UNIVERSITY OF BIRMINGHAM University of Birmingham Research at Birmingham

Variation in the spatial distribution of erector spinae activity during a lumbar endurance task in people with low back pain

Sanderson, Andy; Martinez-Valdes, Eduardo; Heneghan, Nicola R.; Murillo, Carlos; Rushton, Alison; Falla, Deborah

DOI: 10.1111/joa.12935

License: Other (please specify with Rights Statement)

Document Version Peer reviewed version

Citation for published version (Harvard):

Sanderson, A, Martinez-Valdes, E, Heneghan, NR, Murillo, C, Rushton, A & Falla, D 2019, 'Variation in the spatial distribution of erector spinae activity during a lumbar endurance task in people with low back pain', Journal of Anatomy, vol. 234, no. 4, pp. 532-542. https://doi.org/10.1111/joa.12935

Link to publication on Research at Birmingham portal

Publisher Rights Statement:

This is the peer reviewed version of the following article: Sanderson, A., Martinez-Valdes, E., Heneghan, N. R., Murillo, C., Rushton, A. and Falla, D. (2019), Variation in the spatial distribution of erector spinae activity during a lumbar endurance task in people with low back pain. J. Anat.. doi:10.1111/joa.12935, which has been published in final form athttps://doi.org/10.1111/joa.12935. This article may be used for non-commercial purposes in accordance with Wiley Terms and Conditions for Self-Archiving

General rights

Unless a licence is specified above, all rights (including copyright and moral rights) in this document are retained by the authors and/or the copyright holders. The express permission of the copyright holder must be obtained for any use of this material other than for purposes permitted by law.

•Users may freely distribute the URL that is used to identify this publication.

•Users may download and/or print one copy of the publication from the University of Birmingham research portal for the purpose of private study or non-commercial research.

•User may use extracts from the document in line with the concept of 'fair dealing' under the Copyright, Designs and Patents Act 1988 (?) •Users may not further distribute the material nor use it for the purposes of commercial gain.

Where a licence is displayed above, please note the terms and conditions of the licence govern your use of this document.

When citing, please reference the published version.

Take down policy

While the University of Birmingham exercises care and attention in making items available there are rare occasions when an item has been uploaded in error or has been deemed to be commercially or otherwise sensitive.

If you believe that this is the case for this document, please contact UBIRA@lists.bham.ac.uk providing details and we will remove access to the work immediately and investigate.

1	
2	
3	
4	
5	
6	
7	Variation in the spatial distribution of erector spinae activity during a
8	lumbar endurance task in people with low back pain
9	
10	
11 12	Andrew Sanderson, Eduardo Martinez-Valdes, Nicola R. Heneghan, Carlos Murillo, Alison Rushton, Deborah Falla
13 14 15 16 17	Centre of Precision Rehabilitation for Spinal Pain (CPR Spine) School of Sport, Exercise and Rehabilitation Sciences, College of Life and Environmental Sciences, University of Birmingham, UK
19 20 21 22 23 24 25 26	Corresponding Author – Deborah Falla <u>d.falla@bham.ac.uk</u> Centre of Precision Rehabilitation for Spinal Pain (CPR Spine) School of Sport, Exercise and Rehabilitation Sciences, College of Life and Environmental Sciences, University of Birmingham, UK

27 Abstract

- 28 This study aimed to investigate the spatial distribution and redistribution of lumbar erector
- 29 spinae (ES) activity during a lumbar extension endurance task in pain-free participants and
- 30 how this is modified in people with low back pain (LBP). High density surface
- electromyography (HDEMG) was recorded using 13x5 electrode grids placed over the
- 32 lumbar ES in 13 LBP and 13 control participants while completing an Ito test to task failure.
- 33 The root mean square of the HDEMG signals was computed, a topographical map of the
- EMG amplitude generated and the centre of the activity (centroid) determined throughout the
- task. The centroid of the EMG amplitude map was systematically more cranial (F = 6.09, P =
- 36 0.022) for the LBP participants compared to the control subjects. Regression analysis showed
- that the extent of redistribution of ES activity was associated with longer endurance. These
- results show that LBP participants utilised a different motor strategy to perform the
- endurance task, characterised by greater activation of more cranial regions of the ES and less
- 40 redistribution of ES activity throughout the task. This study provides new insight into the
- 41 functional activation of the lumbar ES and how it is modified when people have pain.
- 42 Keywords High density EMG, Ito Test, Erector Spinae, Functional Muscle Activity

43

- Funding Sources This research did not receive any specific grant from funding agencies in
 the public, commercial, or not-for-profit sectors.
- 46 **Competing Interests** There are no competing interests to declare.

48

1

Abbreviation	Meaning	
LBP	Low Back Pain	
EMG	Electromyography	
HDEMG	High Density Electromyography	
ES	Erector Spinae	
MVC	Maximum Voluntary Contraction	
ODI	Oswestry Disability Index	
TSK	Tampa Scale for Kinesiophobia	
PNRS	Pain Numeric Rating Scale	
RPE	Rate of Perceived Exertion	
RMS	Root Mean Squared Amplitude	
MNF	Mean Spectral Power Frequency	
ANCOVA	Analysis of Covariance	
ANOVA	Analysis of Variance	
CON	Control Participant	
SE	Standard Error of the Mean	

52

1. INTRODUCTION

53 Previous anatomical and biomechanical research on the lumbar erector spinae (ES) has focussed on the relationship and the structure of different portions of this muscle group 54 55 (Christophy et al., 2012, Bogduk, 1980, Bogduk, 2005). Bogduk (2005), via several dissection studies, described the origins, insertions and functions of the portions of the 56 lumbar ES, work which was adapted by Christophy and colleagues to produce a 57 biomechanical model of the lumbar musculature (Christophy et al., 2012, Bogduk, 1980). 58 These descriptions concurred that while the portions of the muscle group have different 59 origins and insertions, all play important roles in extending the lumbar spine during 60 61 functional movements. The structures described for the portions of the ES indicate a broad bilateral muscular region lateral to the lumbar spine extending from L5 into the thoracic 62 region (Bogduk, 2005, Christophy et al., 2012). To extend the lumbar spine, the most 63 64 effective motor strategy would be to recruit fibres especially from the caudal portions of the ES, creating a longer lever arm and conferring a biomechanical advantage to the movement 65 (Bogduk, 2005). 66

