults (Rogers et al. 1992). In addition to this altered temporal organization, older adult
88 have altered spatial organization of postural control with increased coactivation of lower limb

89 muscles during standing and reaching (Nagai et al. 2011). It is not known if this coactivation is
90 evident in the trunk and hip musculature during rapid arm raising.

91 Much of the research investigating APAs has utilized perturbations that are induced in the
92 anterior-posterior direction, such as rapid arm flexion. However, postural control in the
93 mediolateral plane is critical to maintaining dynamic stability (Rogers and Mille 2003), and
94 disordered mediolateral postural control is associated with a history of falls (Maki et al. 1994).
95 Research investigating externally-induced mediolateral postural perturbations has demonstrated
96 synergistic APAs in the gluteus medius, external oblique, and paraspinal musculature in healthy
97 young adults (Santos and Aruin 2008). Evidence from the same perturbations suggests that
98 there is no change in the magnitude of trunk and hip APAs in older adults (Claudino et al. 2013).
99 It is still unclear if the temporal and spatial organization of mediolateral APA synergies in the
100 trunk and hip musculature are affected by aging.

101 Neural substrates of postural control are distributed throughout the central nervous system. The
102 structure and function of these substrates is affected by healthy aging. In primary motor cortex,
103 intracortical inhibition during standing is reduced in older adults compared with younger adults,
104 and the extent of this reduction in inhibition is associated with worse postural performance
105 (Papegaaij et al. 2014b). As the motor cortex contributes to preparation of postural adjustments
106 (Tsao et al. 2008; Jacobs et al. 2009a; Chiou et al. 2016, 2018), age-related changes in motor
107 cortex may also be associated with changes in APAs in older adults. In particular, excitability of
108 cortical neural networks is modulated in response to use and with healthy aging. This has been
109 demonstrated by changes in the topographic organization of muscle-specific corticospinal
110 output evoked by transcranial magnetic stimulation (TMS)(Adkins et al. 2005; Plow et al. 2014;
111 Masse-Alarie et al. 2017). TMS studies mapping motor cortical organization during voluntary
112 motor tasks show that older adults demonstrate less distinct topographic representation of
113 muscles, reduced representational volume (Coppi et al. 2014) and shifted representational area

114 (Bernard and Seidler 2012). Therefore, less differentiated and shifted representations of the
115 postural musculature in M1 may underlie the impairments in APAs that are evident in older
116 adults.

117 The primary purpose of this study was to compare temporal and spatial characteristics of
118 anticipatory postural adjustments of the trunk and hip, and the motor cortical representation of
119 trunk and hip musculature, in young adults and healthy older adults. A secondary purpose of
120 this study was to explore if these variables differ in older adults with and without a history of
121 falls. We hypothesized that latency of APAs would be delayed in older adults and that
122 coactivation between muscles would be greater, and that this would be accompanied by
123 reduced differentiation of the trunk and hip musculature motor cortical representation. We
124 further hypothesized that these changes would be more evident in older adults with a history of
125 falls than those with no fall history.

126 **2. METHODS**

127 **2.1 Participants**

128 The study was approved by the Institutional Review Body of the University of Southern
129 California and all participants gave written informed consent before enrollment and data
130 collection. Participants were recruited from the local community. Participants in the older adult
131 group were over 65 years, community-dwelling, independent with activities of daily living and
132 ambulation, able to stand upright without assistance for two minutes and able to follow verbal
133 directions (Newton 2001). A history of falls was determined with a questionnaire (Claudino et al.
134 2013), with a fall defined as an unplanned contact with a support surface below knee level
135 (Takahashi et al. 2006). Fallers were defined as those who had experienced at least one fall in
136 the past year (Hass et al. 2004). Participants in the young adult group were between 18 and 30

137 years old (Isles et al. 2004). Exclusion factors in both groups were a history of disorders
138 affecting balance, significant/persistent low back pain, vestibular disorders, and inability to
139 abduct both arms to at least 90°. As per current TMS recommendations, participants were also
140 excluded if they had metal, electrical or magnetic implants, a personal or family history of
141 epilepsy, or other medical history/use of medications or substances that are known to lower
142 seizure threshold (Rossi et al. 2011).

143 **2.2 Experimental procedure**

144 Balance and mobility were assessed in older adults with the Anticipatory Postural Adjustments
145 section of the BESTest (Horak et al. 2009) and the Timed Up and Go test. Self-selected gait
146 velocity in older adults was calculated from the average of two 10m walking trials.

147 2.2.1 Mediolateral anticipatory postural adjustments

148 Bipolar, disposable surface electromyography electrodes (inter-electrode distance 22mm,
149 Myotronics-Noromed, Inc., Tukwila, USA) were placed on external oblique (EO), thoracic
150 longissimus pars lumborum at the level of L1 (LL) and gluteus medius (GMED) in accordance
151 with established guidelines (Hermens 2000). The electrodes were placed on the same side as
152 the dominant limb. Additionally, electrodes were placed on the deltoid muscles. EMG data were
153 transmitted and digitally sampled at 1500Hz using a wireless telemetry system (base gain 400;
154 TeleMyo DTS Telemetry, Noraxon USA Inc, Scottsdale, USA).

155 Anticipatory postural adjustments were quantified during a rapid arm raise task (Figure 1a). A
156 2lb weight was placed on the wrist of the limb contralateral to the trunk/hip EMG instrumentation
157 (i.e. left arm in an individual who identified their dominant limb as the right) (Horak et al. 1984).
158 As APAs are direction-specific, the contralateral side was selected for the arm raise task as
159 existing research and preliminary data suggested that this would maximize activity in two out of

160 the three muscles under investigation (Santos and Aruin 2008). The weight was used since
161 preliminary data indicated clearer and more consistent APAs in the trunk and hip musculature
162 with external loading. Participants stood barefoot with their feet parallel and heels 10cm apart.
163 In response to an auditory/visual cue, participants abducted the arm to 90° as rapidly as
164 possible. Six trials were collected (Tsao et al. 2010a). The time taken to reach 90° of
165 glenohumeral abduction was monitored utilizing a laser trigger system.

