

Comparing what the clinician draws on a digital pain map to that of persons who have greater trochanteric pain syndrome

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Published

2022

Journal Title

Scandinavian Journal of Pain

Version

Accepted Manuscript (AM)

DOI

[10.1515/sjpain-2021-0135](https://doi.org/10.1515/sjpain-2021-0135)

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1 **Comparing what the clinician draws on a digital pain map to that of persons who have**
2 **greater trochanteric pain syndrome**

3

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26

27 **Category:** Clinical Research

28

29 **Word count:** Total 3163, Abstract 248, Introduction 473, Methods 1265, Results 385,
30 Discussion 1036.

31

32 **Previous presentation of data:** IASP 2021 Virtual World Congress on Pain – Virtual
33 Duration: 9 Jun 2021 - 18 Jun 2021.

34 Plinsinga, M. L., Boudreau, S., Coombes, B., Mellor, R., Hayes, S., & Vicenzino, B. (2021).

35 Clinician Pain Drawings Differ to That of a Person with Greater Trochanteric Pain

36 Syndrome. Poster presented at IASP 2021 Virtual World Congress on Pain,

37 https://d3q7t09ulhfboj.cloudfront.net/storage/app/public/iasp/posters/documents/IASP_Plinsi

38 [nga.pdf#toolbar=0](https://d3q7t09ulhfboj.cloudfront.net/storage/app/public/iasp/posters/documents/IASP_Plinsinga.pdf#toolbar=0)

39

40 **Significance:** Patient drawings are marginally more extensive than the clinician's, and
41 differences in pain drawings are evident in location but not in the area and shape.

42 **Abstract**

43 **Objectives:** To assess the agreements and differences in pain drawings (pain area, shape and
44 location) between individuals who have greater trochanteric pain syndrome (GTPS) and their
45 clinician.

46 **Methods:** In this study, 23 patients with GTPS (21 female, pain duration range 8-24 months)
47 underwent clinical evaluation by a registered physiotherapist. Digital 2d full body pain
48 drawings were independently performed by the clinician during the subjective examination
49 and by the patient following the physical examination. Levels of agreement [LoA] in the pain
50 area were assessed with Bland-Altman plots. Differences in pain drawings were assessed
51 visually by overlaying images, and by quantifying the differences in shape and location with
52 the bounding box, and Jaccard index, respectively.

53 **Results:** Pain areas (/total pixels of the charts) did not differ in size (LoA mean difference
54 less than -0.5%; range -2.35% to 1.56%) or shape (bounding box $p > 0.17$). However, there
55 was minimal overlap in location (Jaccard index range 0.09-0.18 /1 for perfect overlap).

56 **Conclusion:** Clinicians and patients displayed differences in location of pain areas, but not
57 size or shape, when they independently performed digital pain drawings. The reasons that
58 underlie and the clinical impact of these differences remains unclear.

59

60 **Key Words**

61 Lateral hip pain, digital pain drawings, health communication, pain assessment, eHealth.

62

63 **Introduction**

64 In clinical practice of musculoskeletal pain, clinicians typically record information about the
65 size, shape and location of pain areas described by their patients on a body chart (1-4).

66 However, there is limited empirical knowledge on how this information is used to guide
67 clinical decision making and patient management.

68

69 Pain is an unpleasant sensory and emotional experience associated with, or resembling that
70 associated with, actual or potential tissue damage (5). As pain is influenced by biological,
71 psychological and social factors, there is potential for diversity in how the patient reports the
72 pain experience (6). Diversity in pain drawings can also be caused by the assessors'
73 perceptions and ability to draw verbal and physical cues on a body chart.

74

75 Although digital pain drawings are a commonly used tool, to the authors' knowledge, no
76 scientific evidence exists on *how* pain drawings are used in clinical practice. Notably, from
77 the authors' clinical experiences, not the patient, but the clinician is often the one to complete
78 the pain drawing in clinical practice. This has implications for diagnosis and treatment.

79 Specifically, differences in perceived pain distributions between clinician and patient may
80 cause misinterpretation and miscommunication at the diagnostic and treatment stage (7).

81

82

83 Digital pain drawings have been shown to enable patients and clinicians to analytically
84 understand the location, area and distribution of pain (8). Digital pain mapping is rapidly
85 evolving and demonstrates good usability, reliability, and repeatability in acute pain (9, 10)
86 and persistent pain conditions of the knee, neck and back (11-13).

