

1 **Reliability of two methods for identifying the postural**
2 **phase of gait initiation in healthy and post-stroke**
3 **subjects**

4 Andreia S. P. Sousa (PhD)

5 Escola Superior da Tecnologia de Saúde do Instituto Politécnico do Porto, Área
6 Científica de Fisioterapia, Centro de Estudos de Movimento e Atividade Humana, Rua
7 Valente Perfeito, 322 - 4400-330 Vila Nova de Gaia, PORTUGAL
8 E-mail: asp@estsp.ipp.pt, andreia.asps@gmail.com

9
10 Augusta Silva (PhD)

11 Escola Superior da Tecnologia de Saúde do Instituto Politécnico do Porto, Área
12 Científica de Fisioterapia, Centro de Estudos de Movimento e Atividade Humana, Rua
13 Valente Perfeito, 322 - 4400-330 Vila Nova de Gaia, PORTUGAL
14 E-mail: smaugusta@gmail.com

15
16 Rubim Santos (PhD)

17 Escola Superior da Tecnologia de Saúde do Porto, Área Científica de Física, Centro de
18 Estudos de Movimento e Atividade Humana, Rua Valente Perfeito, 322 - 4400-330 Vila
19 Nova de Gaia, PORTUGAL
20 E-mail: rss@estsp.ipp.pt

21
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25 **Corresponding author:**

26 Andreia S. P. Sousa (PhD)
27 Escola Superior da Tecnologia de Saúde do Instituto Politécnico do Porto, Área
28 Científica de Fisioterapia, Centro de Estudos de Movimento e Atividade Humana, Rua
29 Valente Perfeito, 322 - 4400-330 Vila Nova de Gaia, PORTUGAL
30 E-mail: asp@estsp.ipp.pt, andreia.asps@gmail.com

31 **Abstract**

32 This study aims to compare two methods of assessing the postural phase of gait
33 initiation as to intrasession reliability, in healthy and post-stroke subjects. As a
34 secondary aim, this study aims to analyse anticipatory postural adjustments during gait
35 initiation based on the centre of pressure (CoP) displacements in post-stroke
36 participants. The CoP signal was acquired during gait initiation in fifteen post-stroke
37 subjects and twenty-three healthy controls. Postural phase was identified through a
38 baseline-based method and a maximal displacement based method. In both healthy and
39 post-stroke participants higher intra-class correlation coefficient and lower coefficient
40 of variation values were obtained with the baseline-based method when compared to the
41 maximal displacement based method. Post-stroke participants presented decreased CoP
42 displacement backward and toward the first swing limb compared to controls when the
43 baseline-based method was used. With the maximal displacement based method, there
44 were differences between groups only regarding backward CoP displacement. Postural
45 phase duration in medial-lateral direction was also increased in post-stroke participants
46 when using the maximal displacement based method. The findings obtained indicate
47 that the baseline-based method is more reliable detecting the onset of gait initiation in
48 both groups, while the maximal displacement based method presents greater sensitivity
49 for post-stroke participants.

50

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Introduction

58 Gait initiation is an important part of locomotion and has been described as the
59 transient state between two steady states - standing and walking.^{1,2} This transition from
60 a quasi-static state (quiet standing) to a dynamic state (walking) is considered to be
61 governed by a motor program, as stereotyped patterns of activity, soleus inhibition and
62 tibialis anterior activation, and invariant relative timing have been demonstrated.^{3,4}
63 These first phase mechanisms, namely anticipatory postural adjustments, are
64 responsible for moving the centre of pressure (CoP) under the feet backward and toward
65 the first swing limb.⁵⁻⁷ In turn, CoP displacement increases anterior-posterior and
66 medial-lateral components of the ground reaction force, thereby generating momentum
67 in those directions for taking a step before the centre of mass moves out of the base of
68 support.^{4,7} Thus, the central nervous system uses stable, efficient mechanisms for
69 dealing with the inherent instability of upright bipedalism during gait initiation.^{8,9} For
70 this reason, CoP displacement backward and toward the first swing limb has been
71 identified as the postural phase of gait initiation.¹⁰⁻¹⁴