166 2.2.2 Motor cortical representation

167 Topographic organization of muscle representational areas in primary motor cortex were
168 quantified with motor evoked potentials from single-pulse TMS. TMS procedures were
169 conducted and are described here in accordance with current guidelines (Chipchase et al.
170 2012).

171 Motor evoked potentials (MEPs) were elicited using a single-pulse magnetic stimulator
172 (MagStim 200², Magstim Inc, NC) and a 110mm double cone coil (Magstim Inc, NC) (Lagan et
173 al. 2008; Tsao et al. 2008; Fisher et al. 2013). Stimulation was applied on the hemisphere
174 contralateral to the side of EMG instrumentation (i.e. on the left if the dominant limb was the
175 right). The previously described surface EMG electrodes on the external oblique, thoraco-
176 lumbar longissimus and gluteus medius were attached to a pre-amplifier (Motion Lab Systems,
177 15003 Hz, bandpass filter 1 - 1000 Hz, base gain 2000). MEPs were acquired and stored using
178 Signal software (Signal v6, Cambridge Electronic Design Ltd, Cambridge UK). A lycra cap
179 marked with a 1cm grid was placed over the participant's scalp and the location of the vertex
180 determined. To ensure correct and consistent coil placement the Brainsight® Frameless
181 stereotactic image guidance system was used (Rogue Research Inc, Montreal, Canada).
182 Landmarks on each participant's head were co-registered with the Brainsight™ system using an
183 infra-red marker tracking system. The position and orientation of the coil was then tracked

184 relative to the position of these markers and to a 3-D reconstruction of a standard brain MRI.

185 Prior to the TMS data collection, the maximal voluntary isometric contraction (MVIC) for each
186 muscle was determined. Manual resistance was provided to the participant against the lateral
187 border of the dominant limb as they performed hip abduction in side lying (gluteus medius) and
188 at the shoulders as they performed maximal trunk flexion/rotation in supine (external oblique).
189 Due to the small representational area of the trunk and hip musculature, MEPs are not
190 consistently elicited when the muscles are at rest, therefore motor thresholding and mapping
191 was performed during a submaximal contraction for all three muscles (Lagan et al. 2008; Tsao
192 et al. 2010a; Massé-Alarie et al. 2012)

193 *Lumbar longissimus/gluteus medius* - TMS mapping of the lumbar longissimus and gluteus
194 medius were conducted during double-leg bridging in supine (Fisher et al. 2013). Consistent
195 bridge height was ensured by having participants raise the pelvis up to the height of a reference
196 marker placed at a 150% of the vertical distance of their anterior superior iliac spines to the
197 table. Additional resistance to hip abduction was provided by a band placed around the distal
198 thighs (Figure 1c). Each TMS stimulus was delivered as the participant maintained the correct
199 test position and gluteus medius contraction at 20 % MVIC. A consistent level of muscle
200 activation was ensured by providing real-time visual feedback of the root mean square averaged
201 amplitude of the gluteus medius contraction relative to the 20% MVIC activation target.
202 Feedback was provided for amplitude of gluteus medius EMG activity only, as pilot data
203 indicated a consistent activation ratio of approximately 1.6: 1 for the longissimus and gluteus
204 medius during a double-leg bridge at varying heights. Participants received a TMS pulse every
205 5-10 seconds and rested in supine between each stimulus. Commencing approximately 2 cm
206 lateral to and anterior to the vertex (Tsao et al. 2008; Fisher et al. 2013), the optimal site of
207 stimulation, or “hotspot” was determined by systematically stimulating a series of locations using
208 the cap grid reference until the location that consistently produced an MEP was determined.

209 The active motor threshold at the gluteus medius hotspot was quantified as the stimulator
210 intensity that produced at least 5 out of 10 MEPs with an amplitude of at least 100 μ V. The
211 motor cortical representation of gluteus medius and lumbar longissimus were mapped at 120%
212 of the active motor threshold, by delivering stimuli at 24 locations spaced 1cm apart in a 6 by 4
213 grid encompassing the motor cortex (MNI x coordinates -1.04;-30.36; MNI y coordinates -42.34:
214 8.23; Figure 1d) (Mayka et al. 2006). Five stimuli were delivered at each location.(Masse-Alarie
215 et al. 2017)

216 *External oblique* – TMS mapping of the external oblique was conducted during posterior pelvic
217 tilting in supine. A consistent level of muscle activation at 20 % MVIC was ensured by providing
218 visual feedback of the external oblique contraction intensity. TMS stimuli were delivered as the
219 participant maintained a sub-maximal posterior pelvic tilt. Participants rested in the supine
220 position for 5 -10 seconds between each stimulus. Determination of the hot-spot, active motor
221 threshold and mapping was conducted as previously described.

222 **2.3 Data processing and analyses**

223 2.3.1 Mediolateral anticipatory postural adjustments

224 To quantify performance of the rapid arm raise task, reaction time and movement time were
225 calculated. Reaction time was defined as the duration from the cue to onset of deltoid muscle
226 activity. Movement time was defined as the duration of time from onset of deltoid activity to the
227 glenohumeral joint reaching 90° of abduction.