87

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88 To be able to inform and strengthen clinical practice through the use of (digital) pain
89 drawings by clinicians and/or patients, we need to know the similarities and differences in
90 pain drawings between clinicians and patients. A better understanding of the similarities and
91 differences in digital pain drawings between the patient and clinician may improve
92 communication, benefit management and improve patient – clinician interaction/experience
93 satisfaction with treatment (9, 14).

94

95 Greater trochanteric pain syndrome is one of the most commonly treated tendinopathies in
96 middle-aged, overweight females (15). It is classically characterised as pain on the lateral
97 side of the hip over the greater trochanter (16). Some patients also pain in the buttocks, lower
98 back and/or distally down the leg (17). The diverse distribution of pain may reflect different
99 anatomical sources of that pain, that are identified on imaging of greater trochanteric pain
100 syndrome. Alternatively, the diversity in pain distributions may stem from varied pathologies
101 associated with greater trochanteric pain syndrome, such as hip osteoarthritis and referred
102 spinal pain (18-20), central nervous system changes (21, 22), personal characteristics (23,
103 24), or a combination of these factors. Regardless of the mechanisms underpinning the
104 diversity of clinical presentation, it is important that patients and clinicians can evaluate the
105 distribution of pain in a consistent manner.

106

107 To compare patient and clinician representations of pain experienced by individuals who
108 have persistent pain due to greater trochanteric pain syndrome, we explored the agreement
109 between full-body digital pain drawings completed by the patients and a clinician. The
110 specific aims of this study were to 1) assess the agreement between the drawn pain areas, and
111 2) assess any differences in shape and location of pain drawings. We also collated all pain
112 maps to provide an overall representation of this cohort's pain distribution.

113 **Methods**

114 **Study Design**

115 This was a cross-sectional study conducted between May 2017 and September 2018 in the
116 Brisbane metropolitan area, as part of a larger cross-sectional study (22). This study is
117 reported following the Strengthening the Reporting of Observational Studies in
118 Epidemiology (STROBE) guidelines for cross-sectional studies (25). The study was approved
119 by the University of Queensland Human Research Ethics Committee (#2015000219). All
120 participants provided verbal and written consent after being given information on the purpose
121 of the study.

122

123 **Setting**

124 All measurements were conducted between May 16, 2017, and September 11, 2018, in a
125 temperature and noise-controlled environment at the Physiotherapy Department of the
126 University of Queensland, Australia.

127

128 **Participants**

129 Individuals who had lateral hip pain were recruited from the Brisbane metropolitan area with
130 flyers and University advertisements. The sample size could not be calculated a priori in a
131 meaningful way, because there is no published data available on pain drawings between
132 patients and clinicians in greater trochanteric pain syndrome. Authors aimed to recruit as
133 many participants as possible in the available timeline (May 2017 - September 2018).
134 Eligibility for the study was assessed via online screening (MP), telephone screening
135 interview (MP) and a clinical examination conducted by a registered postgraduate qualified
136 musculoskeletal physiotherapist with >20 years of clinical experience (RM).

137

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138 Eligibility criteria for individuals who have greater trochanteric pain syndrome (referred to as
139 patients hereon) have previously been described in detail (22) – in summary included: age
140 18-70 years; lateral hip pain rating $\geq 2/10$ on an 11-point Pain Numeric Rating Scale (PNRS)
141 (0 = no pain and 10 = worst pain imaginable); pain duration \geq three months; and pain on
142 palpation of the gluteal tendon insertion on the greater trochanter plus reproduction of
143 trochanteric pain on at least one of six clinical tests (22, 26). Individuals were excluded if
144 they experienced groin pain, had received a glucocorticoid injection in the last six months,
145 experienced major lower limb trauma in the past year, had any pains that were worse than
146 their hip pain or required treatment for their hip in the past six months. They were also
147 excluded if pain was a result of any other hip joint pathology, if pregnant, or had systemic
148 inflammatory or neurological disorders, uncontrolled diabetes, and fibromyalgia.