72 Disturbance of gait initiation is common in patients with central nervous system
73 impairment, like stroke. In this condition, postural adjustments' dysfunction during the
74 postural phase is related to disturbance in the first step.¹⁵⁻¹⁸ However, despite the
75 importance of the postural phase in gait initiation performance, there has been a poor
76 standardisation of methods to identify the onset of the postural phase of gait initiation,
77 as different variables have been used: centre of mass migration and acceleration, ground
78 reaction force and CoP related variables.^{10-15,18-22} Whereas studies assessing the centre
79 of mass and ground reaction force stated how the event was computed, the same is not
80 observed in studies involving CoP related variables.^{10-15,18,20-22} Since gait initiation is the
81 transition between standing and walking, two methods used in centre of mass
82 displacement evaluation may be transferred to CoP variables: 1) one based CoP

83 displacement during upright standing (baseline-based method), and 2) another based on
84 maximal CoP displacement backward and toward the first swing limb (maximal
85 displacement based method).²⁰⁻²² While the methods used to identify the beginning of
86 the postural phase of gait initiation are poorly standardised, the end of the postural
87 phase has been identified most often as the instant where the CoP reaches its maximum
88 backward and toward the first swing limb positions.^{10,12,13} The methods used in studies
89 assessing the centre of mass can be transferred to CoP variables to identify the onset of
90 the postural phase of gait initiation. However, it is important to know their reliability, as
91 this analysis has not been done yet.

92 Given the postural phase impacts on forward displacement performance, it is
93 important to select a reliable method to assess this particular phase of gait initiation as
94 measurement errors can seriously affect statistical analysis and interpretation.²³ This
95 should be analysed in healthy subjects, but also in subjects with lower performance in
96 gait initiation as post-stroke subjects.^{24,25} Such knowledge has the potential to provide a
97 foundation for answering research questions about the most reliable method to assess
98 the postural control phase of gait initiation in pathologic and non-pathological
99 conditions, and to assess motor control, as the onset of CoP displacement is a key event
100 for electromyography analysis when postural adjustments are investigated.^{26,27} From a
101 clinical point of view, this study contributes to establish how outcomes of interventions
102 can be quantified to assess postural control measures.

103 The aim of the present study was to compare the reliability of CoP displacements
104 during the postural phase of gait initiation calculated by two methods of detection the
105 beginning of the postural phase (baseline-based vs. maximal displacement based
106 methods) in health and post-stroke participants. For this, the intra-session reliability was
107 calculated to assess the variability of each method.²⁸ As secondary aim, this study

108 analysed anticipatory postural adjustments based on the CoP displacements in post-
109 stroke participants. Based on the findings obtained by Breniere (1996) that the natural
110 body frequency (ratio between the amplitude of the centre of mass and of the CoP) is an
111 absolute invariant parameter, specific to human standing and gait, it can be
112 hypothesised that CoP displacement values calculated with the baseline-based method
113 than with the maximal displacement based method are more reliable.²⁹ As to the
114 secondary purpose, based on the results obtained by Brunt (1995), demonstrating weight
115 bearing asymmetry in subjects with stroke, and by Hesse et al. (1997), demonstrating
116 changes in temporal muscle sequence during gait initiation, it can be hypothesised that
117 post-stroke subjects present decreased CoP shift backward and toward the swing limb
118 regardless of the method used.^{16,30}

119

120 **Methods**

121 *Participants*

122 Fifteen patients who had suffered a stroke at least 6 months earlier (8 females, 7
123 males) and 23 healthy participants (12 females, 11 males) participated in this study
124 (demographic descriptors can be found in Table 1). The mean time between their stroke
125 and the time of inclusion in this study was 24.9 ± 11.5 months (6-40 months). All post-
126 stroke participants suffered an ischemic stroke at the subcortical level (internal capsule):
127 8 of them had suffered an infarction in their left hemisphere, whereas 5 had suffered an
128 infarction in their right hemisphere. To be included, patients were required to: (1) have
129 suffered a first-ever ischemic stroke involving the middle cerebral artery territory, as
130 revealed by computed tomography, resulting in hemiparesis; (2) have a Fugl-Meyer
131 (Assessment of Sensorimotor Recovery After Stroke scale) score in the motor
132 subsection below 34;³¹ (3) have the ability to walk, with close supervision if necessary,