228 EMG data were processed in MATLAB[®] using custom-written code. After removal of the DC
229 offset, the EMG signals were band-pass filtered between 40 and 400Hz. This high-pass
230 threshold was set to minimize electrocardiogram (ECG) artifact in the EMG signal. Signals were
231 then full-wave rectified. The latency of the onset of muscle activity for each individual was

232 quantified using the integrated profile or iEMG method (Santello and McDonagh 1998; Allison
233 2003; Smith and Kulig 2016). Onset of activity in each muscle was quantified in ms relative to
234 the onset of the deltoid muscle on the moving arm. Muscle activations were classified as
235 anticipatory postural adjustments if they occurred from 100ms prior to deltoid onset to 50ms
236 after deltoid onset (Figure 1b) (Massé-Alarie et al. 2012). For calculation of coactivation
237 between pairs of muscles, the EMG data were additionally low-pass filtered at 12Hz to obtain a
238 linear envelope and were amplitude normalized to the peak activation occurring in that muscle
239 for that individual throughout the entire arm raise. A coactivation coefficient (CCI) was then
240 calculated for each possible pair of muscles (LL/GMED; GMED/EO; LL/EO) in the same
241 anticipatory postural adjustment time window utilizing equation i)

$$242 \quad i) \sum_{i=1}^N \left(\frac{EMG.low_i}{EMG.high_i} \right) (EMG.low_i + EMG.high_i)$$

243 where N is the number of data points in the anticipatory window. For each instant in time,
244 $EMG.high$ and $EMG.low$ are the amplitude of the signals from each muscle, with $EMG.high$
245 being the muscle with the higher amplitude at that moment and $EMG.low$ being the muscle with
246 the lower amplitude (Nelson-Wong and Callaghan 2010). This index provides a sum of the
247 normalized amplitude of activity for each muscle pair, weighted by the extent of coactivation.

248 2.3.2 Motor cortical representation

249 MEP data were processed in Signal software and MATLAB[®]. Peak-to-peak amplitude of each
250 MEP was extracted from a window 5 to 45ms after the magnetic pulse. Average MEP amplitude
251 was then calculated for each muscle at each grid location. This average amplitude for each
252 location was then normalized to the peak MEP amplitude for that muscle across all grid
253 locations (Tsao et al. 2011; Plow et al. 2014; Masse-Alarie et al. 2017). The center of each
254 muscle representational area was determined by calculating the center of gravity (CoG). The

255 CoG is the amplitude-weighted center of each muscle representational area and is calculated
256 with the following equations:

257 ii) $CoGx = \sum z_i x_i / \sum z_i$

258 iii) $CoGy = \sum z_i y_i / \sum z_i$

259 where x_i and y_i are the medio-lateral and antero-posterior locations respectively and z_i is
260 normalized amplitude (Wassermann et al. 1992; Uy et al. 2002). The CoG, determined using
261 this methodology, is reliable in both young and older adults (Boroojerdi et al., 1999; Uy et al.,
262 2002). Horizontal separation distance between the CoG for each possible pair of muscles was
263 calculated with the Euclidian distance. The volume of the representational area for each muscle
264 was calculated as the sum of the normalized amplitude of MEPs from all grid locations that
265 produced an MEP. To check that the target activation of 20% MVIC had been maintained in
266 gluteus medius and external oblique throughout the experiment, the mean amplitude of EMG
267 activation in the 100ms window immediately prior to the delivery of each stimulus was also
268 calculated.

269 **2.4 Statistical approach**

270 The normality and sphericity of data was assessed using standard procedures (version 24, IBM
271 SPSS Statistics, Armonk, NY). Mann Whitney U tests were utilized to compare reaction time
272 and movement time between groups and active motor threshold for both muscles. Independent
273 t-tests were utilized to compare pre-stimulus activation of GMED and EO.

274 Separate mixed-model ANOVA with between subject factor (group; young adult and older adult)
275 and within subject factor (muscle; lumbar longissimus, gluteus medius and external oblique)
276 were conducted to compare the primary variables for anticipatory postural adjustments and

277 motor cortical representation. Variables for APAs were muscle onset latency and coactivation
278 coefficient between each muscle pair. Variables for motor cortical representation were CoG
279 locations, CoG separation distance and volume of the representational area for the same three
280 muscles. In the case of significant group by muscle interactions, paired post hoc comparisons of
281 a) between groups for each muscle (independent t-tests) and b) within groups for each muscle
282 (paired t-tests) were then made utilizing the Holm-Bonferroni correction for multiple comparisons
283 within each cluster of tests. Estimates of effect sizes for comparisons that reached or
284 approached significance were calculated with an unbiased Cohen's d , with correction for small
285 sample size (d_{unb} , (Fritz et al. 2012)). 0.8 indicates a large effect size, .5 a medium effect size
286 and .3 a small effect size

287 To examine the influence of falls history on all variables, exploratory comparisons between the
288 subgroups of fallers and non-fallers within the older adult group, and between young adults and
289 each subgroup were made with Mann Whitney U tests. Comparisons within subgroups for each
290 muscle were made with the Wilcoxon Signed Ranks Test. Estimates of effect sizes for all non-
291 parametric comparisons were calculated using Cohen's r with 0.5 indicating a large effect size,
292 .3 a medium effect size and .1 a small effect size (Fritz et al. 2012).

293 **3. RESULTS**

294 **3.1 Demographics and balance/mobility tests**

295 Demographics of the young adult and older adult group are provided in Table 1. All of the older
296 adult group participated in regular physical activity. The dominant limb was the right limb for all
297 participants. Therefore, all participants were instrumented with EMG on the right side, utilized
298 their left arm for the arm raising task, and had TMS applied to the left hemisphere. One male

299 older adult with no history of falls did not complete the TMS data collection due to fatigue, and
300 APA data for one female young adult were not recorded due to equipment failure.

301 Scores for the APA section of the BESTest, the TUG time, and self-selected gait speed for the
302 older adults are shown in Table 1.

303 **3.2 Mediolateral anticipatory postural adjustments**

304 Reaction time and movement time were not significantly different between the young and older
305 adult groups (Table 1, $p = 0.740$ and $p = 0.288$ respectively).

306 Muscle onset latency differed between groups, with a significant group by muscle interaction ($F_{(2,21)} = 4.681$, $p = 0.014$). GMED onset was significantly later in older adults than young adults
307 (adjusted $p = 0.039$, unbiased Cohen's $d (d_{unb}) = 1.07$)(Figure 2a & b). Within the older adult
308 group, but not the young adult group, there was a trend for GMED onset being significantly later
309 than LL onset (adjusted $p = 0.069$, $d_{unb} = 1.02$).

311 There was no difference between groups for coactivation index for any of the muscle pairs, with
312 no main effect of group or group by muscle interaction. There was a significant main effect of
313 muscle pairing ($F_{(2,21)} = 8.926$, $p = 0.001$). Post hoc comparisons indicated that there was
314 significantly greater coactivation between LL/GMED than between LL/EO (adjusted $p = .009$,
315 $d_{unb} = 0.44$) (Figure 2c).