Surface electromyography (EMG) is used to measure muscle activity and can be 67 applied as a means to understand variations in neuromuscular control in individuals with 68 musculoskeletal pain (Abboud et al., 2014, Fabian et al., 2005, Falla et al., 2017, Falla et al., 69 2014, Gallina et al., 2011, Gizzi et al., 2015). More recently, high-density surface 70 electromyography (HDEMG) has been utilised to understand and quantify changes in the 71 spatial distribution of muscle activity which was not previously possible with classic bipolar 72 73 surface EMG. Existing research utilising HDEMG has also commonly evaluated changes in the distribution of muscle activity during either sustained or dynamic contractions by 74 quantifying a shift in the centroid of the HDEMG amplitude map, the point which defines the 75 76 barycentre of muscle activation (Falla et al., 2014, Madeleine et al., 2006, Farina et al., 2008,

Falla et al., 2017, Gallina et al., 2013). HDEMG studies on healthy asymptomatic volunteers
have shown that the centre of muscle activity shifts during contraction (Falla and Farina,
2008b, Farina et al., 2008, Tucker et al., 2009) and that this redistribution of muscle activity
has the physiological significance of minimising muscle fatigue and prolonging endurance
(Farina et al., 2008, Gallina et al., 2013, Falla et al., 2014), possibly by preventing overload
on the muscle fibres active at the beginning of the task.

Previous HDEMG investigations have described an association between endurance 83 time and the redistribution of muscle activity in the trapezius in asymptomatic participants 84 (Farina et al., 2008). More Recently HDEMG was applied to evaluate changes in lumbar ES 85 activity in a LBP population (Abboud et al., 2014). Participants completed a force-matching 86 modified Sørenson test (lifting of the unsupported upper body with the legs affixed to a 87 plinth), resisting a load cell around their shoulders which simulated 30% of their maximum 88 89 voluntary contraction (MVC). Increased variability in the position of the centroid of the EMG amplitude map was observed in the healthy controls compared to the LBP group. 90

Despite these observations, the functional relevance of a change in the distribution of
muscular activity remains unclear. We hypothesised that people with LBP would engage
different regions of the lumbar ES during isometric back extension, reflecting less efficient
activation of the ES and, that people with LBP would show less redistribution of ES activity
which would be associated with significantly lower endurance in this group.

96 Thus, the aim of this study was to investigate the spatial distribution of lumbar ES
97 activity and redistribution of activity during an endurance task in participants with chronic
98 LBP and pain-free controls. Moreover, we evaluate the relationship between the extent of
99 redistribution of activity and endurance time, with the hypothesis that those who display a
100 larger redistribution of activity would be able to sustain the contraction for longer. This study

- stands to provide new insight into the functional activation of the lumbar ES and how it is
- 102 modified when people have pain.

103 **2 - METHODS**

This study was an observational, cross-sectional case-control study using a
convenience sample of participants from the staff, students and community of the University
of Birmingham, UK. Data collection took place in a laboratory within the Centre of Precision
Rehabilitation for Spinal Pain, University of Birmingham.

108 2.1 - Participants

LBP participants aged 20-55 were recruited via posters and social media accounts 109 related to the University of Birmingham. Due to the nature of the fatiguing task, it was 110 decided that 55 would be the maximum age of participants eligible for this study. Eligibility 111 criteria included non-specific LBP which had persisted for at least half the days of the 112 previous six months, exceeding the minimum definition for LBP (Dionne et al., 2008). 113 114 Consistent with previous studies, age and gender matched control participants (CON) were recruited in the same way and were included if they had no history of LBP or lower limb 115 disorders. Exclusion criteria for both groups comprised concurrent systemic issues including 116 rheumatic and neuromuscular disorders, a history of chronic respiratory or neurological 117 problems, spinal deformity or surgery, cardiovascular conditions, pregnancy, and healthcare 118 management for LBP in the previous 6 months (a requirement of the University ethical 119 committee). To support a normal distribution for statistical analysis, a planned sample size of 120 30 participants (15 LBP and 15 CON) was chosen, consistent with previous HDEMG studies 121 comparing symptomatic and asymptomatic participants. 122

Ethical approval was granted by the University of Birmingham ethics committee (ERN_17-0782). Participants gave written informed consent prior to data collection and all procedures were completed in accordance with the Declaration of Helsinki. 127 Prior to testing, participants from both groups were required to complete several questionnaires to gather population statistics; including the level of disability, intensity of 128 129 pain, and current level of activity. Participants were asked to complete the Oswestry Disability Index (ODI), as it has previously been shown to be a reliable measure of disability 130 relating to spinal pain (Fairbank and Pynsent, 2000). The Tampa Scale for Kinesiophobia was 131 used to assess any fear surrounding movement related to pain (Miller et al., 1991). A Pain 132 Numeric Rating Scale (0-10) (PNRS) was used to assess current pain at the time of testing, 133 and pain over the prior week (Breivik et al., 2008). Information on the general health of 134 135 participants at the time of testing was collected using the RAND 36 item health survey, which has been shown to be effective and reliable as a measure of health across cultures and gender 136 (Hays et al., 1993, VanderZee et al., 1996). Throughout the endurance task (see below), the 137 138 rate of perceived exertion (RPE) was recorded at 30 s intervals and immediately following task failure using the Borg RPE scale (Borg, 1998). This measure was used to assess the 139 140 perceived exertion of the participants throughout the task and ensure that the task was appropriate for LBP participants. 141

142 2.3 - Experimental Set-Up

Surface EMG signals were recorded from the lumbar ES using 13x5 semi-disposable 2D electrode grids (OT Bioelettronica, Italy). Electrodes were spaced evenly with a 1 mm diameter and an 8 mm inter-electrode distance; one corner electrode was missing in each grid to provide directional reference. Electrodes were positioned over the lumbar ES on the righthand side in control participants and the most painful side in the LBP group. Where equal pain was reported bilaterally, participants were randomly allocated a side (Figure 1).

