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Short-Term Effect of Spinal Manipulation on Pain Perception, Spinal Mobility, and Full Height Recovery in Male Subjects with Degenerative Disc Disease: A Randomized Controlled Trial

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Running Head: Disc Disease after Spinal Manipulation

Title: Short-Term Effect of Spinal Manipulation on Pain Perception, Spinal Mobility, and Full Height Recovery in Male Subjects with Degenerative Disc Disease: A Randomized Controlled Trial

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No reprints are available for this study.

The study protocol was approved by the Ethical and Research Committee of Faculty of Dom Bosco, Curitiba, Paraná, Brasil, with registration number CAAE 0002.0.301.000-11. The study has been registered in the Australian and New Zealand Clinical Trial Registry with registration number ACTRN12613000430730.

1 **Title:** Short-Term Effect of Spinal Manipulation on Pain Perception, Spinal Mobility,  
2 and Full Height Recovery in Male Subjects with Degenerative Disc Disease: A  
3 Randomized Controlled Trial

4

#### 5 **Abstract**

6 **Objective:** To evaluate the short-term effect on spinal mobility, pain perception,  
7 neural mechanosensitivity, and full height recovery after high-velocity low-amplitude  
8 (HVLA) spinal manipulation (SM) in the lumbosacral joint (L5-S1)

9 **Study Design:** Randomized, double-blind, controlled clinical trial with evaluations at  
10 baseline and after intervention

11 **Setting:** University-based physical therapy research clinic

12 **Participants:** Forty male subjects (N=40) (mean age $\pm$  SD; 38  $\pm$  9.14 years)  
13 diagnosed with degenerative lumbar disease at L5-S1 were randomly divided into  
14 two groups: the treatment group (TG) (n = 20) (39  $\pm$  9.12 years) and control group  
15 (CG) (n = 20) (37  $\pm$  9.31 years). All participants completed the intervention and  
16 follow-up evaluations

17 **Interventions:** A single L5-S1 SM technique (pull-move) was performed in the TG,  
18 whereas the CG received a single placebo intervention

19 **Main Outcome Measures:** Measures included assessing the subject's height using  
20 a stadiometer. The secondary outcome measures included perceived low back pain,  
21 evaluated using a visual analogue scale; neural mechanosensitivity, as assessed  
22 using the passive straight leg raise test (SLR); and amount of spinal mobility in  
23 flexion, as measured using the finger to floor distance test (FFD)

24 **Results:** The intra-group comparison indicated a significant improvement in all  
25 variables in the TG ( $p < .001$ ). There were no changes in the CG, except for the FFD

26 (p=.008). In the between-group comparison of the mean differences from pre- to  
27 post-intervention, there was statistical significance for all cases (p<.001)

28 **Conclusions:** An HVLA SM in the lumbosacral joint performed on male subjects  
29 with degenerative disc disease immediately improves self-perceived pain, spinal  
30 mobility in flexion, hip flexion during the passive SLR, and subject's full height.  
31 Future studies should include female subjects and should evaluate the long-term  
32 results

33  
34 **Keywords:** Intervertebral disc degeneration; Lumbar disc disease; Spinal  
35 manipulation.

36  
37 ***List of abbreviations***

38 IVD intervertebral disc

39 LBP low back pain

40 DD disc degeneration

41 SM spinal manipulation

42 HVLA high-velocity low-amplitude

43 ROM range of motion

44 CG control group

45 TG treatment group

46 VAS visual analogue scale

47 SLR passive straight leg raise test

48 FFD finger to floor distance test

49 L5 fifth lumbar vertebra

50 Lumbar intervertebral disc (IVD) disease is one of the main causes for low-back  
51 pain (LBP) among individuals with spinal disorders, affecting approximately 16% of  
52 patients.<sup>1</sup> A total of 80% of the population of industrialized countries experience  
53 episodes of severe LBP during their lives,<sup>2</sup> with disc degeneration (DD) being the  
54 most common pathology in the adult spine and accounting for approximately 90% of  
55 surgery cases.<sup>3</sup> Subjects suffering from symptomatic disc disorders incur the highest  
56 health care expenditure among those with other LBP diagnoses.<sup>4</sup> Nevertheless, the  
57 etiology of LBP appears to be multifactorial, which makes its diagnosis and  
58 management still controversial.<sup>5</sup>

59  
60 Even though spinal manipulation (SM) has been linked with positive changes in  
61 pain central processing mechanisms,<sup>6</sup> there are conflicting reports with regard to the  
62 impact of SM on pain perception in LBP patients.<sup>7,8</sup> However, conclusions are limited  
63 by the scarce number of studies.<sup>8</sup> There appears to be some evidence of the  
64 effectiveness of high-velocity low-amplitude (HVLA) SM in lumbar IVD disorders.<sup>9</sup>  
65 SM has been demonstrated to decrease pain and improve function in symptomatic  
66 lumbar DD.<sup>10</sup> On the contrary, little is known about the neural mechanosensitivity  
67 response of the lower extremities after manipulative treatment.<sup>11,12</sup>

68  
69 The structural disruption of the IVD (loss of the hydrostatic capacity of the  
70 nucleus) during DD may end up leading to a loss of IVD height and a possible  
71 reduction of spinal range of motion.<sup>13</sup> DD, however, has also been positively  
72 correlated with segmental flexibility of the lumbar spine.<sup>14</sup> An IVD height narrowing  
73 has been associated with a history of LBP problems,<sup>15</sup> although no relation between  
74 LBP and IVD height has been concluded in other studies.<sup>16</sup> The cumulative effect of

75 the IVD loss of fluid in response to mechanical stress may change the subject's  
76 measured height (spinal shrinkage).<sup>17</sup> In addition, limitations in hip range of motion  
77 (ROM) in subjects with DD appears to be specially important because hip mobility  
78 influences the loads upon the lower back,<sup>13</sup> and reduced hip ROM may also be  
79 related to LBP.<sup>18</sup>

80

81 Conservative approaches appear to be among the best options for DD  
82 associated with LBP.<sup>13</sup> Therefore, the purpose of the study was to evaluate, in  
83 subjects with lumbar DD, the immediate effect of a lumbosacral HVLA SM on: (a) the  
84 subject's height, (b) self-perceived LBP, (c) neural mechanosensitivity, and (d) spinal  
85 mobility in flexion.

