

**Heightened pain facilitation rather than impaired pain inhibition distinguishes those with moderate/severe disability in work-related neck pain**

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**Published**

2021

**Journal Title**

Pain

**Version**

Submitted Manuscript (SM)

**DOI**

[10.1097/j.pain.0000000000002213](https://doi.org/10.1097/j.pain.0000000000002213)

**Downloaded from**

<http://hdl.handle.net/10072/402114>

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## 1 **1. Introduction**

2 Neck pain spreading to the shoulder region is a prevalent occupational health issue imposing a  
3 substantial economic burden on workers and organizations.<sup>48</sup> This presentation is commonly  
4 termed work-related neck pain and affects 27% to 48% of the general working population  
5 annually,<sup>11</sup> with an even higher prevalence among workers in high-risk occupations. Among  
6 sonographers, whose work involves ultrasound scanning with sustained static postures and  
7 forces and intensive computer-based tasks, it is estimated that 84% experience neck pain, with  
8 55% reporting at least mild disability.<sup>67</sup> Around one in five sonographers prematurely end their  
9 careers due to neck disability and other musculoskeletal disorders.<sup>46</sup> Despite significant  
10 advancements in management, the global cause of disability-adjusted life years for neck pain  
11 has increased rather than decreased over the past years.<sup>48</sup> A potential reason could be the lack  
12 of understanding of the underlying pain mechanisms;<sup>66</sup> increasing knowledge about pain  
13 mechanisms will enable individualized care and improved health outcomes.<sup>6, 55, 60</sup>

14 Quantitative sensory testing (QST) is a non-invasive method to examine neurobiological  
15 pain mechanisms. QST has been used to assess contributions of somatosensory and central pain  
16 modulatory function by analysing an individual's response to innocuous and noxious stimuli.<sup>20</sup>  
17 Thermal and/or mechanical hyperalgesia has been demonstrated over painful neck and shoulder  
18 regions in different occupational groups, such as office workers<sup>29</sup> violin players,<sup>57</sup> butchers,<sup>42</sup>  
19 and secretaries,<sup>24</sup> reflecting sensitization of nociceptive neurons. Our systematic review and  
20 meta-analysis<sup>68</sup> suggested that individuals with non-traumatic neck pain, particularly those  
21 with moderate/severe disability, have widespread hyperalgesia at a remote non-painful site,  
22 indicating nociplastic pain mechanisms involving altered nociceptive processing within the  
23 central nervous system (e.g. central sensitization).<sup>27</sup> However, more research is needed to  
24 confirm which specific central pain modulation mechanisms are involved.<sup>68</sup> Dynamic QST  
25 paradigms such as conditioned pain modulation (CPM), exercise-induced analgesia (EIA), and

1 temporal summation of pain (TSP) can provide more precise and valuable information  
2 regarding pain inhibitory and facilitatory function.<sup>13</sup> Only two studies have examined CPM in  
3 individuals with work-related neck pain, with mixed results.<sup>26,53</sup> Reduced efficacy of EIA has  
4 been shown in people with shoulder myalgia, evidenced by a decreased pressure pain threshold  
5 during and after an isometric exercise of the painful muscle.<sup>35</sup> Heightened TSP has been  
6 reported in other musculoskeletal conditions such as severe knee pain,<sup>50</sup> low back pain<sup>18</sup> and  
7 temporomandibular disorders.<sup>34</sup> Investigation using a more comprehensive battery of  
8 assessment on somatosensory and central pain modulatory function in a high-risk occupation  
9 such as sonographers is warranted to elucidate the pain mechanisms involved and how this  
10 relates to the level of disability.

11 Individuals with more severe neck disability are at greater risk of prolonged recovery and  
12 unfavourable prognosis.<sup>64</sup> Comparisons of subgroups might offer novel insights into the  
13 pathophysiology underlying poor recovery in individuals with more severe disability and help  
14 direct treatments to minimize the personal and economic burden of neck disability. This study  
15 aimed to comprehensively document the somatosensory profile in sonographers with different  
16 neck disability levels. We hypothesized that sonographers with more severe disability would  
17 present thermal and mechanical hyperalgesia at local painful and remote non-painful sites as  
18 well as altered central pain modulation compared to those with no or mild disability.

## 19 **2. Methods**

### 20 ***2.1. Study design***

21 Reporting of this cross-sectional and single-blinded study adheres to the Strengthening the  
22 Reporting of Observational studies in Epidemiology (STROBE) guidelines. Ethical approval  
23 was obtained from the University of Queensland Human Research Ethics Committee (approval

1 #2017001513). The study procedure was clearly explained to participants and signed consent  
2 obtained.

### 3 **2.2. Setting and participants**

4 From June 2018 to August 2019, participants with or without neck pain in the past 12 months  
5 were recruited prospectively from 430 volunteers who participated in an online survey  
6 examining risk factors of pain and disability in sonographers.<sup>67</sup> The sample of the survey were  
7 sonographers from Australia and New Zealand who performed sonography  $\geq 4$  hours/week,  
8 were not pregnant, without fibromyalgia, uncontrolled diabetes or kidney diseases, or without  
9 chemotherapy or radiotherapy for cancers within the past five years. Additional eligibility  
10 criteria for the current study were applied through a telephone interview. Participants were  
11 eligible if they had pain in the neck region, as defined by The Neck Pain Task Force<sup>23</sup>, as the  
12 primary musculoskeletal symptom. Participants were also eligible if they had no neck pain or  
13 severe pain in the low back or upper or lower limbs that required treatment or interfered with  
14 work or home activities in the previous three months. Other general exclusion criteria were a  
15 history of surgery in the spine or upper limbs, trauma (for example, whiplash), irritable bowel  
16 syndrome, inflammatory conditions, neurological disorders or psychiatric disorders. These  
17 additional exclusion criteria were to exclude possible non-work-related sources of neck pain  
18 and confounding conditions that may present somatosensory changes. All participants were  
19 asked to refrain from taking non-steroidal anti-inflammatory drugs or analgesics 24 hours prior  
20 to testing.

### 21 **2.3. Sample size**

22 An estimated total sample size of 111 participants would be required to detect group differences  
23 in pressure pain threshold (PPT), based on calculations undertaken using G\*Power (version  
24 3.1.9.2) with an effect size of 0.30, a significance of 0.05 and a power of 0.80. The effect size

1 was computed using data of PPT measurements derived from two previous studies,<sup>4, 29</sup> which  
2 were the only studies available in the literature on comparing QST across varied levels of neck  
3 disability.

#### 4 **2.4. Measurements**

5 Data collection was performed in a quiet and temperature-controlled room by one researcher  
6 (YX) without knowing the disability levels of participants at the time of measurement. Pilot  
7 testing had been performed on six volunteers who were not sonographers to ensure adequate  
8 intra-rater reliability of QST measures (all intraclass correlation coefficients were  $\geq 0.83$  with  
9 lower bounds of 95% confidence of intervals (CIs)  $\geq 0.66$  and upper bounds  $\geq 0.88$ ). These  
10 volunteers were not included in the study analyses.