Prior to the application of electrodes, the skin in the region lateral to the lumbar spine 149 was prepared by firstly shaving the area if needed and then applying an abrasive paste (SPES 150 Medica, Italy), and finally washing and drying the region. The electrodes were prepared by 151 applying a thin custom double-sided adhesive foam pad to the electrode grid (SPES Medica, 152 Genova, Italy). The cavities of the electrode grids were then filled with an electoconductive 153 paste (SPES Medica, Genova, Italy). As there is no way of differentiating different portions 154 155 of the ES in-vivo, the electrode was placed on the ES in accordance with EMG guidelines and previous studies (Barbero et al., 2012, Falla et al., 2014). The grids were then applied to 156 157 the skin approximately 2cm lateral to the lumbar spinous processes, starting at the level of the L5 and extending to approximately the level of L3, as described previously (Falla et al., 158 2014). Reference electrodes were placed on prepared skin over the right anterior superior 159 160 iliac spine and on the spinous process of the vertebra prominens.

161 A twin-axis SG150B digital goniometer (Biometrics Ltd., Gwent, UK) was applied to the right mid-axillary line of the participant. Only one axis (sagittal plane) was used for 162 163 analysis. The lower sensor was attached to the centre of the iliac crest, with the midline of the sensor in line with the greater trochanter of the femur. With the participant positioned in 164 prone on the plinth, the resting angle was calibrated as 0°, with trunk deviation measured 165 from this point. EMG signals and angular data were sampled at 2048Hz and amplified (400-166 channel EMG amplifier Quattrocento, OT Bioelettronica, Torino, Italy; -3dB, bandwidth 10-167 500 Hz)) by a factor of 150 and converted to digital form by a 16-bit analogue-to-digital 168 converter. Collected signals were stored on a computer hard drive and later analysed using a 169 custom code on MATLAB (The Mathworks Inc., USA). 170

As described previously by Falla et al. (2014), each grid of electrodes recorded 64
monopolar signals. These signals were then processed offline to form horizontal derivatives
across the grids. This was achieved by first filtering the monopolar signals using a 20-350 Hz

174 band-pass filter and then adjacent signals were processed to produce 59 bipolar EMG signals. The amplitude (RMS) and mean spectral power frequency (MNF) for each bipolar derivation 175 were then calculated. The individual RMS and MNF values for each bipolar signal were 176 averaged to produce the mean RMS and MNF values across the grid. The RMS values for 177 each bipolar signal were used to create a topographical map of ES activity. This map was 178 used to determine the location of the x- and y-coordinates of the centroid as described 179 previously (Abboud et al., 2014, Falla et al., 2017, Falla et al., 2014, Farina et al., 2008, 180 Madeleine et al., 2006, Tucker et al., 2009). The location of the centroid was averaged across 181 182 the 10s or 10% epochs for further use in analysis.

The values for the x- and y- coordinates of the centroid were analysed as an absolute shift in mm from the start point quantified in the first 10% epoch (Falla et al., 2014). As movement of the centroid was both cranial and caudal in both groups, to allow for comparison between groups of the absolute shift in the y-coordinate of the centroid, both positive and negative movements were made positive.

188 2.4 - Experimental Procedure

To complete the endurance task, participants were required to maintain an Ito test, as 189 described by Ito et al. (1996) and Muller et al. (2010) until task failure or until 300s. 190 Participants were first asked to lie prone on a plinth, with a firm semi-circular foam pad (18 191 cm diameter) centred below the anterior superior iliac spines. To complete the endurance 192 task, participants were asked to lift their sternum from the plinth, raising their upper body by 193 ~15°. While maintaining this position, participants were asked to keep their arms in line with 194 the body axis and not in contact with the plinth; participants were also required to contract 195 196 their gluteal muscles and retain a neutral neck position. Prior to beginning the task, an

investigator demonstrated the correct position for completion of the Ito test, and participantswere permitted to complete a short 5 s contraction to ensure they had the correct technique.

Throughout the task, the angle of the body axis was monitored visually and 199 participants were alerted if their body axis was approaching the upper or lower acceptable 200 limits $(\pm 10^{\circ})$ (Demoulin et al., 2007). Task failure was determined by a drop in the angle of 201 trunk of greater than 10° at any point. While completing the contraction, participants were 202 timed using a stopwatch, the time was recorded until task failure or until the maximum 203 contraction duration was reached (300 s). Throughout the task, participants were given verbal 204 encouragement and at 30s increments were provided with feedback for how long they had 205 206 sustained the contraction.

207 2.5 - Statistical Analysis

Statistical analysis was performed using SPSS 24 (IBM, USA) with an alpha level set at 0.05. Regression analysis and analysis of covariance (ANCOVA) tests were performed using Prism (GraphPad, USA). Where P-values were reported on SPSS as 0.000, they have been stated as P < 0.001 herein. Effect sizes have been reported where appropriate with ANOVA results, based on guidance by Lakens (2013) in the format of generalised η^2 (η_g^2), alongside η^2 values. For interpretation of these values, effect sizes are defined as small (η^2/η_g^2 = 0.01), medium (η^2/η_g^2 = 0.06), or large (η^2/η_g^2 = 0.14) (Lakens, 2013, Cohen, 1988).

A student t-test was performed in order to identify any differences in endurance times between groups. The questionnaires used to gather sample characteristics were interpreted according to their respective guidelines (Childs et al., 2005, Fairbank and Pynsent, 2000, Hays et al., 1993, Miller et al., 1991). Student t-tests were performed for each group to identify differences between the samples at baseline. To determine if the failure of the task was influenced by fear of movement, the endurance time for each participant was correlatedto their respective TSK score.