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## 88 **Methods**

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### 91 **Design**

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94 This was a controlled, randomized and double-blind clinical trial. All participants  
95 signed an informed consent form, as established by the institutional review board.

96 The study protocol was conducted according to the Declaration of Helsinki.

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### 99 **Randomization Process**

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The random sequence was obtained using the website [www.randomization.com](http://www.randomization.com),<sup>19</sup> and an outside collaborator prevented access to the sequence for those participating in the research.

### **Blinding**

Before randomization, the participants received general information about the study and were informed that there would be different techniques compared. Subjects and evaluators who collected or analyzed data remained unaware of the treatment allocation group.

### **Sample Size**

The sample size was calculated using Granmo version 7.12 software.<sup>a</sup> For a two-sided contrast and accepting an  $\alpha$  value of .05 and a beta risk of .01, eighteen subjects were required per study group to detect a difference equal to or greater than 17.5% in the between-groups comparison of the stadiometry values. A 15% SD was assumed together with an estimated 10% rate of loss to follow-up.



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**126 Study Subjects**

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129 Fifty-six (N=56) male subjects were evaluated for possible participation. Based on a  
130 non-probabilistic convenience sampling, the participants were consecutively  
131 recruited from the principal investigator's clinical consultancy. The research protocol  
132 was implemented at an university-based physical therapy research clinic from March  
133 to October 2012. Of the total number of subjects enrolled, 16 were excluded for  
134 several reasons (figure 1).The final sample included 40 male subjects ( $38 \pm 9.14$   
135 years) with a diagnosis of DD in the lumbosacral joint. The participants were  
136 randomized into two study groups: the Control Group (CG) (n=20) and the Treatment  
137 Group (TG) (n=20). No loss to follow-up was recorded during the data collection or  
138 analysis phases.

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**141 Inclusion/Exclusion Criteria**

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144 The inclusion criteria were as follows: (a) males between 18-55 years; (b)  
145 standardized body mass index (between 20-25 kg/m<sup>2</sup>); (c) imaging evidence (T<sub>2</sub>-  
146 weighted MRI) to ensure clinical diagnosis of DD, based upon the presence or  
147 absence of degeneration in the lumbosacral IVD;<sup>13,20</sup> and (d) LBP (no minimum  
148 intensity of pain was specified),<sup>21</sup> with or without pain radiating to the lower  
149 extremities above the knee, according to categories 1 and 2 of the Quebec Task

150 Force classification system.<sup>22</sup> The exclusion factors were: (a) smoking; (b) history of  
151 alcoholism or alcohol consumption within 24 hours prior to data collection; (c)  
152 professional sportsmen (changes in the IVD response mechanical parameters have  
153 been found in these subjects);<sup>23,24</sup> (d) diagnosis of median, fragmented or migrating  
154 herniation (T<sub>2</sub>-weighted MRI);<sup>13,23</sup> (e) cauda equina syndrome;<sup>25</sup> (f) general  
155 contraindications to SM;<sup>26</sup> (g) surgery for DD; (h) radicular pain and/or radiculopathy  
156 with presence of neurologic signs;<sup>22</sup> and (i) SM treatment within three months before  
157 data collection.

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#### 160 **Data Collection Protocol**

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163 Participants were subjected to the evaluation and intervention protocol together in  
164 one session that lasted approximately an hour. The intervention was conducted  
165 three minutes after the assessment, and the re-evaluation process began three  
166 minutes after the intervention. The therapist in charge of the intervention had over 8  
167 years of clinical experience in the field of manual therapy. The pre-intervention data  
168 collection protocol was conducted in the order stated below. This order was  
169 maintained after the intervention, apart from the stadiometer measurement, which  
170 was performed first in the post-intervention assessment.

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#### 173 **Outcome Measures**

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**176 Evaluation of Self-Perceived LBP**

177 LBP was measured with a visual analogue scale (VAS). The VAS consists of a  
178 horizontal 100-mm line, which ranges from 0 mm (no pain) to 100 mm (severe pain),  
179 where the subjects mark their perceived pain.<sup>27</sup> The VAS is an effective, sensitive  
180 and appropriate tool to measure acute and chronic pain.<sup>27</sup> The subjects were asked  
181 about the current intensity of pain.<sup>28,29</sup>

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**184 Passive Straight Leg Raise Range of Motion**

185 Neural mechanosensitivity was observed by means of the passive straight leg raise  
186 test (SLR).<sup>30</sup> The initial appearance of pain or discomfort was the test end point.<sup>31</sup> In  
187 this position, a goniometer<sup>b</sup> was used to measure the hip flexion ROM. The lower  
188 limb that presented radiating pain was chosen to be assessed. In cases where there  
189 was only midline LBP or equally radicular pain, the SLR ROM from the lower limb  
190 with a worse performance was taken as the reference value. Among other  
191 considerations, the SLR is “positive” when there is identified asymmetry between  
192 lower extremities.<sup>30</sup> The SLR is considered an easy-to use tool, with a reliability  
193 (ICC) of 0.87 (95% CI: 0.69 - 0.95).<sup>30</sup>