133 but without physical assistance as judged by the treating physiotherapist; (4) have the
134 ability to stand with feet apart for 30 seconds or more; and (5) have provided written or
135 verbal informed consent. Patients were excluded for one or more of the following
136 reasons: (1) cognitive deficit that could hinder communication and cooperation (score
137 below 24 in the Mini-Mental State Examination); (2) history of orthopaedic or
138 neurological (other than stroke) disorders, known to affect walking performance and
139 quiet standing position; (3) history of stroke involving the brainstem or cerebellar areas;
140 and (4) taking medication such as antispasticity medication that could affect motor
141 performance and balance. Gait data of post-stroke participants were compared with data
142 obtained from the 23 healthy control participants. All participants in the control group
143 were sedentary and were selected according to the same exclusion criteria which were
144 applied to the post-stroke group; they were excluded if they had suffered any
145 neurological disorder. Participants were considered sedentary if their practice of
146 physical activities was less than three times per week during 20 minutes of continuous
147 vigorous physical activities or less than 5 times per week during 30 minutes of
148 continuous or intermittent moderate physical activities for at least the last 2 years. The
149 study was approved by the local ethics committee and implemented according to the
150 Declaration of Helsinki.

151 *Instrumentation*

152 The values of the vertical (F_z), anterior-posterior (F_x) and medial-lateral (F_y)
153 components of the ground reaction force, as well as the values of the moments of force
154 in the frontal (M_y) and sagittal (M_x) planes, were acquired using a force plate^a at a
155 sampling rate of 1000Hz (FP4060-08 model from Bertec Corporation (USA), connected
156 to a Bertec AM 6300 amplifier ^a and to an analogue board ^b, from Qualysis, Inc.
157 (Sweden)).

158 The force plate signals were analysed with the Acqknowledge software (Biopac
159 Systems, Inc., USA).

160 *Procedures*

161 Data acquisition

162 All participants used their own regular footwear (1.5cm heel) while standing on a
163 force plate, with feet at pelvis width and with their arms by their sides. They were asked
164 to stand as still as possible and to focus on a target 2 meters away and at eye level for 30
165 seconds. After this, participants were asked to walk at self-selected speed over a 5 m
166 walkway, without explicit instructions. If a subject asked which leg to start with, the
167 researcher replied “whatever feels natural for you,” as lower limb preference plays an
168 influential role in the control of frontal plane body motion during gait initiation.³²
169 However, participants were asked to keep the starting leg consistent over all trials. A
170 trial was considered valid when the subject performed at least three steps. Each subject
171 performed three trials with rest periods of 60 seconds between each trial, when the
172 subjects remained seated. Before data acquisition, sufficient time was given so that the
173 participants became familiar with the experimental settings.

174 Data processing

175 Ground reaction force signals were low-pass filtered using a fourth-ordered
176 Butterworth filter by using a zero-phase lag with a cut-off frequency of 20 Hz. The
177 acquired force and moment of force-time series of each trial were used to calculate the
178 CoP fluctuation in the anterior-posterior (AP) and medial-lateral (ML) directions using
179 the following approximations:

$$180 \quad \text{CoP}_{\text{AP}} = \frac{M_y}{F_z}, \quad (1)$$

181
$$\text{CoP}_{\text{ML}} = \frac{M_x}{F_z} \quad (2)$$

182 CoP displacement in AP and ML directions, during the postural phase of gait
183 initiation, was calculated using the difference between maximum CoP backward (first
184 inflection of CoP_{AP}) and toward the swing limb (first inflection of CoP_{ML}) positions
185 and the CoP position associated to the beginning of its displacement for each direction,
186 respectively (Figure 1). Two methods were used to identify the beginning of CoP
187 displacement: (1) a baseline-based method and (2) a maximal displacement based
188 method. In both methods, identification was achieved using a computer program and
189 visual inspection.

190 Baseline-based method

191 The mean of peak-to-peak amplitude and the dispersion time series estimated by
192 standard deviation (SD) of CoP displacement were calculated for the ML and AP
193 directions from 5 to 25 seconds of upright quiet standing. The mean plus 3 times the SD
194 was defined as the threshold for gait initiation onset. The CoP position at the beginning
195 of its displacement backward and towards the swing limb was assessed at the beginning
196 of an interval lasting for at least 50 ms when its absolute value was higher than the
197 threshold (Figure 1).¹⁵ Only changes of CoP displacement with a minimum duration of
198 50 ms were considered, to exclude variations that are not related to gait initiation, as this
199 interval corresponds to the electromechanical delay.³³

200 Maximal displacement based method

201 The CoP position in AP and ML directions was assessed at the beginning of the
202 interval lasting at least 50 ms, when its value was higher than 5% of the magnitude of
203 the first inflection of CoP_{AP} displacement and of the magnitude of the first inflection of
204 CoP_{ML} displacement, respectively (Figure 1).¹⁵