##### 11 *2.4.1. Quantitative sensory testing*

12 To control for any potential effect of one test on subsequent tests,<sup>52</sup> QST was conducted in the  
13 following order: thermal pain threshold, TSP, CPM, PPT and EIA (**Supplementary 1**). These  
14 tests were separated by 5 to 40 minutes to avoid sensitization or habituation due to repetitive  
15 stimuli (**Supplementary 1**). Tests were performed unilaterally at the most affected side for  
16 symptomatic participants or at the scanning hand side for asymptomatic participants. Prior to  
17 data collection, a familiarization trial was performed over an area that was different to the tested  
18 sites to ensure that participants clearly understood the tests. All test stimuli were applied for  
19 three consecutive trials with 30-second intervals and the averages were used for analysis.  
20 Testing protocols and instructions used were based to the German Research Network for  
21 Neuropathic Pain QST protocol.<sup>51</sup>

##### 22 *2.4.1.1. Thermal pain threshold*

1 Cold pain threshold (CPT) and heat pain threshold (HPT) were measured using a Thermostest  
2 unit (Somedic AB, Sweden), with a 2.5 cm x 5 cm thermode probe. Thermal pain thresholds  
3 have shown good to excellent test-retest reliability<sup>44</sup> and are reproducible over 6-9 months in  
4 healthy people.<sup>43</sup> Participants were in a prone or supine lying position for this test. The CPT  
5 was applied in the following order: the neck (cervical spine at the C5-6 level), upper trapezius  
6 (the mid-way between the 7<sup>th</sup> cervical vertebrae and the acromion), the lateral deltoid  
7 (approximately the mid-point between acromion and the deltoid tuberosity), and a remote site  
8 which was the tibialis anterior (approximately 2.5 cm lateral and 5 cm inferior to the tibial  
9 tuberosity).<sup>2, 10, 65</sup> The HPT was then tested on the same sites in the reverse order to avoid  
10 frequent changes of position. For both CPT and HPT, the temperature started at 32°C and  
11 changed at a rate of 1°/s,<sup>51</sup> with cut-off temperatures of 5°C and 50°C to reduce the risk of  
12 frostbite or burns. Participants were instructed to press a stop button if/when they perceived  
13 the cool or warm sensation first became painful to identify CPT and HPT. If the cut-off  
14 temperatures were reached before participants felt the first pain, these were used for analysis.

#### 15 *2.4.1.2. Temporal summation of pain*

16 A single stimulus and then a train of 10 repetitive stimuli at a frequency of 1 Hz (monitored  
17 with a metronome) were applied over an area of 1 cm<sup>2</sup> of the cervical spine at the C5-6 level  
18 using a pinprick stimulator (256mN). The TSP was determined by the difference in perceived  
19 pain on a 0-100 point Numeric Rating Scale (NRS) induced by the single stimulus and the train  
20 of 10 repetitive pinprick stimuli.<sup>28, 50</sup> Reliability of TSP has been demonstrated to be fair to  
21 good.<sup>44</sup>

#### 22 *2.4.1.3. Conditioned pain modulation*

23 The CPM paradigm is based on the “pain-inhibits-pain” mechanism, in which a reduction of  
24 pain perception from a test stimulus is induced by a concurrent application of a noxious

1 stimulus (conditioning stimulus) at another region of the body.<sup>69</sup> Reliability of this method is  
2 fair to excellent.<sup>30</sup> The test stimulus was PPT (described below) at the upper trapezius of the  
3 most affected side. The conditioning stimulus was immersion of the contralateral hand (up to  
4 the wrist) in a circulating cold-water bath (Polyscience, 912, 6L Basic) maintained at  $10 \pm 2^\circ\text{C}$   
5 for a maximum of two minutes. Measurements were performed with participants in a sitting  
6 position. Prior to hand immersion into the cold water, the PPT was tested for three trials with  
7 a 30-second interval between trials. While the hand was immersed in the cold water,  
8 participants were asked to rate the pain level of the hand every 10-seconds using a 0-100 NRS  
9 until the pain level reached at or above 40 out of 100 on the NRS, after which re-evaluation of  
10 PPT on the same site commenced immediately. A cold pain rating of 40 was selected on the  
11 basis that a conditioning stimulus of at least moderate intensity is needed to induce CPM.<sup>49</sup> At  
12 least one PPT measurement was performed every 30-seconds (maximum of three  
13 measurements) while the participant immersed the hand into the cold water before the end of  
14 two minutes. Otherwise, the measurement was regarded as missing.<sup>9</sup> Participants were asked  
15 to rate their pain levels induced by the cold water again before the 2<sup>nd</sup> and 3<sup>rd</sup> PPT  
16 measurements. The CPM efficacy was calculated using the following formula: CPM efficacy  
17 = (average PPT during conditioning stimulus – average PPT before conditioning stimulus). A  
18 positive value reflects endogenous pain inhibition, and the higher the value, the more efficient  
19 an individual's pain inhibition.

#### 20 2.4.1.4. *Pressure pain threshold*

21 A hand-held digital algometer (Somedic Production AB) with a probe size of  $1\text{ cm}^2$  was used  
22 to measure PPT at a rate of 50kPa/s perpendicularly over the same regions that the thermal pain  
23 thresholds were tested. Participants were asked to press a stop button once the sensation of  
24 pressure first becomes pain. Measurement of PPT with a hand-held digital algometer has  
25 demonstrated good to excellent reliability.<sup>44</sup>

1    2.4.1.5. *An exercise-induced analgesia paradigm*

2    Measurement of pain sensitivity following exercise has been used to evaluate endogenous pain  
3    modulation function in humans.<sup>8, 22, 35</sup> The participant's pain response and sensitivity to a  
4    pressure stimulus were examined during a standardized exercise. Participants sat on a chair  
5    with the trunk upright and performed sustained isometric bilateral shoulder abduction at 90°  
6    without external weights until exhaustion (reached to maximum rating of perceived exertion  
7    on a scale of 0 to 10 with inability to maintain shoulder abduction angle of 90°) or up to 3  
8    minutes. Participants rated their pain intensity of the neck-shoulder region using a 0-100 NRS  
9    before and at the end of the exercise. The PPT was measured at the neck, upper trapezius,  
10   deltoid and tibialis anterior before and immediately after the exercise. The EIA was calculated  
11   using the following formula:  $EIA = (\text{average PPT after exercise} - \text{average PPT before exercise})$ .  
12   Thus a positive change in PPT reflected EIA<sup>56</sup> and the higher the value, the more EIA efficacy.  
13   Changes of pain intensity before and after the exercise were also calculated.

14   2.4.2. *Pain, disability and psychological measures*

15   On completion of the QST, participants completed standard questionnaires to collect data on  
16   demographics (e.g. age, gender, body mass index (BMI), physical activity assessed using the  
17   International Physical Activity Questionnaire-Short Form<sup>12</sup>), work-related information, pain  
18   characteristics (e.g. frequency, sick leave and healthcare seeking behavior), and psychological  
19   measures.

20       As sonographers usually present with pain in multiple sites,<sup>67</sup> a novel digital pain drawing  
21   method was used to quantify the location and extent of pain. Participants with symptoms were  
22   asked to shade precisely every area of pain they typically perceived and had experienced in the  
23   previous week, regardless of the intensity, on either female or male body charts displayed on a  
24   digital tablet (iPad 6, Apple Inc, Cupertino, California) with a stylus pen.<sup>3</sup> The characteristics



1 of the stylus pen including the type, size, and colour were standardized. Pain extent (numbers  
2 of pixels shaded inside the body chart expressed as a percentage of the total body chart area)  
3 was calculated to quantify the percentage of total (frontal and dorsal body regions) pain area.<sup>3</sup>  
4 Pain frequency maps were generated to illustrate where pain was most frequently reported. The  
5 worst pain level of the shaded body regions was measured using the 0-100 NRS. Disability  
6 related to neck pain was assessed using the Neck Disability Index (NDI),<sup>61</sup> with a total score  
7 ranging from 0 to 100 (expressed as a percentage). Higher scores indicate greater disability.

8 Psychological factors assessed included depression, anxiety, pain catastrophizing and  
9 fear-avoidance beliefs. Depression and anxiety were assessed using the 8-item Patient Health  
10 Questionnaire<sup>33</sup> and Generalized Anxiety Disorder – 7-item scale,<sup>32</sup> respectively. Total  
11 depression scores ranged from 0 to 24 while anxiety scores ranged from 0 to 21, with a score  
12 of  $\geq 5$  being a cut-off score for depression and anxiety.<sup>31, 32</sup> Pain catastrophizing was evaluated  
13 by the Pain Catastrophizing Scale, with a total score ranging from 0 to 52.<sup>58</sup> Fear-avoidance  
14 beliefs were assessed with two items from the Fear-Avoidance Beliefs Questionnaire<sup>63</sup> (“my  
15 work is too heavy for me” and my work might harm my neck”), with a total score ranging from  
16 0 to 12.