No direct comparison of the values reported for perceived exertion could be made between groups, as the time to task failure varied between groups. Therefore, the initial value after 30 s, the value at the mid-point of endurance, and the level of exertion at task failure were determined for each participant. Significant differences between groups were investigated using a repeated measures analysis of variance (ANOVA).

In order to make comparisons between groups with different times to task failure, the total contraction time for each participant was normalised into 10% epochs of the total endurance time (Farina et al., 2008). Repeated measures ANOVA, with factors of group (CON and LBP) and time (10 epochs) were used to compare differences in EMG variables between groups. Newman-Keuls post-hoc tests were also conducted where appropriate.

To identify trends in the displacement of the y-coordinate of the centroid between groups, a linear regression was performed. To ensure that the results were not affected by the normalisation of time, this regression was performed using absolute endurance times and ycoordinate displacement values calculated from the position of the centroid in the first 10 s epoch. The regression lines for the CON and LBP groups were compared for statistical significance using an ANCOVA.

Finally, in order to assess myoelectric manifestations of muscle fatigue, linear regressions were performed on RMS and MNF variables (Larivière et al., 2002). For each participant, the relationships between RMS and time to task failure, and MNF and time to task failure were computed. In this analysis both absolute values for RMS and MNF across time, and normalised values (using the using the first 10s epoch as a reference) were considered and the resulting slopes extracted. Independent samples t-tests were then

- 244 performed on the slopes for each condition to identify the mean slope for each group and
- identify any differences between these means (Pagé and Descarreaux, 2012, Roy et al., 1995).

3 - RESULTS

247 3.1 - Participants

13 LBP and 13 CON participants successfully completed data collection, population 248 characteristics are reported in Table 1. No significant anthropometric differences were found 249 between groups for BMI, height or weight and the BMI for both groups was within the 250 'normal weight' range (Stenholm et al., 2017). However, as anticipated the LBP group 251 presented with higher levels of disability (ODI -13.16%) and lower general and emotional 252 253 health (RAND 36 item health survey). Prior to data collection, LBP participants reported a current pain level of 1.92 out of 10, but a usual pain of 2.92, characterising the pain within 254 the group as mild or low severity (Breivik et al., 2008). No significant correlation was found 255 between scores on the TSK and the endurance time (R = -0.281, P = 0.165). 256

257 3.2 - Endurance

Significantly lower endurance times were recorded for the LBP group (F = 8.4, P < 0.001) compared to the control group (186.3 \pm 72.3s and 283.0 \pm 33.0s respectively). With 96.7s difference, this equates to the LBP group maintaining the contraction for 65.8% of the total time for the CON group on average. The mean values for initial, middle and final perceived exertion are shown in Figure 2. No significant differences were found between groups for exertion at any point (F = 1.42, P = 0.216).

264 3.3 – Electromyographical Changes

265 3.3.1 – EMG amplitude and mean frequency of the EMG signal

Across the duration of the contraction, the RMS was found to be systematically higher

- for CON than LBP participants (main effect of group; F = 6.09, P = 0.022, $\eta_g^2 = 0.18$, $\eta^2 = 0.$
- 268 0.18) (Figure 3). This higher activation of the ES was visible in the topographical maps of the

EMG amplitude (Figure 4). On average, the CON participants showed a larger distribution of the activity throughout the entire muscle, whereas LBP participants showed a less diffused activation which tended to be more cranial. When this was quantified, an even distribution across the entire grid was observed in 11 CON and 4 LBP participants. Distribution was weighted cranially in 1 CON and 8 LBP participants and distribution was focussed in the middle of the grid for 1 Con and 1 LBP participant.

There were no significant differences between groups for the change in RMS throughout the task (F = 1.42, P = 0.216). There was also no significant increases or decreases in the mean RMS recorded for either group at any point during the endurance task (F = 0.929, P = 0.344). No significant differences between groups were observed for the mean MNF at any point during the contraction (F = 1.118, P = 0.334).

280 3.3.2 – Centroid of the EMG RMS map

No significant differences were found between groups for the position of the xcoordinate of the centroid (medial-lateral direction) throughout the task (initial position – F = 2.27, P = 0.77; shift over the duration of the contraction – F = 2.27, P = 0.77).

The y-coordinate of the centroid (cranial-caudal direction) in CON participants was found to be systematically more caudal than the LBP group (main effect for group; F = 44.00, P < 0.001, $\eta_g^2 = 0.64$, $\eta^2 = 0.65$). The y-coordinate in CON participants was found to be approximately 42.0 mm (± 4.99 mm) cranial of the reference electrode, whereas for the LBP participants, the y-coordinate was approximately 53.6 mm (± 3.64 mm) cranial of the reference electrode. Throughout the endurance task there was a mean difference between the LBP and CON in the y-coordinate position of 11.6mm (Figure 5).

Using the location of the y-coordinate of the centroid in the 1st epoch as a reference 291 point, the displacement was calculated for each 10% epoch. To achieve this, the shift in mm 292 was measured from the position of the y-coordinate in the first epoch, this could be either a 293 294 positive (cranial movement) or negative (caudal) value. No clear direction of shift was found (cranially or caudally) as groups showed both cranial and caudal movements (CON - 6295 cranial, 7 caudal; LBP – 9 cranial, 4 caudal). To better understand the movement of the 296 centroid, all values for displacement were therefore made positive and so net displacement is 297 used for all y-coordinate shift results. At task failure, the mean y-coordinate displacement for 298 299 the CON group was 2.10±0.45mm whereas for the LBP group it was 1.40±0.29mm. Both groups showed a significant displacement of the centroid in the y-axis over time (F=2.5, 300 P=0.004, $\eta_g^2 = 0.22$, $\eta^2 = 0.30$) and a significant displacement within each group (F=9.9, 301 P=0.01) (Figure 6). There was no interaction between groups for the displacement of the y-302 coordinate in the data which had been normalised to task failure (F = 1.709, P = 0.134). 303

The regression analysis performed using absolute values for time showed a significant relationship between the shift in the y-coordinate of the centroid and the time to task failure (Figure 7) for both groups (CON - $r^2 = 0.142$, P < 0.0001; LBP - $r^2 = 0.053$, P = 0.0004). Additionally, ANCOVA analysis showed that there was a significant difference between the regression lines for each group (F=5.597, P=0.0183) indicating that the relationship between y-coordinate shift and time was significantly different between groups (LBP/CON) (Zar, 2010).