316 **3.3 Motor cortical representation**

317 Active motor thresholds, as a percentage of total stimulator output, were not significantly
318 different between the young and older adult groups for either GMED or EO ($p = 0.150$ and $p =$
319 1.000 respectively). The % of MVIC of GMED and EO immediately prior to the delivery of the

320 TMS stimuli was also consistent between the young adult and older adult groups ($p = 0.182$ and
321 0.303 respectively).

322 Motor maps for each muscle in each group are shown in Figure 3. CoGx locations varied by
323 group (group by muscle interaction $F_{(2,21)} = 4.360$, $p = 0.019$). Post hoc comparisons were not
324 significant for any individual muscle. The two groups demonstrated different relative spatial
325 organization of the three muscles. Within the young adults group, LL tended to be more lateral
326 than GMED (adjusted $p = 0.162$, $d_{unb} = 0.60$). Within the older adult group, EO was significantly
327 more lateral than both LL and GMED (adjusted $p = 0.015$ and 0.028 respectively, $d_{unb} = 0.85$
328 and 0.79 respectively, Figure 4a). For COG y location there was a significant main effect of
329 muscle ($F_{(2,21)} = 4.444$, $p = 0.017$). EO was significantly more posterior than LL (adjusted $p =$
330 0.045 , $d_{unb} = 0.52$). There was no main effect of group, or group by muscle interaction (Figure
331 4b).

332 CoG separation distance did not differ between groups. There was a main effect of muscle pair,
333 with LL/GMED separation distance tending toward being smaller than both LL/EO distance and
334 GMED/EO distance (main effect $F_{(2,21)} = 5.059$, $p = 0.020$; post-hoc comparisons adjusted $p =$
335 0.096 in both cases, $d_{unb} = 0.70$ and 0.69 respectively).

336 Volume of motor cortical representational area did not differ between groups. There was a main
337 effect of muscle ($F_{(2,21)} = 3.947$, $p = 0.027$). Volume was significantly larger in the GMED
338 compared with LL (main effect adjusted $p = 0.015$, $d_{unb} = 0.73$).

339 3.4 Subgroup comparisons based on falls history

340 Five out of the twelve older adults reported at least one fall in the preceding year. There was no
341 significant difference in age ($p = 0.684$) or weight ($p = 0.361$) between fallers and non-fallers.
342 BESTest score and TUG performance were the same in fallers and non-fallers ($p = 0.876$ and

343 0.530 respectively). However, fallers had significantly slower gait velocity than non-fallers ($p =$
344 0.016 , effect size $r = 0.49$). Performance of the rapid arm raise task was equivalent between the
345 fallers and non-fallers, with no difference in reaction time or movement time between young
346 adults and older adult fallers ($p = 0.959$ and 0.160 respectively) or young adults and non-fallers
347 ($p = 0.682$ and 0.750).

348 The subgroup analyses comparing young adults with fallers and non-fallers separately showed
349 that age-related changes in mediolateral APAs were most evident in the non-faller group.
350 GMED was significantly later in non-fallers than young adults ($p = 0.022$, $r = 0.52$) but there was
351 no difference in GMED latency between fallers and young adults ($p = 0.234$, Figure 5a). There
352 was also a trend toward significantly less coactivation in the GMED/EO pairing in non-fallers
353 compared with young adults ($p = 0.100$, $r = 0.39$) but no difference in coactivation for any
354 muscle pairing between fallers and young adults ($p > 0.5$ for all comparisons).

355 Active motor threshold of GMED and EO did not differ between the subgroups ($p = 0.931$ and
356 0.662 respectively). Age-related changes in CoG location were most evident in the non-faller
357 group. LL representation was significantly more medial in non-fallers than in young adults ($p =$
358 0.017 , $r = 0.54$) but that there was no difference between the fallers and young adults for any
359 muscle. In the non-fallers, the CoG location for EO was significantly more lateral than both LES
360 and GMED ($p = 0.028$, $r = 0.90$ for both comparisons) but there was no significant difference
361 between COG x locations for the three muscles in the faller group (Figure 5b). There was no
362 significant difference in COG y locations for any muscle between fallers or non-fallers and
363 young adults.

364 Subgroup analyses of separation distance also showed that age-related changes were most
365 evident in the non-fallers. LL /GMED separation distance was significantly less in non-fallers
366 than young adults ($p = .023$, $r = 0.52$) but that there was no difference between fallers and

367 young adults ($p = 0.246$) (Figure 5c). The volume of GMED was significantly smaller in non-
368 fallers than young adults ($p = 0.017$, $r = 0.54$) but that there was no difference between fallers
369 and young adults.

370 **4. DISCUSSION**

371 This study compared the temporal and spatial organization of mediolateral APAs, and the
372 functional representation of the trunk and hip musculature in motor cortex, in healthy young and
373 older adults. For the first time, and in support of our original hypothesis, we found that latency of
374 onset in GMED was delayed in older adults during mediolateral anticipatory postural
375 adjustments. Older adults also demonstrated shifted representational areas for postural
376 musculature in motor cortex. However, the separation distance between the center of gravity for
377 individual muscle representational areas and the volume of each representational area did not
378 differ between the young and older adult groups. The exploratory subgroup analyses indicated
379 that, contrary to our hypotheses, the greatest age-related changes in latency of APAs, muscle
380 coactivation, location of representational area, separation distance and volume of
381 representational area were evident in the non-fallers rather than the fallers. These findings
382 provide some preliminary evidence of potentially adaptive compensations in the non-faller
383 subgroup.