## 17 **2.5. Statistical analysis**

18 All analyses were conducted using R (Version 3.4.2), and the significance level was set at 0.05.

### 19 *2.5.1. Group classification*

20 K-mean cluster analysis was performed to classify participants into 3 groups. This is a  
21 commonly used unsupervised machine learning algorithm for classifying a given dataset into  
22 a predefined number of subgroups (clusters) (K=3 in this study), such that the data points within  
23 the same subgroup are as similar as possible while data points between subgroups are as  
24 different as possible.<sup>41</sup> To better represent the population of working sonographers, NDI scores

1 obtained from the original full sample (430 sonographers who participated in the online survey)  
2 were used to derive the clusters. The cut-off scores for the 3 clusters were: NDI  $\leq$ 8% (no  
3 disability group), NDI=10% - 20% (mild disability group), and NDI  $\geq$ 22% (moderate/severe  
4 disability group). These cut-off scores were replicated when using NDI scores from the current  
5 laboratory sample (92 sonographers). The K-mean cluster analysis was performed because the  
6 conventional NDI cut-off scores for no, mild and moderate/severe disability have not been  
7 validated for non-traumatic neck pain,<sup>39</sup> and may not help to identify homogenous groups of  
8 sonographers based on neck disability. Sonographers with no disability whose neck pain might  
9 not be clinically meaningful was used as a reference group, because previous research found a  
10 high annual prevalence (84%) of neck pain in sonographers,<sup>67</sup> which made it difficult to recruit  
11 a true pain-free healthy control group.

#### 12 2.5.2. *Statistical analysis for group comparisons*

13 Chi-square or Fisher's exact test was performed to compare group differences in categorical  
14 data. For continuous data, normality was checked using the Shapiro-Wilk test and visual  
15 inspection of quantile-quantile plots. Both parametric and non-parametric analyses were  
16 conducted for continuous data that were not normally distributed.<sup>40</sup> Parametric analysis was  
17 used because results were the same across analyses and it enables us to develop multivariate  
18 models and compute effect size (ES) (described below). Furthermore, multivariate and  
19 univariate analysis of variance (MANOVA and ANOVA) are reported to be robust to modest  
20 departures from normality when there were at least 20 degrees of freedom for error in ANOVA  
21 and the violations were not due to outliers.<sup>59</sup> Outliers were examined for each dependent  
22 variable and one to two outliers were found for TPS, CPM, and PPT. These outliers were  
23 carefully checked to ensure that it was not due to encoding error. To avoid possible data  
24 manipulation, outliers were retained in the analyses as results were the same with and without  
25 the outliers.

1 Multiple MANOVAs were performed to compare group differences in variables that are  
2 related conceptually, such as psychological variables (anxiety, depression, pain catastrophizing  
3 and fear-avoidance beliefs), pain thresholds at local sites (the model included cold, heat and  
4 pressure pain thresholds at the neck, upper trapezius and deltoid), and pain thresholds at the  
5 remote site (cold, heat and pressure pain thresholds at the tibialis anterior). When MANOVA  
6 showed a significant result, post-hoc ANOVA was performed to analyse which specific  
7 variable significantly differed between groups. Variables that did not fit in the MANOVA (e.g.  
8 demographics, pain, disability, TSP, CPM, EIA) were analysed by ANOVA. As pain ratings  
9 of the conditioning stimulus were different between groups, ANOVA of CPM was performed  
10 with and without the inclusion of this variable. Considering the potential effect of  
11 psychological variables on QST,<sup>21, 45, 47, 71</sup> models for all QST outcomes were performed with  
12 and without the inclusion of each psychological variable as a covariate. When significant group  
13 differences were found, pairwise comparison was performed with Bonferroni correction, and  
14 ESs and 95% CIs were computed using Hedge's *g* for pairwise group comparisons. An ES was:  
15  $<0.20$  = negligible;  $0.20 - 0.49$  = small;  $0.50 - 0.79$  = moderate; and  $\geq 0.80$  = large effect.<sup>7</sup>

### 16 **3. Results**

17 Ninety-two sonographers were recruited; 31 were classified into the no disability group (NDI  
18 =  $4.06\% \pm 2.90\%$ ), 43 into mild disability group (NDI =  $15.00\% \pm 3.22\%$ ) and 18 into  
19 moderate/severe disability group (NDI =  $30.60\% \pm 13.70\%$ ) (**Figure 1**). These three groups  
20 were comparable ( $P > 0.30$  for all demographics) in age (whole group mean  $\pm$  SD:  $39.91 \pm$   
21  $10.70$  years), gender (83.7% female), BMI ( $23.59 \pm 3.65$  kg/m<sup>2</sup>), physical activity levels  
22 ( $2431.49 \pm 1918.35$  MET-minutes/week) and work-related characteristics (sonography  
23 experience:  $13.37 \pm 10.32$  years; scanning time:  $25.88 \pm 9.52$  hours/week). Almost all  
24 sonographers in the mild and moderate/severe disability groups reported pain in the neck region  
25 at least once a week and a higher proportion of them had sick leave in the past three months,

1 compared to the no disability group (**Table 1**). Regarding psychological characteristics, the  
2 moderate/severe disability group scored significantly higher than no and mild disability groups  
3 on questionnaires of anxiety, depression and pain catastrophizing (**Table 1**). For fear-avoidance  
4 beliefs, significant differences were only found between moderate/severe and no disability  
5 groups (**Table 1**).

### 6 **3.1. Pain distribution and intensity in the previous week**

7 The percentage of participants that reported pain in a specific body region is illustrated using a  
8 heat map in **Figure 2**, with the most frequently reported areas displayed in red. The total area  
9 of bodily pain (as a percentage) was significantly different between all three groups ( $F_{(2, 89)} =$   
10  $18.46, P < 0.01$ ), ranging from  $1.23\% \pm 1.55\%$  (no disability),  $4.68\% \pm 3.21\%$  (mild disability)  
11 and  $7.20\% \pm 5.75\%$  (moderate/severe disability). Worst pain intensity was also significantly  
12 different between all three groups for the neck, shoulder, upper and lower back (**Figure 3**). For  
13 the elbow and wrist/hand, only the moderate/severe group had higher pain intensity than the  
14 no disability group (**Figure 3**). There were no group differences for pain intensity for lower  
15 limb regions.

### 16 **3.2. Thermal and pressure pain thresholds**

17 Significant group differences were only found in CPT ( $F_{(2, 88)} = 6.22, P < 0.01$ ) and PPT ( $F_{(2,$   
18  $89)} = 4.40, P = 0.02$ ) at the tibialis anterior, although there was a trend of group differences at  
19 local sites (**Figure 4**). Pairwise comparisons revealed higher CPT and lower PPT (cold and  
20 mechanical hyperalgesia) at the tibialis anterior in the moderate/severe disability group,  
21 compared to mild disability with moderate effect sizes (CPT: ES (95%CI) = 0.66 (0.09, 1.22);  
22 PPT: -0.68 (-1.24, -0.12)) and compared to no disability groups with large effect sizes (CPT:  
23  $1.06 (0.44, 1.68)$ ; PPT:  $-1.00 (-1.61, -0.39)$ ) (**Supplementary 2**). No group differences were  
24 found in HPT across all tested sites (**Figure 4**).

### 1 **3.3. Temporal summation of pain**

2 There were significant group differences in TSP ( $F_{(2, 89)} = 5.42, P = 0.01$ ), with the  
3 moderate/severe disability group reporting highest TSP (**Figure 5A**). Post-hoc analysis  
4 revealed that the between-group effect was moderate between mild and no disability groups  
5 (ES (95% CI) = 0.67 (0.19, 1.14)), and large between moderate/severe and no disability groups  
6 (0.94 (0.33, 1.54)) (**Supplementary 2**).