149

150 Demographic and clinical data were collected using an online questionnaire and included age
151 (years), sex (male/female), and body mass (kg), standing height (cm), duration of symptoms
152 (months), unilateral or bilateral symptoms, the number of pain regions in the past week
153 (recorded with the Nordic Musculoskeletal Questionnaire) (27), average pain in the past week
154 (11-point PNRS), worst pain in the past week (11-point PNRS), pain during activity in the
155 past week (11-point PNRS), and disability with the Victorian Institute of Sports Assessment
156 – gluteal tendinopathy (VISA-G) questionnaire (28).

157

158 **Acquisition of Digital Pain Drawings**

159 Pain drawings were completed on a 2D digital body chart using the Navigate Pain Android
160 app (Aalborg University, v1). Patients and the clinician drew the area and location of pain on
161 a full-body chart (back, front, left and right) with a Samsung Galaxy Note accessory stylus'
162 pen on a Samsung Galaxy Note 10.1 tablet (Android 4.1.2) with a tip size of 1.5mm. The

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163 digital body charts have been validated against paper drawings in pain-free people (29) and
164 persons with other musculoskeletal pain conditions like patellofemoral pain (3).

165

166 All digital pain drawings were made at the time of the clinical examination. Clinician
167 drawings were made during the subjective interview/ physical examination, based on the
168 patient's verbal description and by physically indicating their areas of pain by using the
169 finger/s to dynamically outline and delineate the extent of the painful areas on their body.

170 After the clinical examination, and when the clinician had completed their pain drawings, the
171 clinician left the room. Another researcher (MP) replaced the clinician in the room and asked
172 the patient to draw their areas of pain. The patient was not informed about the true objective
173 of the study. Specifically, patients were instructed to "draw one or more areas of current pain
174 on the charts as accurately as possible." The patients and clinician were instructed to
175 completely shade the areas of pain and not to use outlines of the perimeter but to completely
176 shade the areas of pain. Drawings were performed independently.

177

178 **Outcome measures**

179 Raw data of the digital pain drawings were exported into Microsoft Excel, and the following
180 outcomes were extracted to compute agreement between drawings.

181

182 *Pain area*

183 The size of the total areas shaded for each view were calculated and expressed in percentage
184 of the total pixels for that view.

185

186 *Shape*

187 The bounding box was used to portray the *overall shape* of the pain drawing. It was
188 calculated by multiplying the maximum length (y) and width (x) distances of the drawn pain
189 area.

190

191 *Location*

192 The Jaccard index was used to determine the area of overlap in pain areas between clinician
193 and patient digital pain drawings. It was expressed as a proportion of the total drawn area of
194 both patient and clinician digital pain drawings. This is visualised in Fig. S1. The Jaccard
195 index is a number between 0 (no overlap) and 1 (complete overlap) of the patient and
196 clinician drawings. All metrics were automatically extracted from the pain drawings using
197 MATLAB® (Version R2017b, Natick, MA, USA) for the analysis.

198

199 *Overlay Images*

200 Overlay images were used to visually describe and inspect differences in body charts between
201 patients. Overlay images were created by superimposing the original pain drawings of all the
202 patients onto each other. The overlay images show the most frequently reported location of
203 pain using a red colour-scheme, reflecting the original drawing colour, and consists of a
204 linearly increasing pink (frequency of one) to dark brown (highest frequency) as described
205 previously by Boudreau et al (8). The colour legend shows the percentage of patients who
206 reported pain in a particular area.

207

208 **Statistical methods**

209 Results were summarised as means (standard deviations) and median (interquartile range)
210 where appropriate. The level of agreement in the pain area (expressed in percentage of total
211 pixels) was presented through Bland-Altman plots with mean differences and limits of

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212 agreement (LoA). Fixed and proportional biases were used to reflect any systematic
213 disagreements between patient and clinician drawings. Fixed bias was represented by the
214 mean of the difference in pixel density from the Bland-Altman plot and proportional bias by a
215 Pearson Correlation Coefficient (PCC) between the difference in pain area and the mean of
216 pain area from the plot (high PCC mean more bias with increased pain area). Pearson
217 correlation coefficients of (negative) 0.0 to 0.3 were regarded as negligible, 0.3 to 0.5 low,
218 0.5 to 0.7 moderate, 0.7–0.9 high, and 0.9 to 1.0 very high proportional bias. Differences in
219 the bounding box(x, y) of patient and clinician drawings were compared with a 2-sided paired
220 t-test. The Jaccard index was presented as a mean (95% confidence interval (CI)) and was
221 compared to 0 with a 1-sample t-test to test the null hypothesis of no differences between
222 clinician and patient drawings reflected by a 100% overlap in drawn areas (Jaccard index =
223 1). All statistics were performed in SPSS (Version 27.0, IBM Statistics, New York). The
224 level of significance was set as $p < 0.05$.
225