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**196 Spine Mobility in Flexion (Finger to Floor Distance Test)**

197 Spinal mobility was assessed using the finger to floor distance test (FFD). The FFD  
198 evaluates the maximum spinal mobility in flexion, and it is a possible indicator of  
199 functional limitation.<sup>32</sup> The FFD was conducted according to the established

200 methodology.<sup>33</sup> This test is considered easy to conduct and has a high degree of  
201 interexaminer reliability ( $r = 0.96$  to  $0.98$ ).<sup>34</sup>

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203

#### 204 **Stadiometer**

205 The stadiometer measures height variations and the amount of IVD compression  
206 caused by pressure on the spinal column.<sup>17</sup> The protocol described by Rodacki et  
207 al<sup>24</sup> was followed. The participants were considered sufficiently trained when after  
208 five consecutive evaluations, the measurements displayed a SD of less than 0.5  
209 mm.<sup>17,24</sup> The person's height was measured after 90 seconds standing upright in the  
210 stadiometer<sup>c</sup> to allow body structures to reach their equilibrium.<sup>24</sup> To prevent postural  
211 adjustments, fixing metal bars were placed on different anatomical points. The  
212 subject also wore safety glasses with a leveling system to stop head movements.  
213 The measuring stick of the digital transducer<sup>d</sup> was positioned on the center of gravity  
214 above the head. The subject remained in the stadiometer at all times during  
215 measurements, instead of using the "in-out" method.<sup>35</sup> The stadiometer is a  
216 noninvasive method that has proven validity<sup>36</sup> and is easier to use and less costly  
217 than MRI.<sup>37</sup>

218

219

#### 220 **Interventions**

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#### 223 **SM in the Treatment Group**

224 The SM technique (pull-move) was performed following previous research  
225 reports.<sup>38,39</sup> The participant was in a side-lying position. The upper body was turned  
226 to introduce rotation and lateral flexion into the lumbosacral spine. Then, the  
227 therapist moved the area of counter-rotation to the segment to be manipulated  
228 (figure 2). The thrust was performed very fast within a short ROM.<sup>38,39</sup>

229

230

### 231 **Placebo Manoeuvre (sham) in the Control Group**

232 In the CG, after placing the subject in side-lying position with hip and knee flexion, no  
233 mechanical tension was added, as no turning of the upper body nor manipulative  
234 thrust were delivered. The position was maintained during the same time as  
235 estimated for the TG.

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237

### 238 **Data Analysis**

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240

241 The descriptive and inferential analysis of the results was performed using the  
242 BioEstat 5 free software program.<sup>e</sup> The mean, SD and 95% confidence interval were  
243 calculated for the different variables. The statistical analysis was conducted  
244 considering significant at a p value < .05. The D'Agostino test evaluated the  
245 normality of the study variables. Only the self-perceived LBP followed a non-normal  
246 distribution (p>.05). The comparison between-groups used the Student t-test for the  
247 quantitative variables, and the Chi-square (X<sup>2</sup>) for the categorical variables. In the  
248 intra-group comparison, the Student t-test was used to analyze the parametric

249 dependent variables, whereas the Mann-Whitney U test was used for the  
250 nonparametric variable. The analysis of variance for repeated measures (ANOVA  
251 test) with the group allowed the inter-group differences to be observed. The  
252 correlation test (Pearson or Spearman) was used to assess the association between  
253 extraneous variables and the dependent variables.

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255

## 256 **Results**

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258

259 The baseline results in regard to the clinical aspects and the outcome measures are  
260 included in table 1. No significant differences in the inter-group comparison were  
261 found ( $p>.05$ ), except in the case of the SLR ( $p=.004$ ).

262

263 Table 2 lists the pre- and post-intervention values and the analysis of the intra-  
264 group changes. All the study variables displayed a significant improvement in the TG  
265 ( $p<.001$ ). On the contrary, there were no intra-group differences in the CG ( $p>.05$ )  
266 except for the FFD ( $p=.008$ ).

267

268 Table 3 reports the between-groups comparison of the mean score changes  
269 after intervention ( $p<.001$  in all cases). The between-group analysis indicated  
270 significance in the case of LBP ( $p<.001$ ;  $F_{1,38}=21.03$ ;  $R^2=0.35$ ), hip flexion ROM  
271 during the SLR ( $p<.001$ ;  $F_{1,38}=50.05$ ;  $R^2=0.56$ ), flexion mobility in the FFD ( $p<.001$ ;  
272  $F_{1,38}=47.63$ ;  $R^2=0.55$ ) and in the stadiometry ( $p<.001$ ;  $F_{1,38}=145.05$ ;  $R^2=0.79$ ).

273

274 In the correlation analysis, a negative correlation was found between the stature  
275 recovery and the increase in mobility during the FFD ( $p<.001$ ;  $r=0.656$ ), as well as  
276 between the height gained and the perceived pain relief ( $p=.001$ ;  $r=-0.499$ ). Similarly,  
277 the changes in the stadiometry positively correlated with improvements in the SLR  
278 ( $p<.001$ ;  $r=0.537$ ).

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280

## 281 Discussion

282

283

284 The lumbosacral SM achieved an immediate reduction in self-perceived LBP.  
285 The minimum important difference is defined as the smallest variation in the outcome  
286 in the domain of interest, indicating meaningful change in clinical status.<sup>40</sup> For VAS, it  
287 has been reported to vary from 20 mm in chronic LBP<sup>41,42</sup> to 35 mm in acute or  
288 subacute LBP.<sup>42</sup> Licciardone et al<sup>43</sup> concluded that a substantial LBP improvement  
289 after SM needs to represent a change  $\geq 50\%$  in regard to VAS score at baseline.  
290 Pain perception decreased by  $17 \pm 16.57$  mm in the TG, which represents a 45.94%  
291 improvement in relation to baseline (table 3). Although the results were close to  
292 clinical significance, they must be cautiously interpreted because subjects were most  
293 likely at different stages of LBP.