### 7 **3.4. Conditioned pain modulation**

8 There were no group differences in the time of exposure to the conditioning stimulus ( $F_{(2, 89)}=0.77, P=0.47$ ), with mean  $\pm$  SD time being  $109.60 \pm 23.02$  seconds for the whole sample.  
9  
10 The mean  $\pm$  SD hand pain ratings (0-100 NRS) during the conditioning stimulus for no, mild  
11 and moderate/severe disability groups were  $64.30 \pm 13.55, 67.85 \pm 11.81$ , and  $74.07 \pm 11.32$ ,  
12 respectively (**Supplementary 2**). There were significant group differences in hand pain rating  
13 ( $F_{(2, 89)} = 3.58, P = 0.03$ ), with the moderate/severe disability group rating significantly higher  
14 pain levels than the no disability group (ES (95%CI) = 0.76 (0.17, 1.36)). Almost all  
15 participants from all groups had increased PPT (decreased mechanical sensitivity) at the upper  
16 trapezius during exposure to the conditioning stimulus (**Figure 5B**). However, no group  
17 differences were found in changes in PPT ( $F_{(2, 89)}=2.35, P=0.10$ ), and the result remained the  
18 same after including hand pain rating as a covariate.

### 19 **3.5. Exercise-induced analgesia**

20 All participants completed the 3-minute exercise, except one in the moderate/severe disability  
21 group who had to interrupt after 2 minutes due to reaching the maximum rating of perceived  
22 exertion. The pain scores prior to exercise were  $5.13 \pm 8.89, 9.51 \pm 9.69$  and  $27.22 \pm 21.57$ ,  
23 with an increase of pain (0-100 NRS) of  $21.55 \pm 22.22, 38.28 \pm 20.39$ , and  $31.78 \pm 17.80$  after

1 exercise for the no, mild and moderate/severe disability groups, respectively. The mild  
2 disability group had significantly higher magnitude of pain increase than the no disability group  
3 (ES (95%CI) = 0.78 (0.30, 1.27)). All groups had increased PPT (decreased mechanical  
4 sensitivity) at all tested sites after exercise, but no significant group differences were found in  
5 changes of PPT at all tested sites (**Figure 6**).

### 6 **3.6. Role of psychological measures as covariates**

7 Group differences in thermal and pressure pain thresholds at the tibialis anterior changed from  
8 statistically significant ( $F_{(6, 174)} = 2.84, P = 0.01$ ) to non-significant when each of the following  
9 psychological variables were included as a covariate in the MANOVA: anxiety ( $F_{(6, 172)} = 2.03,$   
10  $P = 0.06$ ), depression ( $F_{(6, 172)} = 1.72, P = 0.119$ ), pain catastrophizing ( $F_{(6, 172)} = 2.13, P = 0.05$ ),  
11 or fear-avoidance belief ( $F_{(6, 172)} = 2.11, P = 0.06$ ). Results of TSP remained similar after  
12 including depression, pain catastrophizing or fear-avoidance beliefs as a covariate. However,  
13 group differences in TSP changed from statistically significant to non-significant after  
14 including anxiety ( $F_{(2, 88)} = 2.48, P = 0.09$ ) as a covariate. Results of other QST variables did  
15 not change when including psychological measures as covariates.

## 16 **4. Discussion**

17 This is the first study examining somatosensory features in sonographers with different levels  
18 of neck disability. The main finding was that sonographers with moderate/severe neck  
19 disability demonstrated significantly more widespread pain, cold and mechanical hyperalgesia  
20 at a remote site, and higher TSP, compared to those with mild or no disability. The mild  
21 disability group showed significantly higher TSP and pain intensity in response to an isometric  
22 exercise compared to the no disability group. All groups demonstrated normal CPM function  
23 and no significant group differences were found in efficacy of CPM and EIA. These findings

1 offer new insights regarding possible pain mechanisms that may underpin subgroups with  
2 work-related neck disability.

### 3 ***4.1. Comparison of QST between different neck disability levels***

4 Multi-modal hyperalgesia at a site remote from the painful site, as reflected by increased  
5 sensitivity to cold and pressure pain at the tibialis anterior, was found in the moderate/severe  
6 neck disability group compared to both no and mild disability groups. This could be considered  
7 as evidence of nociplastic pain processing characterized by sensitization within the central  
8 nervous system.<sup>14, 19</sup> Although all groups included some participants reporting pain in the lower  
9 limb, the mean pain levels were lower than 10 out of 100 on a NRS (**Figure 3**), which is not  
10 clinically relevant. Furthermore, no group differences in lower limb symptoms were observed  
11 between groups. Therefore, cold and mechanical hyperalgesia at the tibialis anterior is unlikely  
12 to be due to peripheral sensitization of nociceptive neurons. Contrary to our hypothesis,  
13 localized hyperalgesia was not statistically evident in sonographers with mild or  
14 moderate/severe disability, although there was a trend of group differences in CPT and PPT at  
15 the local symptomatic sites. Similar phenomena was observed in other occupations such as  
16 pianists<sup>37</sup> and office workers<sup>26, 29</sup> with neck pain in that hyperalgesia was only found at the  
17 remote sites but not the symptomatic neck-shoulder region. It is possible that the localized  
18 neck-shoulder region has been sensitized in sonographers with no disability due to long-term  
19 exposure to repetitive ultrasound scanning with awkward neck and shoulder postures and static  
20 muscle force, making it more difficult to find statistically significant group differences in CPT  
21 and PPT at local sites. Further investigation of localized hyperalgesia by comparing  
22 sonographers with non-worker healthy controls is warranted to confirm this hypothesis.

23 The presence of nociplastic pain processing in sonographers with moderate/severe  
24 disability is further supported by the finding of a large magnitude of heightened TSP, a human  
25 surrogate model involving a wind-up in the dorsal horn and activation of N-methyl-D-aspartate

1 receptor.<sup>17</sup> In contrast, no evidence was found for impaired CPM in all groups, indicating  
2 endogenous pain inhibition function may be normal in sonographers. Similarly, at the group  
3 level, EIA seemed to be normal in sonographers with moderate/severe disability as decreased  
4 mechanical sensitivity was found after the 3-minute shoulder isometric exercise and changes  
5 in pain rating after exercise did not differ with the no disability group. The EIA efficacy as  
6 demonstrated by changes of PPT had high standard deviations, indicating that some  
7 sonographers may present impaired EIA, but this is unlikely a common feature of sonographers  
8 with disability. Collectively, our findings provide novel evidence that heightened pain  
9 facilitation rather than impaired pain inhibition may underpin the nociplastic pain processing  
10 in sonographers with moderate/severe neck disability.

11 Compared to sonographers with moderate/severe neck disability, signs of central  
12 involvement in those with mild disability is less clear. Sonographers with mild disability did  
13 not show statistically evident thermal and mechanical hyperalgesia compared to those with no  
14 disability. In contrast, they displayed a moderate magnitude of higher TSP, indicating that  
15 heightened pain facilitation may be present, at least for some individuals in this subgroup.  
16 Additionally, sonographers with mild disability showed significantly higher increase in pain  
17 perception during the 3-minute shoulder isometric exercise task compared to those with no  
18 disability. Exercise-induced pain and hyperalgesia were found in fibromyalgia<sup>16</sup> and chronic  
19 whiplash associated disorders,<sup>56</sup> and have been linked to changes in central nervous system  
20 function.<sup>36</sup> However, whether the increased pain perception to exercise points to central  
21 mechanisms in sonographers with mild disability group, or simply reflects a discrepancy in  
22 pain perception compared to other groups, requires further investigation.

#### 23 ***4.2. Potential effect of pain distribution and psychological measures on QST***

24 According to a mechanisms-based classification list from Smart et al,<sup>54</sup> ‘diffuse or widespread  
25 areas of pain’ and ‘maladaptive psychosocial factors’ have been recognized as key signs and



1 symptoms in classifying central sensitization. This is also supported by recent study, involving  
2 patients with knee osteoarthritis, which showed that widespread pain is associated with signs  
3 and symptoms of central sensitization.<sup>38</sup> In our study, sonographers with mild and  
4 moderate/severe disability groups demonstrated more widespread pain than those with no  
5 disability, and those with moderate/severe disability also presented significantly more  
6 widespread pain compared to those with mild disability. This may explain why the  
7 moderate/severe disability group displayed more somatosensory changes associated with  
8 central sensitization while the mild disability group showed a lesser extent of central changes.  
9 However, there was a large amount of variation in the total area of bodily pain within each  
10 group as reflected by the broad standard deviations, suggesting that the area of bodily pain may  
11 not exclusively account for the group differences seen in the somatosensory changes.

