

1 **Agreement between measurements of stance width using motion capture and center of**
2 **pressure in individuals with and without Parkinson's disease**

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31 **Abstract**

32 *Background*

33 Many individuals with Parkinson's disease exhibit narrow stance width during balance and gait. Because
34 of this, stance width is an important biomechanical variable in many studies. Measuring stance width
35 accurately using kinematic markers in parkinsonian patients can be problematic due to occlusions by
36 research staff who must closely guard patients to prevent falls.

37 *Methods*

38 We investigated whether a measure of stance width based on the mediolateral distance between the center
39 of pressure under each foot could approximate stance width measured with kinematic data. We assessed
40 the agreement between estimates of stance width obtained from simultaneous kinematic and center of
41 pressure measures during quiet standing in 15 individuals (n=9 parkinsonian, n=6 age-similar
42 neurotypical). The source data (1363 unique trials) contained observations of stance width varying
43 between 75–384 mm (\approx 25-150% of hip width).

44 *Findings*

45 Stance width estimates using the two measures were strongly correlated ($r = 0.98$). Center of pressure
46 estimates of stance width were 48 mm wider on average than kinematic measures, and did not vary across
47 study groups ($F_{2,12}=1.81$, $P<0.21$). The expected range of differences between the center of pressure and
48 kinematic methods was 14–83 mm. Agreement increased as stance width increased ($P<0.02$).

49 *Interpretation*

50 It is appropriate to define stance width based on center of pressure when it is convenient to do so in
51 studies of individuals with and without Parkinson's disease. When comparing results across studies with
52 the two methodologies, it is reasonable to assume a bias of 48 mm.

53 **Keywords**

54 Postural control; Center of pressure location; Measurement; Methodology; Foot position

55

56 1. Introduction

57 Many individuals with Parkinson's disease (PD) exhibit narrow stance width during balance and gait
58 (1). Clinically, "narrow stance" is a postural abnormality in which the feet are placed substantially medial
59 to the anterior superior iliac spines (ASIS) (2). Stance width is therefore an important variable in many
60 studies of parkinsonian posture and balance (e.g., (3-5)). It is typically treated as a nominal single value or
61 as a range of values described by the mediolateral distance between kinematic markers placed on the
62 heels, or between the medial malleoli (3-5).

63 Due to repeated protective steps, dyskinesias, and other practical concerns when studying
64 parkinsonian balance, it is difficult to control stance width precisely during experiments – and so ideally,
65 stance width should be measured as a continuous covariate throughout an experiment. However, doing so
66 with kinematic markers can be problematic due to occlusions by research staff who must carefully guard
67 patients to prevent falls.

68 Here, we investigated whether a proxy measure of stance width based on the mediolateral distance
69 between the centers of pressure (CoP) beneath each foot could approximate stance width measured
70 kinematically. As typically defined (6), the CoP is the point location of the vertical ground reaction force
71 vector beneath the entire body, and represents a weighted average of all the pressures over the surface
72 area in contact with the ground (6). Whole-body CoP location is often calculated as an important outcome
73 variable in clinical balance studies (5, 7, 8). If bilateral force plates are used, CoP can be calculated
74 separately for each foot (e.g., as it is in instrumented treadmill studies (9)). Since the CoP of each foot
75 must be located within its boundaries, the mediolateral distance between them must be considerably
76 associated with the stance width between the heels during bipedal standing.

77 We used the approach suggested by Bland and Altman (10) to assess agreement between stance width
78 estimated from foot CoP and measured kinematically in neurotypical individuals (NT) and in
79 parkinsonian individuals in the ON (PD-ON) (8) and OFF (PD-OFF) (11) medication states. We
80 quantified the bias and expected range of differences associated with using stance width estimates from
81 foot CoP rather than kinematic measures. Then, we tested whether differences between methods were

82 associated with group membership (NT vs. PD-ON vs. PD-OFF), and whether differences varied with
83 stance width (12).

84

85 **2. Materials and Methods**

86 **2.1 PARTICIPANTS**

87 We used baseline measurements from a convenience sample of participants in previous (3) and
88 ongoing cohort studies investigating the effects of rehabilitation on balance responses (Table 1). PD
89 participants were mild-moderate with bilateral symptoms (Hoehn and Yahr stage 2-3 (13)). All
90 participants provided written informed consent and all study procedures were approved by Institutional
91 Review Boards at the Georgia Institute of Technology and Emory University.