311 Myoelectric manifestations of muscle fatigue showed no differences under any 312 condition. There were no differences in the slopes between groups for absolute RMS (P = 313 0.71), normalised RMS (P = 0.37), absolute MNF (P = 0.48) or normalised MNF (P = 0.79).

314 **4 – DISCUSSION**

This is the first study to assess muscle activation behaviour using HDEMG during a functional position-matching lumbar endurance task in people with and without LBP. The results revealed an altered motor control strategy to a standardised endurance task in people with LBP with evidence of activation of more cranial regions of the lumbar ES with respect to asymptomatic people. Moreover, a relationship was also demonstrated between the extent of redistribution of muscle activity and endurance time which has important implications for the understanding of the neurophysiological responses to fatigue.

322 4.1 –Distribution of Activity

Throughout the endurance contraction, the RMS was found to be significantly higher 323 324 in the CON group than the LBP group. One possible explanation for this disparity in 325 amplitude could be quantified from the systematic differences in the position of the centroid along the y-axis. Throughout the task the y-coordinate of the centroid for the CON group was 326 12mm caudal to that of the LBP group. Previous studies which have induced pain via 327 injection of hypertonic saline, have shown that areas with greater pain show reduced activity 328 and that in an acute painful condition, the muscle activation can shift outside of the painful 329 330 region (Falla et al., 2017, Falla and Farina, 2008a, Madeleine et al., 2006). Although somewhat speculative, it is likely that a more caudal centre of contraction could indicate a 331 more biomechanically favourable contraction through activating a greater number of fibres. 332 In this instance, those with pain appear to have shifted the activity in the ES more cranially. 333 334 A more caudal contraction, which is distributed over a larger area of the muscle would be able to utilise the larger volume of muscles from lower lumbar vertebrae and spread the load 335 336 more effectively across a greater number of muscle fibres creating a longer lever arm (Bogduk, 2005). The longer lever arm would act to minimise the force needed to sustain the 337

contraction and the diffuse activation would reduce localised fatigue, facilitating sustainedendurance.

340 4.2 - Redistribution of lumbar ES activity

During the Ito test, the CON participants showed a greater shift of the centroid of the 341 EMG amplitude map indicating a greater redistribution of lumbar ES activity than the LBP 342 group. It was also shown that the amount of redistribution increased progressively over the 343 duration of the task and that there was an association between the extent of redistribution of 344 activity and endurance time. As previously described by Falla et al. (2014), a redistribution of 345 activity likely prevents localised muscle fatigue through the build-up of metabolic factors and 346 overload on specific regions of the muscle. The task used in Falla et al (2014) was dynamic 347 348 and consisted of periodic contractions, whereas the contraction used here is static and so the tissue would be under further strain due to decreased blood flow and ischemia (Masuda et al., 349 1999). 350

351 The results of this study do not support a direction of shift for this task as there was no clear preference for a direction in either group. However, this study differs from previous 352 studies which used HDEMG to examine the lumbar muscles as it does not involve an external 353 force. Gallina et al. (2013) investigated the significance of the shift in the trapezius muscle 354 and determined that the direction of shift was task dependant. Russ et al. (2018) and Thomas 355 356 et al. (2011) showed that there were specific differences in lumbar endurance between force-357 and position-matching tasks, the reasons for which they were unable to describe. The lack of 358 a clear direction of shift seen in this study, may imply a focus of muscle activity in a more biomechanically favourable point for each participant. As there was no specific point to 359 360 'push' against, the centre of activity for each participant was likely determined by individual anthropomorphic features, for example a greater trunk length to leg length ratio. In this study, 361

it is speculated that as participants were not secured to the plinth or pushing against a point,
the impact of the relative size and weight of the legs compared to the upper body would
impact on the stability of the participant while contracting. Thus the participant might be
likely to sustain a contraction which affords them the optimal stability for their individual
anthropomorphic characteristics.

367 4.3 – Muscular Activity

Biomechanical and anatomical models of the lumbar musculature indicate that the 368 shared insertions of portions the ES cause a diagonal slight overlapping of successive 369 superficial fibres (Bogduk, 2005, Bogduk, 1980). According to anatomical studies, the 370 portions of the ES which are likely to be muscular in the region beginning 2cm lateral to L5 371 372 include the *iliocostalis lumborum pars lumborum* and the *iliocostalis lumborum pars* thoracis, with the muscular portions of the longissimus being too medial or too cranial to be 373 covered by the electrode grid. In the pars lumborum, most deep and lateral fibres are from 374 375 L5, the most superficial and medial fibres are from L1; each successive lamina of fibres 376 slightly overlaps the previous layer (Bogduk, 2005, Christophy et al., 2012). The Ito test used in this study is designed to gain relative isolation of the lumbar musculature, so the 377 378 distribution and redistribution of activity in this portion of the muscle is thought to be key to understanding endurance in this task (Muller et al., 2010). It is therefore suspected and 379 proposed that due to pain in the lumbar region, LBP participants utilised a motor control 380 381 strategy which preferentially activated different portions of the muscle, such as the more cranial *iliocostais lumborum pars thoracis* and thus led to a shorter time to task failure than 382 the CON group. As no imaging was used in this study, the exact distribution of activity 383 among portions of the ES, and what effect any individual variations in muscle architecture or 384 fibre distribution could have on the activation pattern remains unknown (Mannion et al., 385 386 1997, Mannion et al., 2000).