12         Sonographers with moderate/severe disability also showed more psychological  
13 impairments when compared to those with mild and no disability, and this was particularly the  
14 case for anxiety and depression. When scores were dichotomized according to the  
15 recommended cut-off for anxiety and depression,<sup>31, 32</sup> it was found that 61% and 78% of  
16 sonographers in the moderate/severe disability group presented anxious and depressive  
17 symptoms. Group differences in CPT and PPT at the tibialis anterior and TSP were attenuated  
18 and approached significance when assessed psychological factors were entered as covariates  
19 in the analysis. This contrasts with previous studies of people with whiplash-associated  
20 disorders which showed that somatosensory changes were independent of psychological  
21 impairments.<sup>5, 52</sup> In our sample, individuals with trauma were excluded and different  
22 psychological constructs were studied using different questionnaires, which may explain the  
23 discrepancy. Our study suggests that somatosensory changes identified in sonographers with  
24 disability may be partially mediated by psychological factors. However, the direction of this  
25 relationship cannot be established in a cross-sectional design.

1 **4.3. Clinical implications**

2 Our findings have important clinical implications for guiding the design of management  
3 strategies for neck pain in sonographers where the current focus has largely been on ergonomic  
4 interventions.<sup>1, 25</sup> Management might need to address the somatosensory alterations and tailor  
5 interventions to individual sonographers with different disability levels. The findings of  
6 heightened pain facilitation without widespread hyperalgesia in sonographers with mild  
7 disability, suggest that this subgroup is an important group to focus strategies to reduce the  
8 progression to more severe disability. For this group, clinicians should consider that repetitive  
9 noxious or non-noxious stimuli (e.g. by sustained isometric contraction) may lead to  
10 summation of pain. As sonographers typically perform sustained shoulder abduction during  
11 their typical scanning activities,<sup>62</sup> it may be important to monitor their pain levels and institute  
12 frequent periods of rest between scans. Furthermore, activity-based treatment strategies may  
13 need modification to account for increase of pain during a single bout of isometric exercise for  
14 sonographers with mild disability. It might be important to start exercising asymptomatic body  
15 regions to activate endogenous pain inhibitory mechanisms<sup>35</sup> and then progress to exercising  
16 the neck and shoulder muscles following the principles of progressive overload and  
17 periodization with adequate recovery periods.<sup>15, 70</sup> For sonographers with moderate/severe  
18 disability, additional multidisciplinary therapeutic approach including interventions aiming to  
19 reduce anxiety, depression, and maladaptive beliefs about their pain, as well as graded exposure  
20 therapy might be appropriate. Further investigation is needed to identify optimal management  
21 strategies for different subgroups of sonographers to improve disability and reduce professional  
22 attrition.

23 **4.4. Strength and limitations**

1 A strength of this study is that the investigator who conducted QST was not aware of the group  
2 allocation, thus reducing study bias. Moreover, a broad set of QST across multiple modalities  
3 and sites was conducted to characterize the somatosensory profiles. Some limitations should  
4 be considered. First, QST was conducted in a fixed order without randomization which may  
5 increase study bias. However, pain ratings were repeatedly asked to ensure no systematic  
6 increase in pain during the testing occurred and there were sufficient intervals between tests  
7 which should minimize the sensitization due to repeated application of multiple stimuli. Second,  
8 not all sonographers with no disability were completely pain-free and this may have resulted  
9 in an underestimation of differences in somatosensory features in sonographers with varied  
10 disability levels. Third, due to the difficulty in recruiting sonographers with moderate/severe  
11 neck disability, the current sample size was smaller than the estimated size needed, which may  
12 have reduced statistical power potentially explaining the lack of group differences in other  
13 outcomes such as pain thresholds at local sites. Finally, the cause-effect relationship of  
14 observed impairments could not be elucidated from this study due to the nature of a cross-  
15 sectional design.

## 16 **5. Conclusions**

17 This study showed that more widespread pain, remote cold and mechanical hyperalgesia and  
18 heightened TSP, but not dysfunctional CPM and EIA, distinguish sonographers with  
19 moderate/severe neck disability from no disability. Sonographers with mild neck disability  
20 presented a moderate effect of heightened TSP and increased pain ratings in response to  
21 exercise compared to those with no disability. Psychological factors, particularly anxiety and  
22 depression, were found to influence the results of group differences in somatosensory features.  
23 Specific somatosensory changes and psychological factors should be considered when tailoring  
24 management strategies for sonographers with different levels of disability.

1 **Conflict of Interest:** There are no conflicts of interest.

## 1 **References**

- 2 [1] Industry standards for the prevention of work related musculoskeletal disorders in  
3 sonography. *JDMS* 2017;33:370-91.
- 4 [2] Arendt-Nielsen L, Nie H, Laursen MB, Laursen BS, Madeleine P, Simonsen OH,  
5 Graven-Nielsen T. Sensitization in patients with painful knee osteoarthritis. *Pain*  
6 2010;149:573-81.
- 7 [3] Barbero M, Moresi F, Leoni D, Gatti R, Egloff M, Falla D. Test-retest reliability of pain  
8 extent and pain location using a novel method for pain drawing analysis. *Eur J Pain*  
9 2015;19:1129-38.
- 10 [4] Beltran-Alacreu H, López-de-Uralde-Villanueva I, Calvo-Lobo C, Fernández-Carnero J,  
11 La Touche R. Clinical features of patients with chronic non-specific neck pain per  
12 disability level: A novel observational study. *Rev Assoc Med Bras (1992)* 2018;64:700-  
13 9.
- 14 [5] Chien A, Sterling M. Sensory hypoaesthesia is a feature of chronic whiplash but not  
15 chronic idiopathic neck pain. *Man Ther* 2010;15:48-53.
- 16 [6] Chimenti RL, Frey-Law LA, Sluka KA. A mechanism-based approach to physical  
17 therapist management of pain. *Phys Ther* 2018;98:302-14.
- 18 [7] Cohen J. A power primer. *Psychol Bull* 1992;112:155-9.
- 19 [8] Cook DB, Stegner AJ, Ellingson LD. Exercise alters pain sensitivity in Gulf War veterans  
20 with chronic musculoskeletal pain. *J Pain* 2010;11:764-72.
- 21 [9] Coppieters I, De Pauw R, Kregel J, Malfliet A, Goubert D, Lenoir D, Cagnie B, Meeus  
22 M. Differences between women with traumatic and idiopathic chronic neck pain and  
23 women without neck pain: interrelationships among disability, cognitive deficits, and  
24 central sensitization. *Phys Ther* 2017;97:338-53.

- 1 [10] Corrêa JB, Costa LO, de Oliveira NT, Sluka KA, Liebano RE. Central sensitization and  
2 changes in conditioned pain modulation in people with chronic nonspecific low back pain:  
3 a case-control study. *Exp Brain Res* 2015;233:2391-9.
- 4 [11] Côté P, van der Velde G, Cassidy JD, Carroll LJ, Hogg-Johnson S, Holm LW, Carragee  
5 EJ, Haldeman S, Nordin M, Hurwitz EL, Guzman J, Peloso PM. The burden and  
6 determinants of neck pain in workers: results of the Bone and Joint Decade 2000-2010  
7 Task Force on Neck Pain and Its Associated Disorders. *Spine (Phila Pa 1976)*  
8 2008;33:S60-74.
- 9 [12] Craig CL, Marshall AL, Sjöström M, Bauman AE, Booth ML, Ainsworth BE, Pratt M,  
10 Ekelund U, Yngve A, Sallis JF, Oja P. International physical activity questionnaire: 12-  
11 country reliability and validity. *Med Sci Sports Exerc* 2003;35:1396.
- 12 [13] Cruz-Almeida Y, Fillingim RB. Can quantitative sensory testing move us closer to  
13 mechanism-based pain management? *Pain Med* 2014;15:61-72.
- 14 [14] Curatolo M, Arendt-Nielsen L, Petersen-Felix S. Central hypersensitivity in chronic pain:  
15 mechanisms and clinical implications. *Phys Med Rehabil Clin N Am* 2006;17:287-302.
- 16 [15] Daenen L, Varkey E, Kellmann M, Nijs J. Exercise, not to exercise, or how to exercise  
17 in patients with chronic pain? Applying science to practice. *Clin J Pain* 2015;31:108-14.
- 18 [16] Dailey DL, Keffala VJ, Sluka KA. Do cognitive and physical fatigue tasks enhance pain,  
19 cognitive fatigue, and physical fatigue in people with fibromyalgia? *Arthritis Care Res*  
20 (Hoboken) 2015;67:288-96.
- 21 [17] Davies SN, Lodge D. Evidence for involvement of N-methylaspartate receptors in 'wind-  
22 up' of class 2 neurones in the dorsal horn of the rat. *Brain Res* 1987;424:402-6.
- 23 [18] den Bandt HL, Paulis WD, Beckwée D, Ickmans K, Nijs J, Voogt L. Pain mechanisms  
24 in low back pain: a systematic review with meta-analysis of mechanical quantitative