92 **2.2 EXPERIMENT**

93 As in previous studies (3, 14), participants stood barefoot on two laboratory-grade force plates
94 (AMTI-OR6-6-1000, AMTI, Watertown, MA, USA). The force plates were mounted onto a custom
95 translation platform; however, analyses here considered only periods during which the platform was
96 stationary. Force and moment data were sampled at 1080 Hz and used to calculate the locations of the
97 center of pressure beneath each foot using calibration values supplied with the plates (15-17). Kinematic
98 data were collected at 120 Hz using a Vicon motion capture system (Centennial, CO, USA) and a 25-
99 marker set including reflective markers placed on the left and right heels. Average foot CoP locations and
100 heel marker positions were calculated over the first 250 ms of each trial.

101 Stance width was controlled by requesting participants press an object (typically a book) between the
102 medial surfaces of their feet, which was subsequently removed before data collection ($\approx 87\%$ of trials), or
103 by manipulating participant's feet so that kinematic markers on the heels were aligned in the mediolateral
104 direction with tape marks on the floor ($\approx 13\%$).

105 2.3 DATA ANALYSIS

106 Stance width measurements derived from CoP and kinematic data were plotted against each other and
107 examined visually. After visual assessment of outliers, trials were excluded due to: 1) tension in a ceiling-
108 mounted fall arrest tether interfering with CoP calculation (17 trials in one participant), and 2) absent
109 video records preventing trial review (2 trials in one participant). After applying exclusions, 1363 trials
110 (41 – 161 per participant) were available for analysis. Stance widths were expressed in mm and
111 normalized to inter-ASIS distance.

112 Following Bland and Altman (10), correlation between the two measurements was assessed with the
113 Pearson product-moment correlation coefficient r . Differences between methods were calculated for each
114 trial and averaged across trials into a single difference value d_i for each participant. Mean values across
115 methods were calculated for each trial and averaged into a single mean value m_i for each participant. Bias
116 between the two methods was quantified as the mean difference \underline{d} (CoP – kinematic method) and the
117 standard deviation of the differences s . The limits of agreement were calculated as the range $\underline{d}-2s$ to $\underline{d}+2s$.
118 Variation of differences d_i across groups was assessed with one-way ANOVA. Associations between
119 differences d_i and mean values m_i were assessed with r (12). Data processing was performed in Matlab
120 (r2016b, The Mathworks, Natick, MA, USA). Statistical procedures were performed in SAS Studio (3.5,
121 The SAS Institute, Cary, NC, USA) and considered significant at $P = 0.05$.

122 3. Results

123 Stance widths measured from kinematic data varied between 75 – 348 mm, corresponding to 24.9 –
124 154.1% of inter-ASIS distance. CoP and kinematic stance width measurements are presented in Figure
125 1A. The two measures were strongly correlated ($r = 0.98$). The mean difference \underline{d} between methods was
126 48 mm, and the standard deviation of the differences (s) was 17 mm. Differences d_i did not vary across
127 groups ($F_{2,12}=1.81$, $P<0.21$). The limits of agreement, defined as the range $\underline{d}-2s$ to $\underline{d}+2s$ (10), was 14–83
128 mm. A “Bland-Altman plot” of the differences between the two methods d_i against their means m_i is
129 presented in Figure 1B. d_i and m_i were significantly negatively correlated ($r = -0.59$, $P<0.02$).

130 **4. Discussion**

131 Stance width is an important variable in many studies of parkinsonian (4, 5) and neurotypical (18, 19)
132 posture and balance. We found that stance width estimates from foot CoP and kinematic markers were
133 strongly linearly correlated, and that on average, measures of stance width derived from CoP were 48 mm
134 wider than those derived from kinematic markers. This bias that can be explained by the externally-
135 rotated “toe out” posture used by most participants, in which a substantial portion of the foot plantar
136 surface lies lateral to the posterior face of the heel. Overall, these results suggest that foot CoP location, a
137 commonly calculated variable in clinical biomechanics studies (5, 7, 8) can be used to approximate stance
138 width in healthy aging and in individuals with PD in the ON and OFF medication states.

139 We noted that differences between methods were non-negligible – ranging from 14 to 83 mm.
140 However, this precision is adequate to discriminate between nominal stance widths used in the literature,
141 which are typically separated by 100 mm or more (4, 18). Due to the high precision of CoP calculation
142 with laboratory force plates (2-5 mm (17)), the primary source of variability in differences is probably
143 trial-to-trial variability in weight distribution, rather than instrumentation error.