294

295 The SM was delivered to the lumbosacral spine to “open” (gap) the targeted  
296 joint. The side-lying position combined with SM appears to be beneficial, both for  
297 pain reduction and zygapophyseal joint gapping, in subjects with acute LBP.<sup>44</sup>  
298 Nonetheless, although SM has been related to a short-term pain relief in LBP,<sup>43</sup> there

299 is still controversy regarding this aspect. On the one hand, SM has demonstrated an  
300 impact on the central control mechanism and pain regulation.<sup>45</sup> On the other hand, a  
301 recent systematic review concluded that there is low-quality evidence to support that  
302 SM is more effective than sham treatment concerning pain relief.<sup>46</sup> This lack of  
303 evidence is linked to poor quality methodology in many cases.<sup>47</sup> Most studies on SM  
304 lack previous estimates of sample size and a control group.<sup>47</sup> This clinical trial has  
305 taken into account these aspects to increase internal validity.

306  
307 In regards to the SLR, the hip flexion ROM significantly increased in the TG,  
308 compared with the sham intervention. Szlezak et al<sup>48</sup> assessed the immediate effect  
309 of lumbar mobilizations in healthy subjects and found a significant improvement in  
310 the SLR (mean increase of 8.50°). They explained their findings as a consequence  
311 of a possible change in the neurodynamics of lower extremity posterior muscles and  
312 neural structures. The results of the present study in subjects with DD were better  
313 after SM at L5-S1 ( $13.65^\circ \pm 8.62^\circ$ ) (table 3) and surpassed the minimal detectable  
314 change reported for the SLR in LBP patients (5.7°-6.6°).<sup>49</sup> Therefore, an  
315 improvement in the mechanosensitivity of the nervous system appears to be a  
316 plausible reason to understand the observed phenomenon.<sup>30</sup> In this sense, SM has  
317 been linked to short-term inhibitory effects on the human motor system.<sup>50</sup> Sensitizing  
318 maneuvers (ankle dorsiflexion and/or neck flexion) are needed to elucidate that the  
319 limitation during SLR is certainly related to the neural system.<sup>51</sup> In the present study,  
320 SLR was performed with pelvis supported at rest and the ankle and neck in neutral  
321 position with no sensitizing movements added. Another possible explanation for the  
322 findings is a change of distal muscle tone and activity, which has been perceived as  
323 a protective reflexive mechanism to prevent strain of the nerves.<sup>30</sup> Even though it



324 remains controversial, it has also been concluded that HVLA lumbosacral SM  
325 displays a short-term impact on the attenuation of alpha motoneuronal activity.<sup>50</sup>  
326 This seems to be linked to a reduction of muscle tone and pain perception.<sup>50</sup>  
327 However, these aspects (tone and muscle activation) were not measured and  
328 controlled in the study. Though the SLR response was positive in the evaluated  
329 lower limb, future studies should extend the observations to both lower limbs.

330

331 The spinal mobility during FFD was also increased in the TG. SM modulates the  
332 somatosensory system, which inhibits the paravertebral muscle hyperactivity and  
333 improves spine functionality, among other effects.<sup>52</sup> As stated before, the SM  
334 contributed to enhanced hip flexion mobility with a possible impact on FFD. Previous  
335 studies concluded that the mean difference in the FFD after intervention should be  
336 greater than 4.5 cm<sup>49</sup> or 10 cm<sup>53</sup> for the result to have clinical significance and  
337 predict improvement in disability. The FFD increased an average of  $3.67 \pm 2.09$  cm  
338 in the TG (table 3), and only one of the TG subjects improved above 10 cm.  
339 Therefore, the results cannot be considered as clinically relevant. Other factors that  
340 may influence the FFD have not been taken into account, such as hip ROM, pelvic  
341 alignment at the hip and hamstring and/or calf muscles tension.<sup>32</sup>

342

343 FFD also improved in the intra-group analysis in the CG ( $p=.008$ ) (table 2), which  
344 underwent a sham intervention with the subject placed in a side-lying position. As  
345 suggested, paraspinal muscles may relax during maintained side-posture  
346 positioning,<sup>44</sup> with the results previously referred to the FFD. However, only the  
347 mechanical effect from the thrust appears to be a key element to the effectiveness of  
348 the SM.<sup>38,45</sup> There were only a few minutes between pre- and post-intervention

349 measurements. Therefore, it is also possible that the post-intervention evaluation  
350 could have just improved as a consequence of the learning process from the pre-  
351 intervention assessment. To increase the internal validity and predictive value of  
352 studies, the combination of FFD and SLR has been suggested as the best option in  
353 evaluating subjects with acute/subacute LBP.<sup>49</sup>

354

355 A significant height change was found in the TG, even though the mean absolute  
356 amount of increase was very small ( $3.98 \pm 1.46$  mm, table 3). Similar height recovery  
357 was observed after superficial heat treatment in LBP patients ( $4.2 \pm 2.4$  mm)<sup>54</sup> but  
358 also after sustained lumbar flexion and extension postures in pain-free participants  
359 (between 3.15 mm – 5.84 mm).<sup>37,55</sup> In the latter studies, measurements were made  
360 with the subject in the seated position. In addition, age, which seems to be linked to  
361 DD progression,<sup>5,13</sup> and may influence the stadiometry results,<sup>24</sup> was different among  
362 study samples. Thus, it is difficult to compare between studies.