205 The threshold's selection was adapted from methods used on previous studies that

206 have used the same criterion for other biomechanical variables, and on the fact that it
207 provided a good agreement with visual inspection.^{21,22,34}

208 *Statistical analysis*

209 The acquired data were analysed using the Statistic Package Social Science (SPSS)^c
210 software version 22, from *IBM Company* (USA). Reliability measures of CoP
211 displacement assessed from each method were calculated for healthy (n=23) and post-
212 stroke participants (n=15). The Intra-Class Correlation Coefficient (ICC_{2,1}) with a 95%
213 Confidence Interval (CI) was used because it considers random effects over time and
214 expresses relative reliability of the measures of CoP displacement obtained with each
215 method.²³ Specifically, a two-way ANOVA model with a random subject effect was
216 used to estimate the intra-session reliability. The following range of reliability
217 coefficients were used to report the degree of reliability: 0.00 to 0.25 – little, if any
218 correlation; 0.26 to 0.49 – low correlation; 0.50 to 0.69, moderate correlation; 0.70 to
219 0.89, high correlation and 0.90 to 1.00, very high correlation.³⁵ The Coefficient of
220 Variation (CV) was used to express absolute reliability and was calculated per subject,
221 by dividing SD by the mean of three trials.

222 Shapiro–Wilk test results and histogram analysis have shown that data were
223 normally distributed. The statistical difference between ICCs was evaluated through the
224 application of Fisher’s Z transformation, with significance determined with the t
225 statistic. The paired samples T-test was used to compare the CV, CoP displacement and
226 postural phase duration values obtained with each method. The independent samples T-
227 test was used to compare mean values of CoP displacement and CV values between
228 healthy and post-stroke groups. Because of the reduced sample, the Wilcoxon test was
229 used to compare CoP displacement between post-stroke participants that initiated gait

230 with the ipsilesional limb, $n=6$ and those who initiated gait with the contralesional limb,
231 $n=9$. Cohen's d was calculated to assess effect size and power analysis $(1-\beta)$ was
232 performed to give an indication of the power of hypothesis tests and the magnitude of
233 the differences that researchers are able to detect in those settings. A 0.05 significance
234 level was used for inferential analysis

235 **Results**

236 Higher reliability was obtained in CoP displacement values during the postural
237 phase of gait initiation when using the baseline based method, in both healthy and post-
238 stroke participants.

239 In healthy participants, when the baseline-based method was used, CoP
240 displacement measures presented high to very high correlation, while values obtained
241 with the maximal displacement based method presented high correlation (Table 2).
242 Despite a tendency to lower values of ICC in the maximal displacement based method,
243 no significant differences were observed (CoP_{AP} , $p=.104$; CoP_{ML} , $p=.164$). When
244 analyzing CV values, statistically significant differences between methods occurred in
245 both CoP_{AP} ($p=.001$, $(1-\beta)=.99$, Cohen's $d=1.11$) and CoP_{ML} ($p=.005$, $(1-\beta)=.45$,
246 Cohen's $d=.37$), with higher values in the maximal displacement based method (Table
247 2).

248 In post-stroke participants, CoP displacement calculated using the baseline-based
249 method presented high to very high intra-session correlation values, while moderate to
250 high intra-session correlation values were obtained using the maximal displacement
251 based method. However, there were no significant differences (CoP_{AP} , $p=.278$; CoP_{ML} ,
252 $p=.194$). The differences in reliability between the methods were more pronounced in

253 CV values, as higher values were observed in CoP_{AP} displacement ($p=.007$, $(1-\beta)=.99$,
254 Cohen's $d=1.24$) in the maximal displacement based method (Table 2).

255 Generally, CoP displacement was lower in post-stroke participants when compared
256 to healthy participants (Table 2). Specifically, the post-stroke group presented lower
257 CoP displacement backwards ($p=.031$) and towards the first swing limb ($p=.001$) when
258 the baseline-based method was used. Despite decreased CoP displacement, post-stroke
259 participants presented generally higher values of absolute variability (Table 2). When
260 the maximal displacement based method was used, statistical differences were only
261 observed in CoP_{AP} displacement ($p=0.007$) and higher absolute variability was observed
262 in CoP_{AP} displacement ($p=.004$) in post-stroke subjects when compared to health
263 participants (Table 2). No significant differences were observed in ICC values between
264 healthy and post-stroke participants in both methods (Table 2).