388 The LBP group demonstrated endurance which fell significantly short of the CON group. Similar findings have been demonstrated in previous studies investigating lumbar 389 390 endurance to task failure, however the absolute endurance times reported here were significantly higher than those reported following a Sørenson test (Abboud et al., 2014, 391 Jubany et al., 2017). This difference could be attributed in part to the differences between 392 393 force-matching tasks previously used, and position-matching such as the Ito test used here (Russ et al., 2018). However, Muller et al. (2010) reported lower endurance times for a 394 position-matching Sørenson test when compared directly with an Ito test. In this instance it 395 396 may also be relevant to consider the biomechanical and myoelectrical differences between the Ito and Sørenson positions. As discussed previously, muscle activation in the Ito test is 397 focussed on the lumbar region while the Sørenson has shared activation between the lumbar 398 399 and hip extensors, possibly contributing to differences in endurance time (Muller et al., 2010). Additionally though both tasks are measures of lumbar endurance, each requires a 400 401 different position to be held; with the Ito test requiring spinal extension to be sustained and 402 the Sørenson requiring an unsupported neutral spine maintained against gravity (Muller et al., 2010). Due to this, it is likely that the point at which the participant's centre of mass is 403 404 supported may be lower in the Ito test, producing a lower moment.

At task failure, both LBP and CON participants reported a mean RPE of between 18.3 - 18.5, indicating that both groups reached a similar level of exertion. Analysis of the MNF results and the indices measuring the myoelectric manifestations of fatigue revealed that there were no significant differences between the CON and LBP groups. Previous HDEMG studies evaluating fatigue of the lumbar ES have shown greater myoelectric manifestations of fatigue than these results suggest, however other studies also did not find significant differences between groups (Abboud et al., 2014, Tucker et al., 2009). This could be somewhat explained

by recent studies which have demonstrated that frequency variables, including MNF, do not 412 accurately predict motor unit recruitment during contractions (Merletti and Farina, 2016, 413 Vecchio et al., 2017). Additionally, it has been shown in the knee extensors that myoelectric 414 manifestations of fatigue are only seen when the exertion exceeds 40% of the MVC (de 415 Ruiter et al., 2012). Two exercises in a study by Plamondon et al. (2002) were similar in 416 position and function to the Ito test, in that study these exercises were found to be between 417 418 26-32% of a participant's MVC. As the current study did not assess the functional capacity of the participants, the results for MNF may be affected by the task being below 40% of an 419 420 MVC for some participants.

The results of this study coalesce to indicate that the LBP participants utilise a different motor control strategy to complete the task. This strategy was characterised by a reduced activation of the ES which was focussed more cranially and throughout the task showed less redistribution of activity. It appears that participants used less favourable portions of the ES to complete the task which lead to shorter endurance times.

426 4.5 - Strengths and Limitations

A strength of this study was its use of HDEMG to present a more comprehensive 427 characterisation of ES activity during an endurance test, a test which can be easily replicated 428 in a clinical environment. In addition, the Ito test presented here has previously been found to 429 430 better isolate the lumbar musculature than the Sørenson test (Muller et al., 2010). As no significant differences were found in the RPE between groups, it is supported that the Ito test 431 432 was a suitable test for this population. However, it should be considered that as no clear guidelines for task failure have been validated for the Ito test, the task failure criteria of $\pm 10^{\circ}$ 433 434 could be perceived to affect the redistribution of activity during the task. However, as systematic differences were seen between groups for all values related to the RMS, we are 435

436 confident that the differences between groups are valid, but could hinder comparison to other lumbar endurance tasks. To mitigate this effect, where possible, generalised effect sizes have 437 been reported with ANOVA results, which have been interpreted in line with the guidelines 438 439 suggested by Cohen (1988), and reiterated by Lakens (2013), whereby effect sizes are defined as small ($\eta^2 = 0.01$), medium ($\eta^2 = 0.06$), or large ($\eta^2 = 0.14$). However it has been 440 suggested that these benchmarks for η^2 may not be as accurate in repeated measures 441 conditions. Therefore we also included the η_g^2 values, which have been proposed to allow 442 better comparisons between studies (Lakens, 2013). 443

Since participants could not be under current active management by a healthcare professional 444 (a requirement of the University Ethical Committee), the LBP group presented with low 445 levels of current pain and mild disability. Although the sample size was relatively small and 446 the LBP participants presented with relatively mild LBP, significant group differences were 447 448 revealed and even greater group differences may be expected when testing patients with even greater pain severity or longer pain duration (Arendt-Nielsen and Graven-Nielsen, 2008, 449 450 Mannion et al., 2000). Finally, synergistic muscles were not covered by the HDEMG grid. 451 This limitation was imposed in an attempt to reduce the effect of crosstalk between overlapping muscles of different architecture which may have confounded the results 452 (Martinez-Valdes et al., 2018). Further studies using mixed methodologies, including 453 intramuscular electrodes and motor unit decomposition, may provide clearer information 454 about individual muscle contributions to this task. 455

5 - CONCLUSION

457	Asymptomatic people display a spatial redistribution of lumbar ES activity during an
458	endurance task and this adaptation is reduced in people with LBP. Moreover, people with
459	LBP engage more cranial regions of the lumbar ES during trunk extension; likely reflecting
460	an inefficient motor strategy.
461	
462	
463	The authors declare no conflict of interest.
464	
465	

466 **Author Contributions**

- 467 AS, EMV and DF contributed to study conception and design. AS and CM acquired the data.
- 468 AS, EMV and DF performed the data analysis. All authors contributed to interpretation of the
- data. Drafting of the manuscript was performed by AS, EMV and DF. All authors
- 470 participated in revising the manuscript and approving the final submission.