384 In our cohort of healthy, active older adults, performance of the rapid arm raising task did not
385 differ from the young adults in terms of reaction time or movement time. This finding is
386 consistent with existing research indicating that simple (non-choice) reaction time is preserved
387 in older adults (Rogers et al. 1992; Bleuse et al. 2006) and that the velocity of movement is also
388 consistent under low-loading conditions (Bleuse et al. 2006). Despite this similarity in task
389 performance, older adults demonstrated altered temporal organization of the APA synergy. In
390 the young adult group, onset of activity in GMED was prior to that of the trunk muscles. This is

391 consistent with findings from previous studies of anterior-posterior arm raising (Mank'kovskii et
392 al. 1980; Horak et al. 1984). In contrast, GMED was activated last in the older adult group. To
393 our knowledge, previous research examining mediolateral APAs in older adults has exclusively
394 utilized predictable, externally induced perturbations rather than voluntary limb movement. This
395 previous research demonstrated no difference in the magnitude of trunk and hip APAs in older
396 adults with and without a history of falls compared with young adults but did not investigate
397 onset timing or coactivation (Claudino et al. 2013). Taken together, these results support a
398 hypothesis that the temporal organization of APAs and their amplitude scaling are separate
399 constructs with distinct neural substrates and that they may be differently influenced by aging
400 (Bleuse et al. 2006; Jacobs et al. 2009b; Huang and Brown 2013).

401 During rapid mediolateral arm raising, reactive forces and moments caused by the motion of the
402 arm result in trunk/pelvis flexion, trunk side bending and pelvis rotation toward the side of the
403 moving limb (Hodges et al. 1999). APAs in the contralateral GMED, EO and LL resist these
404 forces/moments. In particular, appropriate activation in GMED is critical to stabilize the trunk
405 and pelvis (Santos and Aruin 2008) and to maintain dynamic mediolateral balance in standing
406 (Granata et al. 2005). Therefore it is important to determine why postural GMED onset is
407 delayed in older adults. Studies have demonstrated reduced peak torque and rate of torque
408 development with aging in GMED (Rogers and Mille 2003). Underlying this is Type II fiber
409 atrophy and fatty infiltration that is most evident in older adults with a history of falls (Sato et al.
410 2002; Inacio et al. 2014). Therefore, we speculate that delayed GMED APAs in the present
411 study are reflective of a central nervous system strategy that possibly compensates for impaired
412 GMED muscle composition by reducing the use of this muscle. However, it is also possible that
413 delayed GMED APAs are purely a result of altered muscle fiber composition.

414 Interestingly, the present study did not demonstrate age-related increases in coactivation
415 between the trunk and hip musculature during APAs. A majority of earlier work has

416 demonstrated that older adults utilize greater muscle coactivation, but this has been reported
417 between agonists and antagonists in the lower limb during static standing or walking rather than
418 between synergists during APAs (Hortobágyi and Devita 2006; Hortobágyi et al. 2009; Nagai et
419 al. 2011). Agonist/antagonist coactivation serves to stiffen joints in the presence of impaired
420 postural control, and it is possible that this occurred in the present study in other lower limb
421 muscles or between pairs of trunk and hip muscles that were not measured.

422 Altered temporal organization of APA synergies in the trunk and hip musculature in older adults
423 was accompanied by shifts in the representational areas of these muscles in motor cortex. In
424 young adults, the CoG for LL was more lateral than that of GMED. In contrast, in older adults,
425 and particularly the non-fallers, the CoG for EO was more lateral than both LL and GMED. The
426 spatial organization and excitability of representational areas for movement or muscles in motor
427 cortex is highly plastic and is modulated by use or training (Remple et al. 2001; Perez et al.
428 2004; Adkins et al. 2005; Tennant et al. 2012). Therefore, reduced postural utilization of GMED
429 in older adults may be accompanied by merging of the LL and GMED representational areas.
430 These novel findings in older adults are similar to evidence of pain-related adaptations in trunk
431 muscle APAs and reorganized trunk muscle cortical representation in individuals with low back
432 pain (Tsao et al., 2008). As the alteration in motor cortical representation was not accompanied
433 by systematic changes in volume of representational areas or separation distance across our
434 older adult group, it is unlikely that our findings are an artifact of the known reduced brain
435 volume in older adults (Jäncke et al. 2015).

436 Dedifferentiation of the representational areas for the three muscles was not consistently
437 evident in our older adult group. Existing evidence from voluntary motor tasks has suggested
438 that older adults compensate for reduced gray and white matter volume by increased and
439 diffuse activation of multiple motor areas and both hemispheres during movement (Seidler et al.
440 2010; Bernard and Seidler 2012). Ours is the first study to specifically examine if

441 dedifferentiation of representational areas occurs between muscles within the motor cortex of a
442 single hemisphere. The non-faller subgroup did have less spatial differentiation between the
443 representational areas of LL and GMED. Therefore, our findings suggest that age-related
444 dedifferentiation of representational areas is specific to individual muscles rather than a
445 generalized characteristic of muscle representations in motor cortex. Greater overlap between
446 individual muscle representational areas may facilitate task-specific synergistic activity in
447 muscles that are frequently activated together (Masse-Alarie et al. 2017). In support of this,
448 across both groups, the smaller separation distance between LES and GMED was
449 accompanied by greater coactivation between those muscles during APAs. The subgroup
450 analysis also showed that increased distance between EO and GMED was accompanied by
451 decreased coactivation between those two muscles in the non-faller group.

452 As we did not follow these individuals over time, it is not possible to identify a causal or temporal
453 relationship between adaptations in APAs, changes in motor cortical representational areas,
454 and falls. However, our subgroup analyses suggest two possibilities. The first is that the
455 significant adaptations evident in the non-faller group represent an adaptive response to altered
456 GMED peripheral muscle characteristics. The adaptive response is evident as a lesser role for
457 GMED in APAs and is accompanied by merging of the LL and GMED representational areas.
458 The alternative interpretation is that the findings from the non-faller group are representative of
459 normal age-related changes, and that the faller group had developed adaptations that make
460 them more consistent with young adults as an attempt to improve postural control following a
461 fall. However, since our faller group demonstrated impaired motor behavior, including
462 decreased gait velocity, compared with the non-faller group, the latter explanation seems less
463 likely.