265 Nine post-stroke participants initiated gait with their contralesional limb, while six
266 initiated gait with their ipsilesional limb. Globally, a trend to decreased CoP
267 displacement and increased absolute variability was observed in both limbs of post-
268 stroke participants, when compared to healthy participants, in both methods (Figure 2).
269 No differences were observed between post-stroke participants that initiated gait with
270 ipsilesional and contralesional limbs in the baseline-based method (AP, $p=.877$, $(1-$
271 $\beta)=.13$, Cohen's $d=.14$; ML, $p=.643$, $(1-\beta)=.09$, Cohen's $d=.10$) and in the maximal
272 displacement based method (AP, $p=.09$, $(1-\beta)=.34$, Cohen's $d=.3$; ML, $p=.643$, $(1-$
273 $\beta)=.18$, Cohen's $d=.24$).

274 When both methods were compared as to CoP displacement, significant differences
275 were only observed in CoP displacement towards the first swing limb in subjects with
276 stroke ($p=0.039$). Higher values were obtained using the maximal displacement based

277 method (Table 2). In general, a trend to higher durations of postural phase were
278 obtained with this method for CoP displacement towards the first swing limb (baseline-
279 based method, 431 ± 209 ms (healthy), 563 ± 281 ms (post-stroke); maximal displacement
280 based method, 504 ± 196 ms (healthy), 631 ± 371 ms (post-stroke)), while a trend to higher
281 duration of postural phase was obtained for CoP backwards displacement using the
282 baseline-based method (baseline-based method, 548 ± 259 ms (healthy), 618 ± 252 ms
283 (post-stroke); maximal displacement based method, 366 ± 187 ms (healthy), 509 ± 346 ms
284 (post-stroke)). Despite this tendency, statistically significant differences between
285 methods were obtained for the duration of the postural phase in ML direction in subjects
286 with stroke ($p=0.028$).

287 **Discussion**

288 Generally, both methods were reliable for identifying the postural control phase of
289 gait initiation in healthy subjects. This low within-subject variability, associated with
290 the non-significant differences observed between CoP displacement obtained with the
291 two methods, demonstrates that both methods can be used to identify the postural phase
292 of gait initiation in healthy participants. These high values of intra-session reliability are
293 consistent with the evidence that the initiation of gait is accomplished by stereotyped
294 patterns of activity and consequently stereotyped trajectory of CoP displacement.^{3,4}
295 However, the results obtained as to absolute reliability favour the use of the baseline-
296 based method over the maximal displacement based method.

297 Higher differences between methods were obtained in post-stroke participants.
298 Lower values of absolute variability were obtained with the baseline-based method
299 associated with a trend to higher values of ICC, when compared to the maximal
300 displacement based method. These findings seem to corroborate the high values of
301 reliability for CoP displacement parameters obtained in upright standing in post-stroke

302 subjects.^{36,37} In fact, this has been demonstrated to occur despite the standing balance
303 of post-stroke participants is characterised by increased CoP displacement and an
304 asymmetrical weight bearing distribution in favour of the ipsilesional limb.^{36,37}

305 It has been shown that increased CoP movements during quiet standing in post-
306 stroke subjects seem partly related to increased body sway and partly to exaggerated
307 corrective ankle mechanisms.^{38,39} Based on this, it seems reasonable that the baseline-
308 based method is associated with lower values of CoP displacement towards the first
309 swing limb during the postural phase of gait initiation when compared to the maximal
310 displacement based method. As there is higher CoP displacement during standing in
311 post-stroke participants, the baseline standard deviation is higher. As a consequence, it
312 takes longer to mark the onset of CoP displacement. In the present study, increased
313 duration of the postural phase in ML direction was obtained in the post-stroke group
314 when the maximal displacement based method was used. The greater the displacement
315 of CoP in baseline, the lower will be the window of gait initiation extracted by the
316 baseline method. As a consequence, CoP displacement values would be lower when
317 compared to the maximal displacement method. In this case, the maximal displacement
318 based method would possibly be more accurate to identify the CoP displacement
319 towards the first swing limb onset as the baseline method is more influenced by the
320 amount of sway during standing (baseline). In fact, when compared to healthy
321 participants, different results have been obtained in post-stroke participants with each
322 method. While lower CoP backwards displacement and lower CoP displacement
323 towards the first swing limb was observed when the baseline-based method was used,
324 no differences occurred in CoP_{ML} displacement with the maximal displacement based
325 method. The differences between control and post-stroke groups as to CoP_{ML}
326 displacement assessed with the baseline-based method may be caused by differences in

327 baseline due to differences in body sway. However, the results obtained in the present
328 study do not allow us to confirm this hypothesis.