471

473 **Reference List**

474 Abboud J, Nougarou F, Page I, Cantin V, Massicotte D, Descarreaux M (2014) Trunk motor variability 475 in patients with non-specific chronic low back pain. Eur J Appl Physiol, 114, 2645-54. 476 Arendt-Nielsen L, Graven-Nielsen T (2008) Muscle pain: sensory implications and interaction with 477 motor control. *Clin J Pain*, **24**, 291-8. 478 Barbero M, Merletti R, Rainoldi A (2012) Atlas of Muscle Innervation Zones, Springer-Verlag Mailand. 479 Bogduk N (1980) A reappraisal of the anatomy of the human lumbar erector spinae. J Anat, 131, 525-480 40. 481 Bogduk N (2005) Clinical anatomy of the lumbar spine and sacrum, Elsevier/Churchill Livingstone, 482 Edinburgh. 483 Borg G (1998) Borg's perceived exertion and pain scales, Human kinetics. 484 Breivik H, Borchgrevink PC, Allen SM, et al. (2008) Assessment of pain. Br J Anaesth, 101, 17-24. 485 Childs JD, Piva SR, Fritz JM (2005) Responsiveness of the numeric pain rating scale in patients with 486 low back pain. Spine (Phila Pa 1976), 30, 1331-4. 487 Christophy M, Faruk Senan NA, Lotz JC, O'Reilly OM (2012) A musculoskeletal model for the lumbar 488 spine. *Biomech Model Mechanobiol*, **11**, 19-34. 489 Cohen J (1988) Statistical Power Analysis for the Behavioral Sciences, Routledge Academic, New 490 York, NY. 491 de Ruiter CJ, Maas EA, Wesseling MG, de Haan A (2012) Knee extensor fatigue threshold is related to 492 whole-body VO2max. Med Sci Sports Exerc, 44, 1366-74. 493 Demoulin C, Crielaard JM, Vanderthommen M (2007) Spinal muscle evaluation in healthy individuals 494 and low-back-pain patients: a literature review. Joint Bone Spine, 74, 9-13. 495 Dionne CE, Dunn KM, Croft PR, et al. (2008) A consensus approach toward the standardization of 496 back pain definitions for use in prevalence studies. Spine (Phila Pa 1976), **33**, 95-103. 497 Fabian S, Hesse H, Grassme R, Bradl I, Bernsdorf A (2005) Muscular activation patterns of healthy 498 persons and low back pain patients performing a functional capacity evaluation test. 499 Pathophysiology, 12, 281-7. 500 Fairbank JC, Pynsent PB (2000) The Oswestry Disability Index. Spine (Phila Pa 1976), 25, 2940-52; 501 discussion 2952. 502 Falla D, Cescon C, Lindstroem R, Barbero M (2017) Muscle Pain Induces a Shift of the Spatial 503 Distribution of Upper Trapezius Muscle Activity During a Repetitive Task: A Mechanism for 504 Perpetuation of Pain With Repetitive Activity? *Clin J Pain*, **33**, 1006-1013. 505 Falla D, Farina D (2008a) Neuromuscular adaptation in experimental and clinical neck pain. J 506 Electromyogr Kinesiol, 18, 255-61. 507 Falla D, Farina D (2008b) Non-uniform adaptation of motor unit discharge rates during sustained 508 static contraction of the upper trapezius muscle. Exp Brain Res, 191, 363-70. 509 Falla D, Gizzi L, Tschapek M, Erlenwein J, Petzke F (2014) Reduced task-induced variations in the 510 distribution of activity across back muscle regions in individuals with low back pain. Pain, 511 155, 944-53. 512 Farina D, Leclerc F, Arendt-Nielsen L, Buttelli O, Madeleine P (2008) The change in spatial 513 distribution of upper trapezius muscle activity is correlated to contraction duration. J 514 Electromyogr Kinesiol, 18, 16-25. 515 Gallina A, Merletti R, Gazzoni M (2013) Uneven spatial distribution of surface EMG: what does it 516 mean? Eur J Appl Physiol, 113, 887-94. 517 Gallina A, Merletti R, Vieira TM (2011) Are the myoelectric manifestations of fatigue distributed 518 regionally in the human medial gastrocnemius muscle? J Electromyogr Kinesiol, 21, 929-38. 519 Gizzi L, Muceli S, Petzke F, Falla D (2015) Experimental Muscle Pain Impairs the Synergistic Modular 520 Control of Neck Muscles. PLoS One, 10, e0137844. 521 Hays RD, Sherbourne CD, Mazel RM (1993) The RAND 36-Item Health Survey 1.0. Health Econ, 2, 522 217-27.