- 1 sensory testing outcomes in people with nonspecific low back pain. *J Orthop Sports Phys*  
2 *Ther* 2019;49:698-715.
- 3 [19] Fernández-Carnero J, Fernández-de-Las-Peñas C, de la Llave-Rincón AI, Ge HY,  
4 Arendt-Nielsen L. Widespread mechanical pain hypersensitivity as sign of central  
5 sensitization in unilateral epicondylalgia: a blinded, controlled study. *Clin J Pain*  
6 2009;25:555-61.
- 7 [20] Fillingim RB, Loeser JD, Baron R, Edwards RR. Assessment of chronic pain: domains,  
8 methods, and mechanisms. *J Pain* 2016;17:T10-20.
- 9 [21] Garrigós-Pedron M, La Touche R, Navarro-Desentre P, Gracia-Naya M, Segura-Ortí E.  
10 Widespread mechanical pain hypersensitivity in patients with chronic migraine and  
11 temporomandibular disorders: relationship and correlation between psychological and  
12 sensorimotor variables. *Acta Odontol Scand* 2019;77:224-31.
- 13 [22] Ge HY, Nie H, Graven-Nielsen T, Danneskiold-Samsøe B, Arendt-Nielsen L.  
14 Descending pain modulation and its interaction with peripheral sensitization following  
15 sustained isometric muscle contraction in fibromyalgia. *Eur J Pain* 2012;16:196-203.
- 16 [23] Guzman J, Hurwitz EL, Carroll LJ, Haldeman S, Côté P, Carragee EJ, Peloso PM, van  
17 der Velde G, Holm LW, Hogg-Johnson S, Nordin M, Cassidy JD. A new conceptual  
18 model of neck pain: linking onset, course, and care: the Bone and Joint Decade 2000-  
19 2010 Task Force on Neck Pain and Its Associated Disorders. *Spine (Phila Pa 1976)*  
20 2008;33:S14-23.
- 21 [24] Hägg GM, Aström A. Load pattern and pressure pain threshold in the upper trapezius  
22 muscle and psychosocial factors in medical secretaries with and without shoulder/neck  
23 disorders. *Int Arch Occup Environ Health* 1997;69:423-32.
- 24 [25] Harrison G, Harris A. Work-related musculoskeletal disorders in ultrasound: can you  
25 reduce risk? *Ultrasound* 2015;23:224-30.

- 1 [26] Heredia-Rizo AM, Petersen KK, Madeleine P, Arendt-Nielsen L. Clinical outcomes and  
2 central pain mechanisms are improved after upper trapezius eccentric training in female  
3 computer users with chronic neck/shoulder pain. *Clin J Pain* 2019;35:65-76.
- 4 [27] IASP. Task Force on Taxonomy. IASP Terminology [25/09/2020]. Available from:  
5 <https://www.iasp-pain.org/Education/Content.aspx?ItemNumber=1698>.
- 6 [28] Izumi M, Petersen KK, Laursen MB, Arendt-Nielsen L, Graven-Nielsen T. Facilitated  
7 temporal summation of pain correlates with clinical pain intensity after hip arthroplasty.  
8 *Pain* 2017;158:323-32.
- 9 [29] Johnston V, Jimmieson NL, Jull G, Souvlis T. Quantitative sensory measures distinguish  
10 office workers with varying levels of neck pain and disability. *Pain* 2008;137:257-65.
- 11 [30] Kennedy DL, Kemp HI, Ridout D, Yarnitsky D, Rice AS. Reliability of conditioned pain  
12 modulation: a systematic review. *Pain* 2016;157:2410-9.
- 13 [31] Kroenke K, Spitzer RL, Williams JB. The PHQ-9: validity of a brief depression severity  
14 measure. *J Gen Intern Med* 2001;16:606-13.
- 15 [32] Kroenke K, Spitzer RL, Williams JB, Monahan PO, Löwe B. Anxiety disorders in  
16 primary care: prevalence, impairment, comorbidity, and detection. *Ann Intern Med*  
17 2007;146:317-25.
- 18 [33] Kroenke K, Strine TW, Spitzer RL, Williams JB, Berry JT, Mokdad AH. The PHQ-8 as  
19 a measure of current depression in the general population. *J Affect Disord* 2009;114:163-  
20 73.
- 21 [34] La Touche R, Paris-Alemany A, Hidalgo-Pérez A, López-de-Uralde-Villanueva I,  
22 Angulo-Díaz-Parreño S, Muñoz-García D. Evidence for central sensitization in patients  
23 with temporomandibular disorders: a systematic review and meta-analysis of  
24 observational studies. *Pain Pract* 2018;18:388-409.



- 1 [35] Lannersten L, Kosek E. Dysfunction of endogenous pain inhibition during exercise with  
2 painful muscles in patients with shoulder myalgia and fibromyalgia. *Pain* 2010;151:77-  
3 86.
- 4 [36] Lima LV, Abner TSS, Sluka KA. Does exercise increase or decrease pain? Central  
5 mechanisms underlying these two phenomena. *J Physiol* 2017;595:4141-50.
- 6 [37] Linari-Melfi M, Cantarero-Villanueva I, Fernández-Lao C, Fernández-de-Las-Peñas C,  
7 Guisado-Barrilao R, Arroyo-Morales M. Analysis of deep tissue hypersensitivity to  
8 pressure pain in professional pianists with insidious mechanical neck pain. *BMC*  
9 *Musculoskelet Disord* 2011;12:268.
- 10 [38] Lluch Girbés E, Dueñas L, Barbero M, Falla D, Baert IA, Meeus M, Sánchez-Frutos J,  
11 Aguilera L, Nijs J. Expanded distribution of pain as a sign of central sensitization in  
12 individuals with symptomatic knee osteoarthritis. *Phys Ther* 2016;96:1196-207.
- 13 [39] MacDermid JC, Walton DM, Avery S, Blanchard A, Etruw E, McAlpine C, Goldsmith  
14 CH. Measurement properties of the neck disability index: a systematic review. *J Orthop*  
15 *Sports Phys Ther* 2009;39:400-17.
- 16 [40] Maclachlan LR, Collins NJ, Hodges PW, Vicenzino B. Psychological and pain profiles  
17 in persons with patellofemoral pain as the primary symptom. *Eur J Pain* 2020;24:1182-  
18 96.
- 19 [41] MacQueen J, editor Some methods for classification and analysis of multivariate  
20 observations. *Proceedings of the Fifth Berkeley Symposium on Mathematical Statistics*  
21 *and Probability, Volume 1: Statistics*; 281-97; Berkeley, Calif.: University of California  
22 Press; 1967.
- 23 [42] Madeleine P, Lundager B, Voigt M, Arendt-Nielsen L. Sensory manifestations in  
24 experimental and work-related chronic neck-shoulder pain. *Eur J Pain* 1998;2:251-60.