329 Globally, the findings of the present study support the use of the baseline-based
330 method to assess in a reliable way the onset of gait initiation. However, because the
331 baseline method is more influenced by the amount of sway during standing, the
332 maximal displacement based method seems to present greater sensitivity in identifying
333 the beginning of CoP_{ML} displacement in post-stroke subjects. A higher variability of
334 CoP_{ML} displacement during standing can explain the higher sensitivity of maximal
335 displacement for detecting the beginning of CoP displacement towards the first swing
336 limb. However, future studies are required to confirm this hypothesis.

337 The lower CoP displacement backwards and towards the first swing limb, observed
338 in post-stroke participants, impairs posture stability and motor performance.⁴⁰ The
339 decrease of CoP backward displacement during the postural phase leads to lower
340 generation of forward momentum of the centre of mass and a consequent impairment of
341 gait velocity and step length.^{7,18} The dynamic stability is also compromised, as the
342 reduction of CoP shift towards the swing-leg side increase the extent to which the centre
343 of mass falls toward the swing-leg side during step execution, reducing the ML stability
344 during gait initiation.^{8,41}

345 The decreased backward CoP shift has been associated with a decreased inhibition
346 of the soleus and the gastrocnemius and lower tibialis anterior activity associated with a
347 delayed onset.^{11,17} The dysfunction of the postural phase of the contralesional limb can
348 result from tibialis anterior activation deficit as a consequence of affection of the lateral
349 cortico-spinal system and from impairments in anticipatory postural adjustments as a
350 result of a deregulation of supplementary motor area and premotor cortex. A
351 deregulation of supplementary motor area and premotor cortex is typically found in

352 post-stroke subjects showing damage in the territory of the middle cerebral area.⁴²⁻⁴⁴ It
353 should be noted that a lesion in the premotor cortex affects the anticipatory postural
354 adjustments of bilateral lower extremities in step initiation.⁴³ Postural control
355 dysfunction of the ipsilesional limb has been demonstrated in other functional tasks and
356 particularly in participants with sub-cortical injuries located at the internal capsule
357 level.⁴⁵⁻⁴⁷ In fact, injuries located at this region are typically associated with dysfunction
358 of the ventral–medial systems and may justify changes in the activity of the ipsilesional
359 soleus muscle.⁴⁸ The neuronal connection of the soleus and the tibialis anterior need
360 further discussion, but it can constitute a possible explanation to the dysfunction of the
361 postural phase of gait initiation in the ipsilesional limb. Under this perspective, future
362 research should attempt to investigate the activation patterns of ankle muscles during
363 gait initiation of post-stroke participants.

364 The results obtained in this study indicate that both methods are reliable tools to
365 assess the postural phase of gait initiation in healthy and post-stroke participants. Higher
366 values of CoP displacements reliability were obtained with the baseline-based method.
367 However, because the baseline method is more influenced by the amount of sway
368 during standing, the maximal displacement based method seems to present greater
369 sensitivity in identifying the beginning of CoP_{ML} displacement in post-stroke subjects.
370 Both methods demonstrated that post-stroke participants present decreased CoP
371 displacement during the postural phase of gait initiation. From a clinical point of view,
372 these results indicate that attention should be given to the postural phase of gait
373 initiation in the rehabilitation of post-stroke participants.

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511

512 **Suppliers**

513 a. Bertec Corp, 6171 Huntley Rd, Ste J, Columbus, OH 43229.

514 b. Qualysis AB, Packhusgatan 6, 411 13 Gothenburg, Sweden.

515 c. SPSS Inc, 233 S Wacker Dr, 11th Fl, Chicago, IL 60606.

516

517 **Figure Captions**

518 Figure 1: Representation of CoP displacement during the beginning of gait initiation
519 with the right limb as first swing limb. The gray box in part a) represents an
520 approximation of postural control phase of gait initiation. The onset of CoP
521 displacement for AP and ML determined by the baseline-based method (grey line) and
522 the maximal displacement based method (dark line) are more precisely represented in
523 part b). Dashed lines represent the threshold obtained with each method. The gray box
524 in part b) represents postural control phase of gait initiation obtained by both methods.

525 Figure 2: Mean (bars) \pm SD (error bars) of CoP displacement and CV obtained in
526 ipsilesional and contralesional limbs of post-stroke subject and healthy controls during
527 the postural phase of gait initiation with the baseline-based method and the maximal
528 displacement based method.

529