363

364 The increase in paravertebral muscle activity in chronic LBP has been  
365 associated with greater compressive loads and a lower possibility of height  
366 recovery,<sup>56</sup> although this correlation remains just as an hypothesis.<sup>57</sup> Nevertheless,  
367 the inverse process may be inferred. The impact of SM on “gapping”<sup>44</sup> and on  
368 diminishing paravertebral hyperactivity<sup>52</sup> may produce changes in stadiometry. It  
369 remains uncertain if a single SM in a single spinal joint (L5/S1) is sufficient to explain  
370 the immediate height recovery. SM effects are not isolated to the targeted level, as  
371 lumbar SM is just specific and accurate half the time.<sup>58</sup> Likewise, standing posture is  
372 also dependent on other factors, such as pelvic alignment at the hip and hamstring  
373 tension, among others, which were not controlled.

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375

**376 Study Limitations**

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379 The subjects were not evaluated for LBP duration. Thus, they might be at different  
380 stages of LBP. The research only included male subjects and assessed short-term  
381 effects. It would be of great interest to determine whether there are gender-related  
382 effects. Future studies should include males and females and medium- to long-term  
383 results. Furthermore, fear-avoidance beliefs, which have been correlated with lumbar  
384 flexion in LBP, have not been evaluated.<sup>59</sup>

385

386

**387 Conclusions**

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390 A side-lying SM technique in the lumbosacral region decreases self-perceived LBP  
391 in the short term and produces an immediate improvement in spinal mobility in  
392 flexion, the subject's height and hip flexion mobility during the SLR in male subjects  
393 with DD.

394

**395 References**

396

- 397 1. Martin BI, Deyo RA, Mirza SK, Turner JA, Comstock BA, Hollingworth W,  
398 Expenditures and health status among adults with back and neck problems.  
399 JAMA 2008;299:656-64.
- 400 2. Larivière C, Gangnon D, Loisel P. A biomechanical comparison of lifting  
401 techniques between subjects with and without chronic low back pain during  
402 freestyle lifting and lowering tasks. Clin Biomech 2002;17:89-98.
- 403 3. An HS, Anderson PA, Houghton VM, Iatridis JC, Kang JD, Lotz JC, Natarajan  
404 RN, Oegema TR Jr, Roughley P, Setton LA, Urban JP, Videman T, Andersson  
405 GB, Weinstein JN. Introduction: disc degeneration: summary. Spine (Phila Pa  
406 1976) 2004;29:2677-8.
- 407 4. Luo X, Pietrobon R, Sun SX, Liu GG, Hey L. Estimates and patterns of direct  
408 health care expenditures among individuals with back pain in the United States.  
409 Spine (Phila Pa 1976) 2004;29:79-86.
- 410 5. Negrini S, Zaina F. The chimera of low back pain etiology. Am J Phys Med  
411 Rehabil 2013;92:93-7.
- 412 6. Schmid A, Brunner F, Wright A, Bachmann LM. Paradigm shift in manual  
413 therapy? Evidence for a central nervous system component in the response to  
414 passive cervical joint mobilisation. Man Ther 2008;13:387-96.
- 415 7. Kuczynski JJ, Schwieterman B, Columber K, Knupp D, Shaub L, Cook CE.  
416 Effectiveness of physical therapist administered spinal manipulation for the  
417 treatment of low back pain: a systematic review of the literature. Int J Sports  
418 Phys Ther 2012;7:647-62.
- 419 8. Rubinstein SM, Terwee CB, Assendelft WJ, de Boer MR, van Tulder MW.  
420 Spinal manipulative therapy for acute low back pain: an update of the  
421 cochrane review. Spine (Phila Pa 1976) 2013;38:E158-77.

- 422 9. Lisi AJ, Holmes EJ, Ammendolia C. High-velocity low amplitude spinal  
423 manipulation for symptomatic lumbar disk disease: a systematic review of the  
424 literature. *J Manipulative Physiol Ther* 2005;28:429-42.
- 425 10. Burton AK, Tillotson KM, Cleary J. Single-blind randomised controlled trial of  
426 chemonucleolysis and manipulation in the treatment of symptomatic lumbar disc  
427 herniation. *Eur Spine J* 2000;9:202-7.
- 428 11. Stern PJ, Côté P, Cassidy JD. A series of consecutive cases of low back pain  
429 with radiating leg pain treated by chiropractors. *J Manipulative Physiol Ther*  
430 1995;18:335-42.
- 431 12. Pollard H, Ward G. The effect of upper cervical or sacroiliac manipulation on hip  
432 flexion range of motion. *J Manipulative Physiol Ther* 1998;21:611-6.
- 433 13. Beattie PF. Current understanding of lumbar intervertebral disc degeneration: a  
434 review with emphasis upon etiology, pathophysiology, and lumbar magnetic  
435 resonance imaging findings. *J Orthop Sports Phys Ther* 2008;38:329-40.
- 436 14. Tanaka N, An HS, Lim TH, Fujiwara A, Jeon CH, Haughton VM. The relationship  
437 between disc degeneration and flexibility of the lumbar spine. *Spine J* 2001;1:47-  
438 56.
- 439 15. Battie MC, Videman T, Levalahti E, Gill K, Kaprio J. Heritability of low back pain  
440 and the role of disc degeneration. *Pain* 2007;131:272-80.
- 441 16. Videman T, Battié MC, Gibbons LE, Maravilla K, Manninen H, Kaprio J.  
442 Associations between back pain history and lumbar MRI findings. *Spine (Phila*  
443 *Pa 1976)* 2003;28:582-8.
- 444 17. Eklund JA, Corlett EN. Shrinkage as a measure of the effect of load on the  
445 spine. *Spine (Phila Pa 1976)* 1984;9:184-94
- 446 18. Porter JL, Wilkinson A. Lumbar-hip flexion motion. A comparative study between