226 **Results**

227 **Participants**

228 Out of 535 volunteers, 260 underwent telephone screening, 105 underwent clinical
229 examination, and 23 patients with greater trochanteric pain syndrome were included (22). In
230 total, 23 digital pain drawings were completed by all 23 patients and the clinician.

231 Demographics of our study population can be found in Table 1. Consistent with the greater
232 trochanteric pain syndrome population, the majority of patients were female (n=21). Fig. S2
233 shows examples of individual patient-clinician charts, including charts with minimal
234 differences, charts with large differences, and charts closest to the mean difference in the
235 area.

236

237 **Similarities in area between pain drawings using Bland-Altman plots**

238 The Bland-Altman plots comparing patient and clinician drawings showed a mean difference
239 between -0.45% and -0.12% of the total pixel counts for all charts. Mean differences (LoA)
240 showed systematic, negative fixed biases for the back (-0.45%; LoA -2.36%, 1.56%), front (-
241 0.32%; LoA -1.08%, 0.44%), left (-0.28%; LoA -2.11%, 1.56%), and right (-0.12%; LoA -
242 1.49%, 1.24%) full body drawings, indicating that clinician pain areas were marginally
243 smaller compared to patients' (Fig. 1) (30). A high proportional bias was found for the front
244 charts (PCC -0.88, $p < 0.001$). This indicates that differences between patient-clinician pain
245 drawings for the front charts (e.g., fixed negative biases) increases with larger areas of pain.
246 Low and negligible proportional biases were observed for the left (PCC -0.37), back (PCC -
247 0.11), and right (PCC 0.06) charts (all $p > 0.09$). but not for back, left and right charts.

248

249 **Differences between pain drawings in shape (bounding box) and location (Jaccard**
250 **index)**

251 Overlays illustrating the pain distribution for the female patients (n=21) and the clinician
252 drawings are shown in Fig. 2. Visual inspection of the location and overall pain area suggest
253 minor differences in location and shape of patient and clinician drawings.

254
255 No differences in the bounding box(x,y) between participant and clinician drawings were
256 found for any of the chart views (all $p > 0.17$, Table 2). The mean Jaccard indexes were 0.09
257 (95% CI 0.04, 0.16) for the back view, 0.11 (95%CI -0.01, 0.23) for the front, 0.15 (95% CI
258 0.01, 0.29) for the left and 0.18 (95% CI 0.00, 0.35) for the right view (all $p < 0.05$), indicating
259 the overlap of a patient and clinician drawings were very small compared to the sum total of
260 the areas drawn.

261

262 **Discussion**

263 Our findings suggest differences in *location of pain drawings* reflected by a minimal overlap
264 in patient and clinician drawings based on the Jaccard index, but do not suggest differences in
265 the *area and shape* of the pain drawings reflected by the Bland-Altman plots and bounding
266 box respectively.

267

268 Three other studies have reported similarities in patient and clinician pain drawings; two with
269 the Jaccard index (9, 31), and one by categorizing drawn pain areas (32) (see Table S1 for a
270 Table of comparison). The Jaccard index measures the overlap between two drawings (Fig.
271 S1), and our study found a mean overlap in drawn areas ranging from 9%-18%. To our
272 knowledge, no clinical important difference for the Jaccard index exists, although an overlap
273 between of less than 20% seems very minimal. Two studies that have used the Jaccard index
274 to compare overlap between patient and clinician drawings, finding Jaccard indexes ranging
275 from 19-22% (Table S1) (31). In our study, lateral images had Jaccard index values of 15-
276 18%, which seem in line with the above studies (Table S1). A minimal overlap could be
277 explained by the clinician being more focussed on the treatment area (in our case the lateral
278 hip), placing the drawing in a slightly different location, and by potentially dismissing other
279 areas of pain that the patient focuses on, which is supported by the individual drawings
280 shown in Fig. S2. Jaccard indexes of the back and front full-body charts in our study were
281 much lower (9% and 11%, respectively), which may be explained by the fact that different
282 pain conditions were mapped, that the clinical examination was focused on the assessment of
283 greater trochanteric pain syndrome, and by the timing of the drawings. Another study
284 reported accuracy in pain drawings of 49% between full-body paper drawings of 36 chronic
285 pain patients and their doctor, analysed as drawings being either "same" or "different" (32).
286 Differences in findings may have been influenced by many factors, including the type of