- 1 [43] Malmström EM, Stjerna J, Högestätt ED, Westergren H. Quantitative sensory testing of  
2 temperature thresholds: possible biomarkers for persistent pain? *J Rehabil Med*  
3 2016;48:43-7.
- 4 [44] Marcuzzi A, Wrigley PJ, Dean CM, Adams R, Hush JM. The long-term reliability of  
5 static and dynamic quantitative sensory testing in healthy individuals. *Pain*  
6 2017;158:1217-23.
- 7 [45] Mason KJ, O'Neill TW, Lunt M, Jones AKP, McBeth J. Psychosocial factors partially  
8 mediate the relationship between mechanical hyperalgesia and self-reported pain. *Scand*  
9 *J Pain* 2018;18:59-69.
- 10 [46] Masson B, Robinson C, Brinsmead S, Hassall L, Chamberlin S. The 2014 ASA  
11 workplace health and safety survey results. In: Association AS, editor. 2014.
- 12 [47] McKernan LC, Finn MTM, Carr ER. Personality and affect when the central nervous  
13 system is sensitized: an analysis of central sensitization syndromes in a substance use  
14 disorder population. *Psychodyn Psychiatry* 2017;45:385-409.
- 15 [48] Murray CJ, Barber RM, Foreman KJ, Abbasoglu Ozgoren A, Abd-Allah F, Abera SF,  
16 Aboyans V, Abraham JP, Abubakar I, Abu-Raddad LJ, Abu-Rmeileh NM, Achoki T,  
17 Ackerman IN, Ademi Z, Adou AK, Adsuar JC, Afshin A, Agardh EE, Alam SS, Alasfoor  
18 D, Albittar MI, Alegretti MA, Alemu ZA, Alfonso-Cristancho R, Alhabib S, Ali R, Alla  
19 F, Allebeck P, Almazroa MA, Alsharif U, Alvarez E, Alvis-Guzman N, Amare AT,  
20 Ameh EA, Amini H, Ammar W, Anderson HR, Anderson BO, Antonio CA, Anwari P,  
21 Arnlöv J, Arsic Arsenijevic VS, Artaman A, Asghar RJ, Assadi R, Atkins LS, Avila MA,  
22 Awuah B, Bachman VF, Badawi A, Bahit MC, Balakrishnan K, Banerjee A, Barker-  
23 Collo SL, Barquera S, Barregard L, Barrero LH, Basu A, Basu S, Basulaiman MO,  
24 Beardsley J, Bedi N, Beghi E, Bekele T, Bell ML, Benjet C, Bennett DA, Bensenor IM,  
25 Benzian H, Bernabé E, Bertozzi-Villa A, Beyene TJ, Bhala N, Bhalla A, Bhutta ZA,

1 Bienhoff K, Bikbov B, Biryukov S, Blore JD, Blosser CD, Blyth FM, Bohensky MA,  
2 Bolliger IW, Bora Başara B, Bornstein NM, Bose D, Boufous S, Bourne RR, Boyers LN,  
3 Brainin M, Brayne CE, Brazinova A, Breitborde NJ, Brenner H, Briggs AD, Brooks PM,  
4 Brown JC, Brugha TS, Buchbinder R, Buckle GC, Budke CM, Bulchis A, Bulloch AG,  
5 Campos-Nonato IR, Carabin H, Carapetis JR, Cárdenas R, Carpenter DO, Caso V,  
6 Castañeda-Orjuela CA, Castro RE, Catalá-López F, Cavalleri F, Çavlin A, Chadha VK,  
7 Chang JC, Charlson FJ, Chen H, Chen W, Chiang PP, Chimed-Ochir O, Chowdhury R,  
8 Christensen H, Christophi CA, Cirillo M, Coates MM, Coffeng LE, Coggeshall MS,  
9 Colistro V, Colquhoun SM, Cooke GS, Cooper C, Cooper LT, Coppola LM, Cortinovis  
10 M, Criqui MH, Crump JA, Cuevas-Nasu L, Danawi H, Dandona L, Dandona R,  
11 Dansereau E, Dargan PI, Davey G, Davis A, Davitoiu DV, Dayama A, De Leo D,  
12 Degenhardt L, Del Pozo-Cruz B, Dellavalle RP, Deribe K, Derrett S, Des Jarlais DC,  
13 Dessalegn M, Dharmaratne SD, Dherani MK, Diaz-Torné C, Dicker D, Ding EL, Dokova  
14 K, Dorsey ER, Driscoll TR, Duan L, Duber HC, Ebel BE, Edmond KM, Elshrek YM,  
15 Endres M, Ermakov SP, Erskine HE, Eshrati B, Esteghamati A, Estep K, Faraon EJ,  
16 Farzadfar F, Fay DF, Feigin VL, Felson DT, Fereshtehnejad SM, Fernandes JG, Ferrari  
17 AJ, Fitzmaurice C, Flaxman AD, Fleming TD, Foigt N, Forouzanfar MH, Fowkes FG,  
18 Paleo UF, Franklin RC, Fürst T, Gabbe B, Gaffikin L, Gankpé FG, Geleijnse JM, Gessner  
19 BD, Gething P, Gibney KB, Giroud M, Giussani G, Gomez Dantes H, Gona P, González-  
20 Medina D, Gosselin RA, Gotay CC, Goto A, Gouda HN, Graetz N, Gugnani HC, Gupta  
21 R, Gupta R, Gutiérrez RA, Haagsma J, Hafezi-Nejad N, Hagan H, Halasa YA, Hamadeh  
22 RR, Hamavid H, Hammami M, Hancock J, Hankey GJ, Hansen GM, Hao Y, Harb HL,  
23 Haro JM, Havmoeller R, Hay SI, Hay RJ, Heredia-Pi IB, Heuton KR, Heydarpour P,  
24 Higashi H, Hajar M, Hoek HW, Hoffman HJ, Hosgood HD, Hossain M, Hotez PJ, Hoy  
25 DG, Hsairi M, Hu G, Huang C, Huang JJ, Husseini A, Huynh C, Iannarone ML, Iburg

1 KM, Innos K, Inoue M, Islami F, Jacobsen KH, Jarvis DL, Jassal SK, Jee SH, Jeemon P,  
2 Jensen PN, Jha V, Jiang G, Jiang Y, Jonas JB, Juel K, Kan H, Karch A, Karema CK,  
3 Karimkhani C, Karthikeyan G, Kassebaum NJ, Kaul A, Kawakami N, Kazanjan K, Kemp  
4 AH, Kengne AP, Keren A, Khader YS, Khalifa SE, Khan EA, Khan G, Khang YH,  
5 Kielling C, Kim D, Kim S, Kim Y, Kinfu Y, Kinge JM, Kivipelto M, Knibbs LD, Knudsen  
6 AK, Kokubo Y, Kosen S, Krishnaswami S, Kuate Defo B, Kucuk Bicer B, Kuipers EJ,  
7 Kulkarni C, Kulkarni VS, Kumar GA, Kyu HH, Lai T, Lalloo R, Lallukka T, Lam H,  
8 Lan Q, Lansingh VC, Larsson A, Lawrynowicz AE, Leasher JL, Leigh J, Leung R, Levitz  
9 CE, Li B, Li Y, Li Y, Lim SS, Lind M, Lipshultz SE, Liu S, Liu Y, Lloyd BK, Lofgren  
10 KT, Logroscino G, Looker KJ, Lortet-Tieulent J, Lotufo PA, Lozano R, Lucas RM,  
11 Lunevicius R, Lyons RA, Ma S, Macintyre MF, Mackay MT, Majdan M, Malekzadeh R,  
12 Marcenes W, Margolis DJ, Margono C, Marzan MB, Masci JR, Mashal MT,  
13 Matzopoulos R, Mayosi BM, Mazorodze TT, McGill NW, McGrath JJ, McKee M,  
14 McLain A, Meaney PA, Medina C, Mehndiratta MM, Mekonnen W, Melaku YA,  
15 Meltzer M, Memish ZA, Mensah GA, Meretoja A, Mhimbira FA, Micha R, Miller TR,  
16 Mills EJ, Mitchell PB, Mock CN, Mohamed Ibrahim N, Mohammad KA, Mokdad AH,  
17 Mola GL, Monasta L, Montañez Hernandez JC, Montico M, Montine TJ, Mooney MD,  
18 Moore AR, Moradi-Lakeh M, Moran AE, Mori R, Moschandreas J, Moturi WN, Moyer  
19 ML, Mozaffarian D, Msemburi WT, Mueller UO, Mukaigawara M, Mullany EC,  
20 Murdoch ME, Murray J, Murthy KS, Naghavi M, Naheed A, Naidoo KS, Naldi L, Nand  
21 D, Nangia V, Narayan KM, Nejjari C, Neupane SP, Newton CR, Ng M, Ngalesoni FN,  
22 Nguyen G, Nisar MI, Nolte S, Norheim OF, Norman RE, Norrving B, Nyakarahuka L,  
23 Oh IH, Ohkubo T, Ohno SL, Olusanya BO, Opio JN, Ortblad K, Ortiz A, Pain AW,  
24 Pandian JD, Panelo CI, Papachristou C, Park EK, Park JH, Patten SB, Patton GC, Paul  
25 VK, Pavlin BI, Pearce N, Pereira DM, Perez-Padilla R, Perez-Ruiz F, Perico N, Pervaiz