144 There are two notable limitations to this approach. First, differences between methods were highest at
145 the narrow stance widths preferred by PD subjects, a fact that should be considered carefully during study
146 design. Second, because these participants were allowed to adopt a comfortable “toe out” orientation
147 during testing, the agreement between the methods in experimental paradigms in which foot orientation is
148 enforced (e.g., parallel (4); 20° (18)) remains to be established.

149 **5. Conclusion**

150 In summary, these results suggest that: 1) it is appropriate in studies of individuals with and without
151 PD to define stance width based on CoP, and 2) when comparing results across studies with the two
152 methods, it is reasonable to assume a bias of 48 mm.

153

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164 **Competing Interests**

165 The author has declared that no competing interests exist.

166

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210 **Figure legends**

211 Figure 1. Comparison of stance width measurements from kinematic and CoP data. A: Plot of results of
212 one method (CoP, ordinate) against those of the other (kinematics, abscissa). Marker shapes designate
213 study group and participants are coded by color. B: “Bland-Altman” (10) plot of limits of agreement
214 between the two methods. The CoP method introduces an absolute bias \underline{d} of 48 mm and an expected
215 range of deviations 14-83 mm. Color and marker codes are as in part A.

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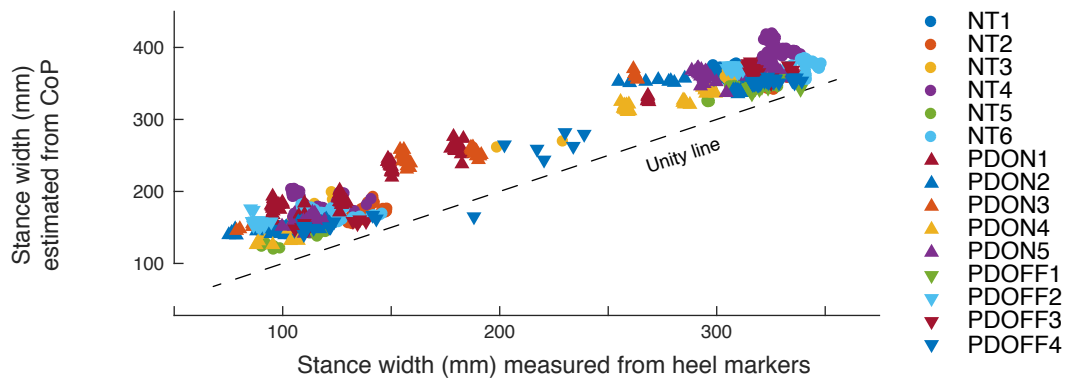
218 **Tables**

219 Table 1. Demographic, clinical, and anthropometric features of the study population.

Participant	Hoehn & Yahr Stage	Age	Sex	Height, m	Weight, kg	Inter-ASIS distance, cm	Left leg length, cm	Right leg length, cm
Neurotypical								
NT1	-	54	F	1.62	66.7	31.3	86.5	88.5
NT2	-	56	F	1.78	74.8	24.7	98.0	98.0
NT3	-	58	M	1.64	67.2	25.5	82.5	82.0
NT4	-	64	M	1.80	95.2	22.0	91.0	91.8
NT5	-	70	F	1.57	51.0	22.3	82.0	82.0
NT6	-	77	M	1.85	81.9	27.8	106.0	105.0
PD-ON								
PDON1	2	68	M	1.80	80.9	28.5	93.5	94.0
PDON2	2	69	F	1.55	74.8	30.1	84.0	83.0
PDON3	3	73	F	1.80	62.7	21.1	90.0	91.0
PDON4	2.5	79	M	1.68	68.2	27.5	91.0	90.0
PDON5	3	79	M	1.70	74.4	24.0	89.5	90.0
PD-OFF								
PDOFF1	3	75	F	1.54	50.3	22.3	83.0	82.0
PDOFF2	2	53	M	1.75	86.2	25.1	90.4	89.7
PDOFF3	2	54	F	1.63	66.0	26.2	88.0	88.0
PDOFF4	3	82	F	1.68	59.9	25.5	94.8	94.1

220 Abbreviations: ASIS, anterior superior iliac spine; PD-ON, PD participants in the ON medication
 221 state; PD-OFF, PD participants after 12+ hours of withdrawal of antiparkinsonian medications.

1A. Comparison of stance width (mm) measured from heel markers and estimated from CoP



1B. Agreement between methods