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287 patients (acute pain (9) versus non-acute/persistent pain (31, 32)), electronic versus paper
288 drawings, experience in completing pain drawings, the ability to perceive and draw pain, the
289 type of charts and the instructions given to patients and clinicians (Table S1). These factors
290 highlight the importance for clinicians and patients to communicate differences and
291 similarities in their pain drawings to ensure information is not lost or changed.

292

293 In our study, clinician drawings were, on average, smaller compared to patient drawings. We
294 could speculate that this negative fixed bias could result from patients drawing their current
295 areas of pain, whereas the clinician projected a potential diagnosis. In this study, the bias may
296 have been exaggerated because the clinician made drawings during the clinical examination
297 and the patient did theirs after the clinical examination. The patients plausibly had been cued
298 into their areas of pain after the physical examination as the clinical examination was
299 focussed on stressing (provoking pain) the associated structures involved in greater
300 trochanteric pain syndrome as well as those likely to refer to the hip region. The differences
301 in pain drawings may reflect mis-communication between the patient and clinician, although
302 differences like negative fixed biases (e.g. smaller clinician drawings compared to patient
303 drawings) are to be expected when people perform tasks on unfamiliar devices (33). In this
304 study, all patients and the clinician were instructed on how to complete the drawing but were
305 unfamiliar with the device and/or application. One study revealed that fixed biases could be
306 overcome by repeated use (33), and therefore future research should incorporate attempts to
307 familiarise patients and clinicians with the device and technology before further investigating
308 these biases.

309

310 The use of clinician and patient drawings may be a beneficial tool in clinical practice.

311 Shaballout et al. (9) studied pain drawings in an acute pain setting. They had patients and

312 clinicians independently draw the pain on charts and then discussed both drawings during the
313 consultation. This study showed that doctors felt they had improved understanding of the
314 patient's pain and to a lesser extent this impacted on their clinical decisions, even though
315 drawings showed fair to good similarity (9). An earlier study by Cummings et al. (32)
316 showed that patient drawings alone are insufficient in clinical decision making and that a
317 shared-decision approach to communication between patient and clinician over the pain
318 distributions is desirable (32). Further, in clinical practice it is often the clinician – not the
319 patient – that completes the pain drawings. Our findings of differences between patient and
320 clinician pain drawings reinforces the notion that at the least, the patient should be making
321 the drawing, and perhaps both the patient and clinician should be completing pain drawings
322 as suggested by Shallabout et al. (7) and Cummings et al. (31). This may facilitate
323 conversation between clinician and patient and subsequently improve diagnosis and
324 treatment. Further research should examine if allowing the patient and clinician to discuss
325 their pain maps does indeed lead to better clinical decisions and improved care. This
326 approach is supported in other aspects of health care that show detailed communication
327 between patient and clinician about the clinical examination and associated tests (e.g.,
328 images, blood tests) will improve quality of care (34).

329

330 **Strengths, limitations, future research**

331 This study has several limitations that need to be kept in mind when inferring more broadly,
332 they are: (i) the small number of patients (n=23), (ii) only one clinician, (iii) a single study
333 site, and (iv) patients completed their pain drawing after a full clinical examination whereas
334 the clinician completed their pain drawing early in the examination at the interview. The
335 extent to which these factors impact the generalisability of our findings is unknown. To
336 overcome these limitation future studies would benefit from a larger sample of patients and

337 more than one clinician – providing greater estimates of any biases within our data. Future
338 studies that utilise more extensive data sets of digital pain drawings also provide an
339 opportunity to involved advanced statistical modelling and machine learning methods (29) as
340 clinician support tools during consultations with patients.

341

342 **Conclusions**

343 Our findings suggest that clinicians and patients will draw somewhat different pain
344 distributions, with differences in location but not in the area or shape of the digital pain
345 drawings. Perceived patient-clinician inconsistencies the location of pain may influence
346 decision-making and subsequent management, and therefore should be acknowledged and
347 addressed when using (digital) pain drawings in clinical practice. The impact on the decision-
348 making and management of patients remains to be determined.