- 447 asymptomatic and chronic low back pain in 18- to 36-year-old men. Spine (Phila Pa  
448 1976). 1997;22:1508-13.
- 449 19. Dallal GE. Randomization plan generators. Available at  
450 <http://www.randomization.com>. Accessed January 19, 2012.
- 451 20. Beattie PF, Arnot CF, Donley JW, Noda H, Bailey L. The immediate reduction in  
452 low back pain intensity following lumbar joint mobilization and prone press-ups is  
453 associated with increased diffusion of water in the L5-S1 intervertebral disc. J  
454 Orthop Sports Phys Ther 2010;40:256-64
- 455 21. Shirado O, Doi T, Akai M, Hoshino Y, Fujino K, Hayashi K, Marui E, Iwaya T.  
456 Multicenter randomized controlled trial to evaluate the effect of home-based  
457 exercise on patients with chronic low back pain: the Japan low back pain  
458 exercise therapy study. Spine (Phila Pa 1976) 2010;35:811-9.
- 459 22. Werneke MW, Hart DL. Categorizing patients with occupational low back pain by  
460 use of the Quebec Task Force Classification system versus pain pattern  
461 classification procedures: discriminant and predictive validity. Phys Ther  
462 2004;84:243-54.
- 463 23. Rodacki C, Fowler N, Rodacki A, Birch K. Stature loss and recovery in pregnant  
464 women with and without low back pain. Arch Phys Med Rehabil 2003;84:507-12.
- 465 24. Rodacki CL, Fowler NE, Rodacki AL, Birch K. Technical Note: Repeatability of  
466 measurement in determining stature in sitting and standing postures.  
467 Ergonomics 2001;44:1076-85.
- 468 25. Negrelli WF. Hérnia Discal: Procedimentos de tratamento. Acta Ortop Bras  
469 2001;9:39-45.
- 470 26. Gibbons P, Tehan P. Spinal manipulation: indication, risks and benefits. J Bodyw  
471 Mov Ther 2001;5:110-9.

- 472 27. Carlsson AM. Assessment of chronic pain. I. Aspects of the reliability and validity  
473 of the visual analogue scale. *Pain* 1983;16:87-101.
- 474 28. Paatelma M, Kilpikoski S, Simonen R, Heinonen A, Alen M, Videman T.  
475 Orthopaedic manual therapy, McKenzie method or advice only for low back pain  
476 in working adults: a randomized controlled trial with one year follow-up. *J Rehabil*  
477 *Med* 2008;40: 858-63.
- 478 29. Senna MK, Machaly SA. Does maintained spinal manipulation therapy for  
479 chronic nonspecific low back pain result in better long-term outcome? *Spine*  
480 (Phila Pa 1976) 2011;36:1427-37.
- 481 30. Boyd BS, Wanek L, Gray AT, Topp KS. Mechanosensitivity of the lower  
482 extremity nervous system during straight-leg raise neurodynamic testing in  
483 healthy individuals. *J Orthop Sports Phys Ther* 2009;39:780-90.
- 484 31. Rebain R, Baxter GD, McDonough S. A systematic review of the passive straight  
485 leg raising test as a diagnostic aid for low back pain (1989 to 2000). *Spine* (Phila  
486 Pa 1976) 2002;27:388-95.
- 487 32. Ohtsuki K, Suzuki T. A comparison of the immediate changes in subjects with  
488 chronic lower back pain effected by lower back pain exercises and direct  
489 stretching of the tensor fasciae latae, the hamstring and the adductor magnus. *J*  
490 *Phys Ther Sci* 2012;24:707-09.
- 491 33. Méndez-Sánchez R, Albuquerque-Sendín F, Fernández-de-las-Peñas C, et al.  
492 Immediate effects of adding a sciatic nerve slider technique on lumbar and lower  
493 quadrant mobility in soccer players: a pilot study. *J Altern Complement Med*  
494 2010;16:669-75
- 495 34. Horre T. Finger-to-floor distance and schober test: validity criterion for these  
496 tests? *Manuelle Ther* 2004;8:55-65.

- 497 35. Stothart JP, McGill SM. Stadiometry: on measurement technique to reduce  
498 variability in spine shrinkage measurement. *Clin Biomech* 2000;15:546-8.
- 499 36. Kourtis D, Magnusson ML, Smith F, Hadjipavlou A, Pope MH. Spine height and  
500 disc height changes as the effect of hyperextension using stadiometry and MRI.  
501 *Iowa Orthop J* 2004;24:65-71.
- 502 37. Owens SC, Brismée JM, Pennell PN, Dedrick GS, Sizer PS, James CR.  
503 Changes in spinal height following sustained lumbar flexion and extension  
504 postures: a clinical measure of intervertebral disc hydration using stadiometry. *J*  
505 *Manipulative Physiol Ther* 2009;32:358-63.
- 506 38. Kirkaldy-Willis WH, Cassidy JD. Spinal manipulation in the treatment of  
507 low-back pain. *Can Fam Physician* 1985;31:535-40.
- 508 39. Hondras MA, Long CR, Cao Y, Rowell RM, Meeker WC. A randomized  
509 controlled trial comparing 2 types of spinal manipulation and minimal  
510 conservative medical care for adults 55 years and older with subacute or chronic  
511 low back pain. *J Manipulative Physiol Ther* 2009;32:330-43.
- 512 40. Jaeschke R, Singer J, Guyatt GH. Measurement of health status. Ascertaining  
513 the minimal clinically important difference. *Control Clin Trials* 1989;10:407-15.
- 514 41. Hagg O, Fritzell P, Nordwall A. The clinical importance of changes in outcome  
515 scores after treatment for chronic low back pain. *Eur Spine J* 2003;12:12-20.
- 516 42. Ostelo RW, de Vet HC. Clinically important outcomes in low back pain. *Best*  
517 *Pract Res Clin Rheumatol* 2005;19:593-607.
- 518 43. Licciardone JC, Kearns CM, Minotti DE. Outcomes of osteopathic manual  
519 treatment for chronic low back pain according to baseline pain severity: Results  
520 from the OSTEOPATHIC Trial. *Man Ther* 2013; 18:533-40