464 There are some limitations to the present study. Although the sample size was small it was
465 based on a *a priori* power analysis. Further, our group comparisons are supported by a

466 conservative approach to hypothesis testing and demonstrate large effect sizes. Challenges in
467 recruiting male older adults who met the inclusion/exclusion criteria for TMS resulted in an
468 unequal sex distribution. However, in the young adult group there were no differences between
469 males and females for any of the variables, and we are not aware of any research indicating
470 sex-related differences in APAs or motor cortical representations in older adults. Finally, the
471 results of this study may not extrapolate to other postural motor behaviors as multiple task-
472 dependent factors influence the temporal and spatial organization of APAs. These include the
473 speed and direction of movement, self-paced versus external cuing, and whether the
474 perturbation is induced by a voluntary movement or by an anticipated external perturbation
475 (Horak et al. 1984; Santos and Aruin 2008).

476 This study demonstrates for the first time that motor cortical representation of trunk and hip
477 musculature is altered in healthy older adults and that this is accompanied by disordered
478 anticipatory postural adjustments. Understanding age-related changes in anticipatory postural
479 adjustments, and the neural correlates of these changes will assist in optimizing interventions to
480 maintain and improve balance in older adults.

481

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489

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645 **FIGURE LEGENDS**

646

647 **Figure 1.** a) Experimental set up for standing rapid arm raise showing participant instrumented
648 with surface EMG electrodes on deltoid, contralateral lumbar longissimus, gluteus medius and
649 external oblique (not pictured). b) Window for anticipatory postural adjustments from 100ms
650 before to 50ms after onset of deltoid. Task reaction time calculated as time from go signal to
651 deltoid onset. Task movement time calculated as time from deltoid onset to 90 degrees shoulder
652 abduction (end of trial). c) Experimental set up for TMS mapping of gluteus medius and lumbar
653 longissimus. Participant is performing a double-leg bridge while applying an abduction force to
654 the band placed around the distal thighs. d) 6 by 4 grid for mapping centered over motor cortex
655 using stereotactic image guidance, with exemplar motor evoked potentials from 4 grid locations
656 for the external oblique muscle.

657 **Figure 2.** a) Exemplar EMG data from a single trial for a young adult and older adult indicating
658 onset of deltoid activation (red line). b) Group data for onset latency of contralateral lumbar
659 longissimus (LL), gluteus medius (GMED) and external oblique (EO) relative to onset of deltoid
660 activation (DELTA). Negative values indicate onsets in postural muscles that occurred prior to
661 onset in DELTA. Note significant difference in GMED onset between young and older adults (*p =
662 0.039). c) Group data for the sum of the normalized amplitude of activity for each muscle pair,
663 weighted by the extent of coactivation (coactivation index, CCI). Muscle pairs are lumbar
664 longissimus/gluteus medius (LL/GMED), lumbar longissimus/external oblique (LL/EO), and
665 gluteus medius/external oblique (GMED/EO). Note significant difference between CCI of
666 LL/GMED and LL/EO (*p = 0.009).

667 **Figure 3.** Averaged motor maps for the young adult group (top) and older adult group (bottom)
668 showing location of the representational area for external oblique (EO), lumbar longissimus (LL)
669 and gluteus medius (GMED) mapped on a 6 by 4cm grid. The colorbar indicates average
670 normalized MEP amplitude. Average location of center of gravity for each group is
671 superimposed in black on each map.

672 **Figure 4.** Location of center of gravity (CoG) for lumbar longissimus (LL), gluteus medius
673 (GMED) and external oblique (EO) in the young adult group and the older adult group. a) CoG x
674 location. Note that EO is significantly more lateral than LL and GMED in the older adult group
675 (*p = 0.015 and 0.028 respectively). b) CoG y location. Note that EO is significantly more
676 posterior than LL in both groups (*p = 0.045).

677 **Figure 5.** Subgroup comparisons based on falls history. a) Individual data for onset latency of
678 contralateral gluteus medius (GMED) relative to onset of deltoid activation. GMED was
679 significantly later in non-fallers than young adults (p = 0.022). b) Individual data for center of
680 gravity x location (CoG x location) for lumbar longissimus (LL), gluteus medius (GMED) and
681 external oblique (EO). LL representation was significantly more medial in non-fallers than in
682 young adults (*p = 0.017). In the non-fallers, the CoG location for EO was significantly more
683 lateral than both LL and GMED (*p = 0.028 for both comparisons).

684

685 **Table 1.** Demographics and balance/mobility test performance for young adults (n = 13) and
686 older adults (n = 12). Values are means ± standard deviation.

	Young adults	Older adults
Age (years)	25.75 (2.09)	72.42 (8.16)
Sex (number of females)	8	10
Mass (kg)	62.46 (9.82)	67.24 (11.75)
BESTest APA score (%)	-	80.56 (11.23)
Timed up and Go Test (s)	-	7.91 (1.56)
Self-selected gait velocity		1.27 (0.18)
Reaction time (s)	0.25 (0.04)	0.27 (0.06)
Movement time (s)	0.38 (0.06)	0.41 (0.09)