349

350

351

352 **Acknowledgements:** The authors would like to thank Albert Cid Royo for helping with the
353 pain map overlays.

354 **Research funding:** This study was supported by the National Health and Medical Research
355 Council (NHMRC) Program Grant (#631717) (BV) and the Talent Management Grant,
356 Aalborg University (SAB). MLP was supported by the International Postgraduate Research
357 Scholarship (IPRS)/University of Queensland Centennial Scholarship (UQcent).

358 **Author contributions:** Melanie Plinsinga: conceptualization, methodology, data collection,
359 data-analysis, writing – first draft; Shellie Boudreau: conceptualization, methodology, data-
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361 editing; Rebecca Mellor: methodology, data collection, review & editing; Sandi Hayes: data-
362 analysis, review & editing; Bill Vicenzino: conceptualization, methodology, data-analysis,
363 review & editing. *All authors have accepted responsibility for the entire content of this*
364 *manuscript and approved its submission.*

365 **Competing interests:** Authors state no conflict of interest.

366 **Informed consent:** Informed consent has been obtained from all individuals included in this
367 study.

368 **Ethical approval:** Research involving human subjects complied with all relevant national
369 regulations, institutional policies and is in accordance with the tenets of the Helsinki
370 Declaration (as amended in 2013) and has been approved by the authors' Institutional Review
371 Board (The University of Queensland Human Research Ethics Committee #2015000219).

372

373

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480 **Figure legends**

481

482 **Fig. 1.** Bland-Altman plots showing the level of agreement in pain area between patient and
483 clinician pain drawings for full-body views of the (A) back, (B) front, (C) left side, (D) right
484 side. The limits of agreement are marked with dashed lines.

485

486 **Fig. 2.** Visual comparison of patient (left column) and clinician (right column) drawings for
487 female full-body charts of (A) the back, (B) front, (C) left and (D) right views (n=21). Men
488 overlay images are presented separately to the females, because the male and female avatars
489 are unique and could thus not be overlaid (Fig. S3 for male overlay images).

490

491 **Table 1.** Characteristics of the 23 patients who had greater trochanteric pain syndrome
 492 participants. Data represented as mean (Standard Deviation) unless otherwise indicated.

Characteristics	Patients
Age years	50 (10)
Women n, (%)	21 (91%)
Body mass kg/m ²	28.89 (7.37)
Median (Q1-Q3) duration of symptoms months	12 (8-24)
Unilateral n, (%)	14 (61%)
VISA-G /100	60.36 (9.14)
Median pain severity in past week (Q1-Q3)	
Average PNRS /10	4 (3-5)
Worst PNRS /10	6 (5-7)
During activity PNRS /10	5 (5-8)
Median number of pain regions (Q1-Q3)	2 (2-4.25)

493 Abbreviations: PNRS, pain numeric rating scale; Q1, first quartile; Q3, third quartile.

494

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495 **Table 2.** Mean differences (95% confidence intervals), p-value (2-sided paired t-test) in
 496 bounding box (BBx, y) values between patient and clinician drawings (mean (Standard
 497 Deviation)).

Bounding Box	Patient	Clinician	Difference
Back view			
BBx	144.91 (100.75)	116.48 (96.83)	-28.44 (-70.21, 13.24), 0.17
BBy	236.70 (226.20)	222.04 (228.44)	-14.65 (-133.15, 103.84), 0.80
Front view			
BBx	110.74 (111.04)	76.48 (97.76)	-24.26 (-85.57, 17.05), 0.18
BBy	184.83 (175.81)	123.04 (167.39)	-61.78 (-153.10, 29.53), 0.18
Left view			
BBx	68.61 (56.58)	63.65 (46.03)	-4.96 (-24.33, 14.42), 0.60
BBy	188.00 (210.77)	137.91 (168.45)	-50.09 (-147.99, 47.82), 0.30
Right view			
BBx	81.74 (58.92)	72.35 (49.43)	-9.39 (-37.49, 18.70), 0.50
BBy	170.04 (168.39)	152.17 (192.14)	-17.87 (-118.24, 82.50), 0.72

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