- 521 44. Cramer GD, Cambron J, Cantu JA, Dexheimer JM, Pocius JD, Gregerson D,  
522 Fergus M, McKinnis R, Grieve TJ. Magnetic resonance imaging zygapophyseal  
523 joint space changes (gapping) in low back pain patients following spinal  
524 manipulation and side-posture positioning: a randomized controlled mechanisms  
525 trial with blinding. *J Manipulative Physiol Ther* 2013;36:203-17.
- 526 45. Pickar JG. Neurophysiological effects of spinal manipulation. *Spine J*  
527 2002;2:357-71.
- 528 46. Rubinstein SM, van Middelkoop M, Assendelft WJ, de Boer MR, van Tulder MW.  
529 Spinal manipulative therapy for chronic low-back pain: an update of a Cochrane  
530 review. *Spine (Phila Pa 1976)* 2011;36:825-46.
- 531 47. Posadzki P. Is spinal manipulation effective for pain? an overview of systematic  
532 reviews. *Pain Med* 2012;13:754-61.
- 533 48. Szlezak AM, Georgilopoulos P, Bullock-Saxton JE, Steele MC. The immediate  
534 effect of unilateral lumbar Z-joint mobilisation on posterior chain neurodynamics:  
535 a randomised controlled study. *Man Ther* 2011;16:609-13.
- 536 49. Ekedahl H, Jönsson B, Frobell RB. Fingertip-to-floor test and straight leg raising  
537 test: validity, responsiveness, and predictive value in patients with  
538 acute/subacute low back pain. *Arch Phys Med Rehabil* 2012;93:2210-5
- 539 50. Dishman JD, Bulbulian R. Spinal reflex attenuation associated with spinal  
540 manipulation. *Spine* 2000;25:2519-25.
- 541 51. Boyd BS. Measurement properties of a hand-held inclinometer during straight  
542 leg raise neurodynamic testing. *Physiotherapy* 2012;98:174-9.
- 543 52. Colloca CJ, Keller TS, Gunzburg R. Biomechanical and neurophysiological  
544 responses to spinal manipulation in patients with lumbar radiculopathy. *J*  
545 *Manipulative Phys Ther* 2004;27:1-15.

- 546 53. Akaha H, Matsudaira K, Takeshita K, Oka H, Hara N, Nakamura K. Modified  
547 measurement of finger-floor distance. *J Japanese Soc Lumbar Spine Disord*  
548 2008;14:164-9.
- 549 54. Lewis SE, Holmes PS, Woby SR, Hindle J, Fowler NE. Short-term effect of  
550 superficial heat treatment on paraspinal muscle activity, stature recovery, and  
551 psychological factors in patients with chronic low back pain. *Arch Phys Med*  
552 *Rehabil* 2012;93:367-72.
- 553 55. Gerke DA, Brismée JM, Sizer PS, Dedrick GS, James CR. Change in spine  
554 height measurements following sustained mid-range and end-range flexion of  
555 the lumbar spine. *Appl Ergon* 2011;42:331-6.
- 556 56. Healey EL, Burden AM, McEwan IM, Fowler NE. The impact of increasing  
557 paraspinal muscle activity on stature recovery in asymptomatic people. *Arch*  
558 *Phys Med Rehabil* 2008;89:749-53.
- 559 57. Lewis S, Holmes P, Woby S, Hindle J, Fowler N. The relationships between  
560 measures of stature recovery, muscle activity and psychological factors in  
561 patients with chronic low back pain. *Man Ther* 2012;17:27-33.
- 562 58. Ross JK, Bereznick DE, McGill SM. Determining cavitation location during  
563 lumbar and thoracic spinal manipulation: is spinal manipulation accurate and  
564 specific? *Spine* 2004;29:1452-7.
- 565 59. George SZ, Fritz JM, McNeil DW. Fear-avoidance beliefs as measured by the  
566 Fear-Avoidance Beliefs Questionnaire: change in Fear-Avoidance Beliefs  
567 Questionnaire is predictive of change in self-report of disability and pain intensity  
568 for patients with acute low back pain. *Clin J Pain* 2006;22:197-203.
- 569  
570

571 **Suppliers**

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582 **Figure Legends**

583

584 **Figure 1.** Flowchart Diagram According to CONSORT Statement for the Report of  
585 Randomized Controlled Trials.

586

587 **Figure 2.** Pull-move technique in the Treatment Group.