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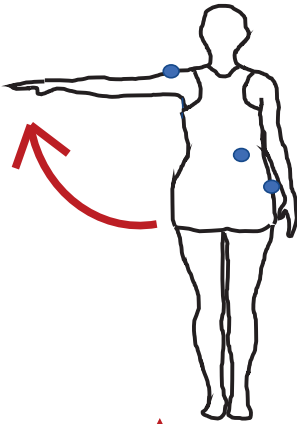
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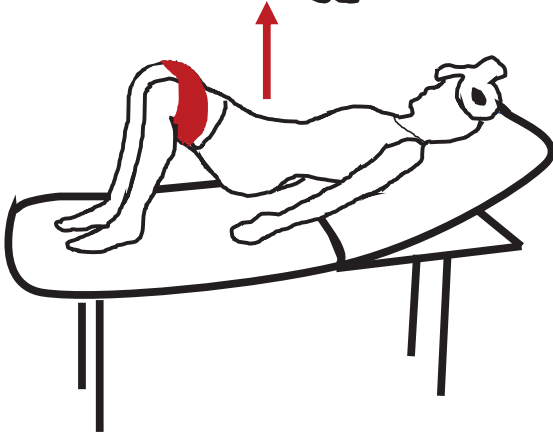
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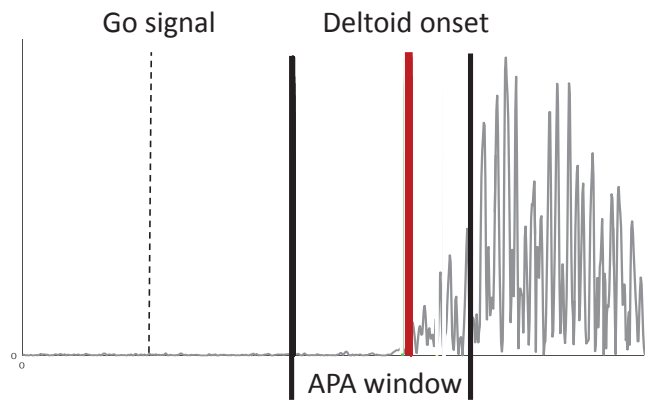
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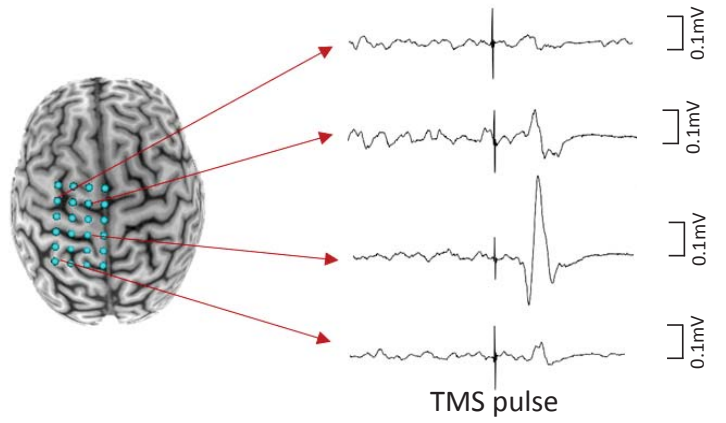
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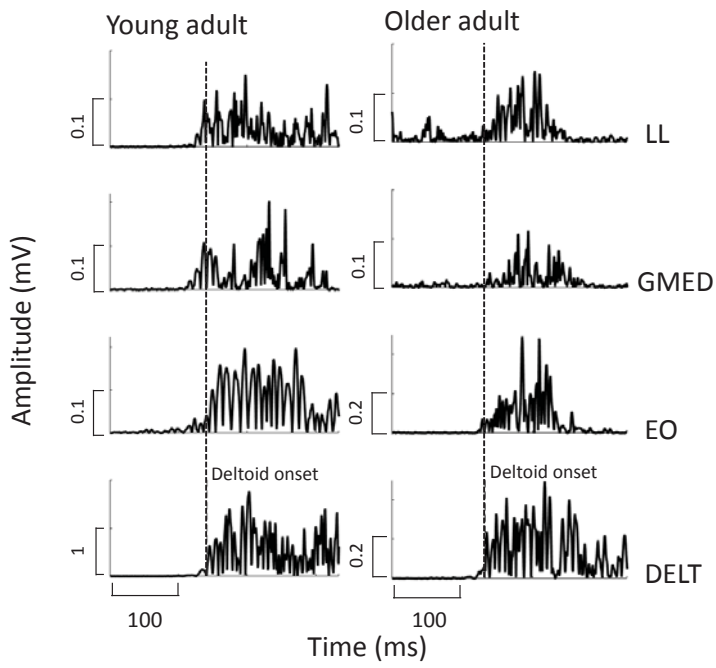
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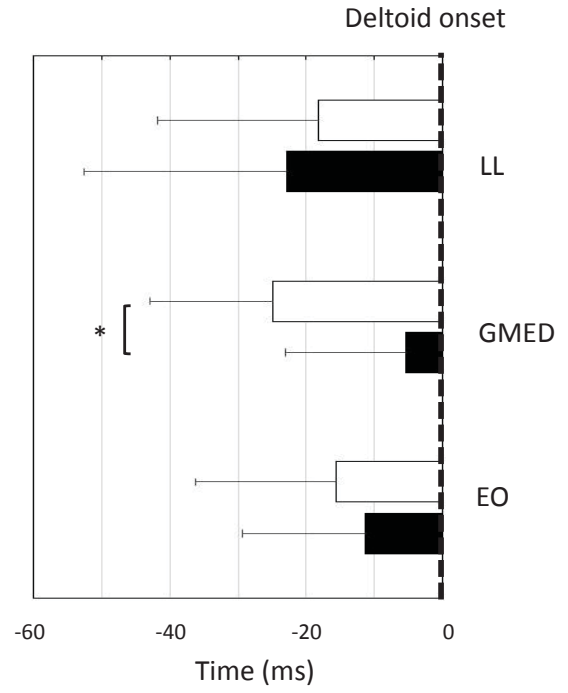
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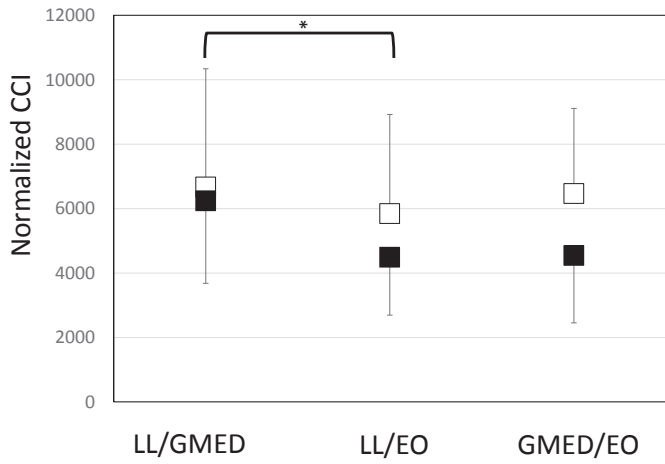
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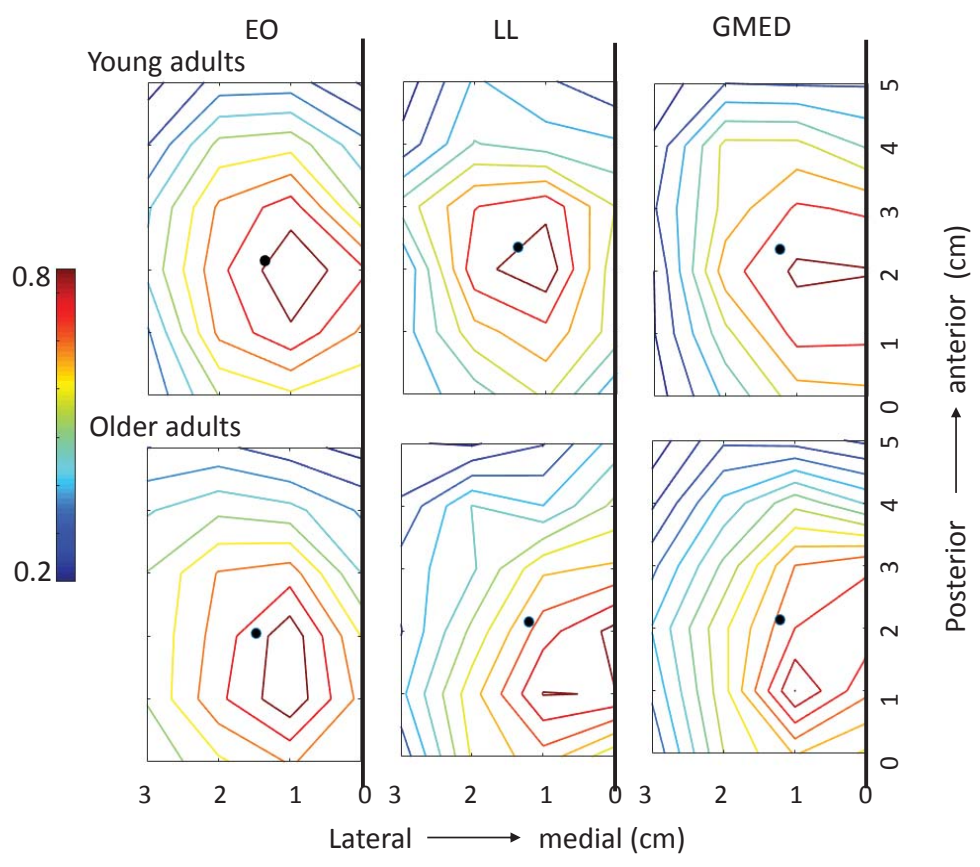