- Ito T, Shirado O, Suzuki H, Takahashi M, Kaneda K, Strax TE (1996) Lumbar trunk muscle endurance
 testing: an inexpensive alternative to a machine for evaluation. *Arch Phys Med Rehabil*, **77**,
 75-9.
- Jubany J, Marina M, Angulo-Barroso R (2017) Electromyographic and Kinematic Analysis of Trunk
 and Limb Muscles During a Holding Task in Individuals With Chronic Low Back Pain and
 Healthy Controls. *PM R*, **9**, 1106-1116.
- Lakens D (2013) Calculating and reporting effect sizes to facilitate cumulative science: a practical
 primer for t-tests and ANOVAs. *Frontiers in psychology*, 4, 863-863.
- Larivière C, Arsenault AB, Gravel D, Gagnon D, Loisel P (2002) Evaluation of measurement strategies
 to increase the reliability of EMG indices to assess back muscle fatigue and recovery. *Journal of Electromyography and Kinesiology*, **12**, 91-102.
- Madeleine P, Leclerc F, Arendt-Nielsen L, Ravier P, Farina D (2006) Experimental muscle pain
 changes the spatial distribution of upper trapezius muscle activity during sustained
 contraction. *Clin Neurophysiol*, **117**, 2436-45.
- Mannion AF, Dumas GA, Cooper RG, Espinosa FJ, Faris MW, Stevenson JM (1997) Muscle fibre size
 and type distribution in thoracic and lumbar regions of erector spinae in healthy subjects
 without low back pain: Normal values and sex differences. *Journal of Anatomy*, **190**, 505513.
- Mannion AF, Kaser L, Weber E, Rhyner A, Dvorak J, Muntener M (2000) Influence of age and
 duration of symptoms on fibre type distribution and size of the back muscles in chronic low
 back pain patients. *Eur Spine J*, 9, 273-81.
- Martinez-Valdes E, Negro F, Falla D, De Nunzio AM, Farina D (2018) Surface electromyographic
 amplitude does not identify differences in neural drive to synergistic muscles. J Appl Physiol
 (1985), 124, 1071-1079.
- Masuda K, Masuda T, Sadoyama T, Inaki M, Katsuta S (1999) Changes in surface EMG parameters
 during static and dynamic fatiguing contractions. *J Electromyogr Kinesiol*, **9**, 39-46.
- Merletti R, Farina D (2016) *Surface Electromyography: Physiology, Engineering, and Applications,* John Wiley & Sons.
- Miller RP, Kori SH, Todd DD (1991) The Tampa Scale: a Measure of Kinisophobia. *The Clinical journal* of pain, 7, 51.
- Muller R, Strassle K, Wirth B (2010) Isometric back muscle endurance: an EMG study on the criterion
 validity of the Ito test. *J Electromyogr Kinesiol*, **20**, 845-50.
- Pagé I, Descarreaux M (2012) Trunk muscle fatigue during a lateral isometric hold test: what are we
 evaluating? *Chiropr Man Therap*, **20**, 12.
- Plamondon A, Serresse O, Boyd K, Ladouceur D, Desjardins P (2002) Estimated moments at L5/S1
 level and muscular activation of back extensors for six prone back extension exercises in
 healthy individuals. *Scand J Med Sci Sports*, **12**, 81-9.
- Roy SH, De Luca CJ, Emley M, Buijs RJ (1995) Spectral electromyographic assessment of back muscles
 in patients with low back pain undergoing rehabilitation. *Spine (Phila Pa 1976)*, **20**, 38-48.
- Russ DW, Ross AJ, Clark BC, Thomas JS (2018) The Effects of Task Type on Time to Task Failure During
 Fatigue: A Modified Sorensen Test. J Mot Behav, 50, 96-103.
- Stenholm S, Head J, Aalto V, et al. (2017) Body mass index as a predictor of healthy and disease-free
 life expectancy between ages 50 and 75: a multicohort study. *International Journal Of Obesity*, 41, 769.
- Thomas JS, Ross AJ, Russ DW, Clark BC (2011) Time to task failure of trunk extensor muscles differs
 with load type. *J Mot Behav*, 43, 27-9.
- Tucker K, Falla D, Graven-Nielsen T, Farina D (2009) Electromyographic mapping of the erector
 spinae muscle with varying load and during sustained contraction. *J Electromyogr Kinesiol*,
 19, 373-9.

- VanderZee KI, Sanderman R, Heyink JW, de Haes H (1996) Psychometric qualities of the RAND 36 Item Health Survey 1.0: a multidimensional measure of general health status. *Int J Behav Med*, 3, 104-22.
- Vecchio AD, Negro F, Felici F, Farina D (2017) Associations between motor unit action potential
 parameters and surface EMG features. **123**, 835-843.
- 577 Zar JH (2010) *Biostatistical Analysis,* Prentice Hall.

578

Tables

Characteristic		LBP	Control	P-Value
Age (Years)		27.39 ±9.7	26.46 ±5.0	-
Gender (# Males)		6	7	-
Height (cm)		168.75 ±9.7	170.38 ±6.7	-
Weight (kg)		70.97 ±12.4	69.11 ±12.7	-
BMI		24.78	23.78	-
ODI (%) *		13.16% ±8%	0.00%	< 0.001
TSK		25.31 ±4.89	22.31 ±7.20	-
DNIPS	Current Pain *	1.92 ±1.44	0	< 0.001
FINKS	Usual Pain *	2.92 ±1.98	0	< 0.001
	Physical Functioning *	82.52 ±10.64	99.30 ±2.52	< 0.001
RAND 36 Item	Emotional Wellbeing *	69.85 ±17.33	82.46 ±7.58	0.024
Health Survey	Pain *	68.46 ±16.79	95.00 ±8.6	< 0.001
	General Health *	64.62 ±20.15	82.31 ±10.53	0.010

Table 1 – Mean participant characteristics separated by group, showing the standard deviation where
 appropriate. Where significant differences occur, the characteristic is marked with an asterisk and a P-value
 is displayed.

586 Figures

Figure 1 – Depicting (A) the approximate positioning of the HDEMG grid 2cm lateral to the L5 Spinous
process on the lumbar ES of the participant and (B) a schematic of the electrode grid showing the x- and yaxes, reference electrode and inter-electrode distance (not to scale).

Figure 2 – The mean values for the initial, mid-point and final values (and SE) for the RPE as reported by
 participants during the endurance task. No significant differences were found between groups for exertion
 during the task.

Figure 3 - Average RMS values for LBP and CON participants across the duration of the endurance
contractions (and SE), shown in 10% epochs of the participants' total endurance times. No interactions or
differences in shift were found between groups, but the CON group was found to be systematically higher
throughout the contraction.

Figure 4 - Representative RMS topographical maps for CON (A) and LBP (B) participants during the endurance task. The centroid is depicted by the crosshair and the scale is indicated in μ V.

Figure 5 - Absolute mean locations (and SE) of the y-coordinate of the centroid for CON and LBP groupthroughout the endurance contraction

Figure 6 - Displacement of the y-coordinate of the centroid from the position in the first 10% epoch (and
SE), showing a significant displacement of the y-coordinate for both the CON and LBP group.

Figure 7 - Linear regression analysis on the shift in y-coordinate of the centroid, showing significant
 variation in the shift of the y-coordinate over the length of the endurance contraction (F=5.597, P=0.0183).
 Two CON points where shift was more than 6mm not shown.