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**Table 1.** Physical and clinical baseline characteristics of the sample\*

	Control Group (n=20)	Treatment Group (n=20)	P value
Age (yrs)	37 ± 9.31	39 ± 9.12	.541
Weight (kg)	76.65 ± 3.77	80.1 ± 8.88	.118
Height (m)	1.76 ± 0.04	1.79 ± 0.05	.159
LBP (VAS) (mm)	29.0 ± 26.33	37.1 ± 36.14	.429
SLR (degrees)	48.05 ± 11.19	39.10 ± 7.01	.004
FFD (cm)	9.9 ± 4.37	14.02 ± 9.58	.091
Stadiometry (mm)	- 0.0 ± 0.13	- 0.0 ± 0.13	1.00

Abbreviations: LBP, self-perceived low back pain; VAS, visual analogue scale; SLR, straight leg raise test, degrees of hip flexion; FFD, finger to floor distance test.

\* Data are reported as mean ± SD

**Table 2.** Pre and post-intervention values and intra-group differences in each study group \*

	Control Group			Treatment Group		
	Pre - Int	Post - Int	p	Pre - Int	Post – Int	p
LBP (VAS) (mm)	29.0 ± 26.33	29.10 ± 26.37	1.00	37.1 ± 36.14	20.01 ± 22.47	<.001
SLR (degrees)	48.05 ± 11.19	47.59 ± 10.19	1.00	39.10 ± 7.01	52.75 ± 9.53	<.001
FFD (cm)	9.9 ± 4.37	9.55 ± 4.54	.008	14.02 ± 9.58	10.35 ± 8.35	<.001
Stadiometry (mm)	- 0.0 ± 0.13	+ 0.02 ± 0.13	.142	- 0.0 ± 0.13	3.98 ± 1.46	<.001

Abbreviations: Pre – Int, Pre-Intervention; Post – Int, Post-Intervention; LBP, low back pain; VAS, visual analogue scale; SLR, straight leg raise test, degrees of hip flexion; FFD, finger to floor distance test; p value, intra-group comparison between pre- and post- intervention results

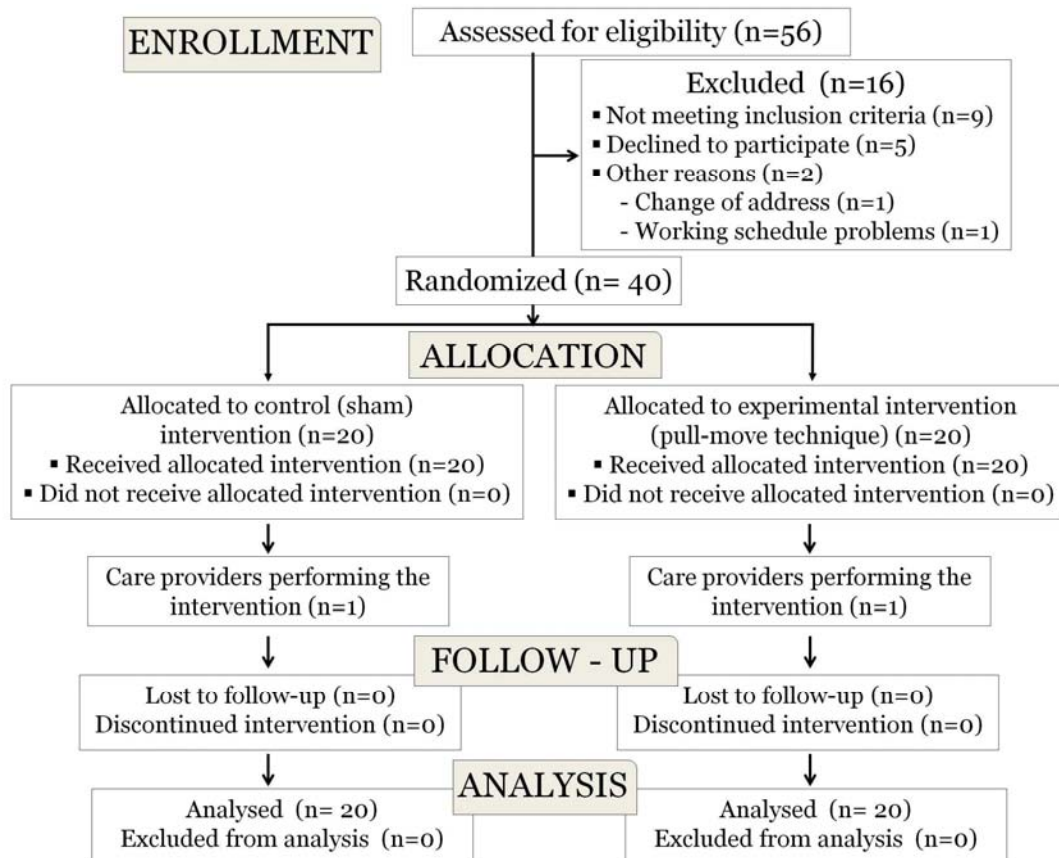
\*Values are reported as mean ± SD

**Table 3.** Between – group comparison of the mean differences from pre to post-intervention\*

	Control Group	Treatment Group	p
LBP (VAS) (mm)	0.10 ± 0.04 (0.01 / 0.08)	17.0 ± 16.57 (9.24 / 24.75)	<.001
SLR (degrees)	0.46 ± 0.39 (0.13 / .60)	-13.65 ± 8.62 (-17.68 / -9.61)	<.001
FFD (cm)	0.35 ± 0.48 (.12 / .57)	3.67 ± 2.09 (2.69 / 4.65)	<.001
Stadiometry (mm)	-0.03 ± 0.09 (-.07 / .01)	-3.98 ± 1.46 (-4.67 / -3.30)	<.001

Abbreviations: LBP, self-perceived low back pain; VAS, visual analogue scale; SLR, straight leg raise test, degrees of hip flexion; FFD, finger to floor distance test; p value, intergroup comparison of the mean values between pre and post-intervention

\* Values are reported as mean ± standard deviation (95% confidence interval)







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