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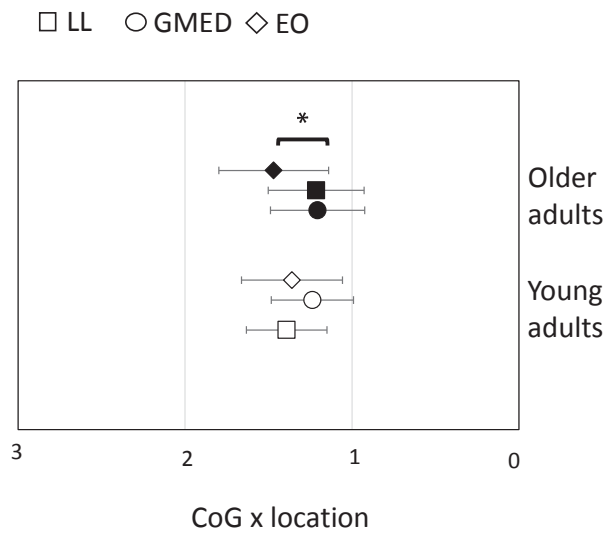


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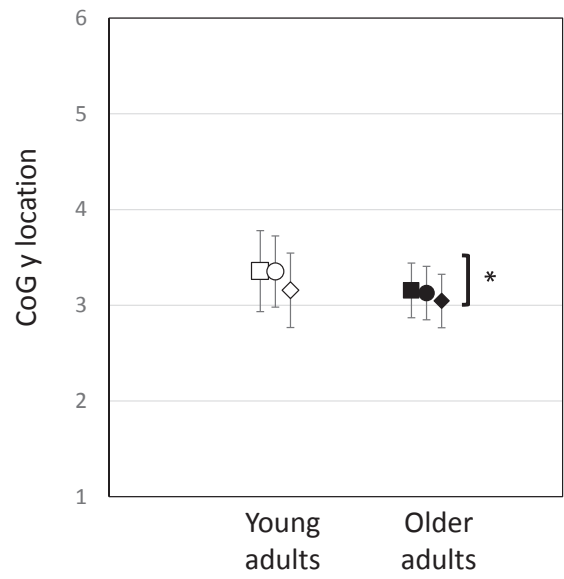




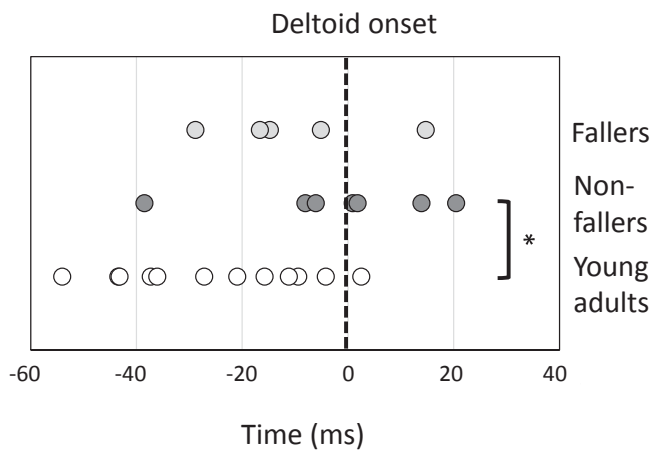
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