1 A, Pesudovs K, Peterson CB, Petzold M, Phillips MR, Phillips BK, Phillips DE, Piel FB,  
2 Plass D, Poenaru D, Polinder S, Pope D, Popova S, Poulton RG, Pourmalek F,  
3 Prabhakaran D, Prasad NM, Pullan RL, Qato DM, Quistberg DA, Rafay A, Rahimi K,  
4 Rahman SU, Raju M, Rana SM, Razavi H, Reddy KS, Refaat A, Remuzzi G, Resnikoff  
5 S, Ribeiro AL, Richardson L, Richardus JH, Roberts DA, Rojas-Rueda D, Ronfani L,  
6 Roth GA, Rothenbacher D, Rothstein DH, Rowley JT, Roy N, Ruhago GM, Saeedi MY,  
7 Saha S, Sahraian MA, Sampson UK, Sanabria JR, Sandar L, Santos IS, Satpathy M,  
8 Sawhney M, Scarborough P, Schneider IJ, Schöttker B, Schumacher AE, Schwebel DC,  
9 Scott JG, Seedat S, Sepanlou SG, Serina PT, Servan-Mori EE, Shackelford KA, Shaheen  
10 A, Shahraz S, Shamah Levy T, Shangguan S, She J, Sheikhabaei S, Shi P, Shibuya K,  
11 Shinohara Y, Shiri R, Shishani K, Shiue I, Shrimel MG, Sigfusdottir ID, Silberberg DH,  
12 Simard EP, Sindi S, Singh A, Singh JA, Singh L, Skirbekk V, Slepak EL, Sliwa K, Soneji  
13 S, Søreide K, Soshnikov S, Sposato LA, Sreeramareddy CT, Stanaway JD, Stathopoulou  
14 V, Stein DJ, Stein MB, Steiner C, Steiner TJ, Stevens A, Stewart A, Stovner LJ,  
15 Stroumpoulis K, Sunguya BF, Swaminathan S, Swaroop M, Sykes BL, Tabb KM,  
16 Takahashi K, Tandon N, Tanne D, Tanner M, Tavakkoli M, Taylor HR, Te Ao BJ,  
17 Tediosi F, Temesgen AM, Templin T, Ten Have M, Tenkorang EY, Terkawi AS,  
18 Thomson B, Thorne-Lyman AL, Thrift AG, Thurston GD, Tillmann T, Tonelli M,  
19 Topouzis F, Toyoshima H, Traebert J, Tran BX, Trillini M, Truelsen T, Tsilimbaris M,  
20 Tuzcu EM, Uchendu US, Ukwaja KN, Undurraga EA, Uzun SB, Van Brakel WH, Van  
21 De Vijver S, van Gool CH, Van Os J, Vasankari TJ, Venketasubramanian N, Violante  
22 FS, Vlassov VV, Vollset SE, Wagner GR, Wagner J, Waller SG, Wan X, Wang H, Wang  
23 J, Wang L, Warouw TS, Weichenthal S, Weiderpass E, Weintraub RG, Wenzhi W,  
24 Werdecker A, Westerman R, Whiteford HA, Wilkinson JD, Williams TN, Wolfe CD,  
25 Wolock TM, Woolf AD, Wulf S, Wurtz B, Xu G, Yan LL, Yano Y, Ye P, Yentür GK,

- 1 Yip P, Yonemoto N, Yoon SJ, Younis MZ, Yu C, Zaki ME, Zhao Y, Zheng Y, Zonies  
2 D, Zou X, Salomon JA, Lopez AD, Vos T. Global, regional, and national disability-  
3 adjusted life years (DALYs) for 306 diseases and injuries and healthy life expectancy  
4 (HALE) for 188 countries, 1990-2013: quantifying the epidemiological transition. *Lancet*  
5 2015;386:2145-91.
- 6 [49] Niri R-R, Granovskiy Y, Yarnitskiy D, Sprecherl E, Granotl M. A psychophysical study  
7 of endogenous analgesia: The role of the conditioning pain in the induction and  
8 magnitude of conditioned pain modulation. *Eur J Pain* 2011;15:491-7.
- 9 [50] Petersen KK, Arendt-Nielsen L, Simonsen O, Wilder-Smith O, Laursen MB. Presurgical  
10 assessment of temporal summation of pain predicts the development of chronic  
11 postoperative pain 12 months after total knee replacement. *Pain* 2015;156:55-61.
- 12 [51] Rolke R, Baron R, Maier C, Tölle TR, Treede RD, Beyer A, Binder A, Birbaumer N,  
13 Birklein F, Bötefür IC, Braune S, Flor H, Hüge V, Klug R, Landwehrmeyer GB, Magerl  
14 W, Maihöfner C, Rolko C, Schaub C, Scherens A, Sprenger T, Valet M, Wasserka B.  
15 Quantitative sensory testing in the German Research Network on Neuropathic Pain  
16 (DFNS): standardized protocol and reference values. *Pain* 2006;123:231-43.
- 17 [52] Scott D, Jull G, Sterling M. Widespread sensory hypersensitivity is a feature of chronic  
18 whiplash-associated disorder but not chronic idiopathic neck pain. *Clin J Pain*  
19 2005;21:175-81.
- 20 [53] Shahidi B, Maluf KS. Adaptations in evoked pain sensitivity and conditioned pain  
21 modulation after development of chronic neck pain. *Biomed Res Int* 2017;2017:8985398.
- 22 [54] Smart KM, Blake C, Staines A, Doody C. Self-reported pain severity, quality of life,  
23 disability, anxiety and depression in patients classified with 'nociceptive', 'peripheral  
24 neuropathic' and 'central sensitisation' pain. The discriminant validity of mechanisms-  
25 based classifications of low back ( $\pm$ leg) pain. *Man Ther* 2012;17:119-25.

- 1 [55] Smart KM, O'Connell NE, Doody C. Towards a mechanisms-based classification of pain  
2 in musculoskeletal physiotherapy? *Phys Ther Rev* 2008;13:1-10.
- 3 [56] Smith A, Ritchie C, Warren J, Sterling M. Exercise-induced hypoalgesia is impaired in  
4 chronic whiplash-associated disorders (WAD) with both aerobic and isometric exercise.  
5 *Clin J Pain* 2020;36:601-11.
- 6 [57] Steinmetz A, Jull GA. Sensory and sensorimotor features in violinists and violists  
7 with neck pain. *Arch Phys Med Rehabil* 2013;94:2523-8.
- 8 [58] Sullivan MJL, Bishop SR, Pivik J. The Pain Catastrophizing Scale: development and  
9 validation. *Psychol Assess* 1995;7:524-32.
- 10 [59] Tabachnick BG, Fidell LS. *Using multivariate statistics*, 5th ed. Boston, MA: Allyn &  
11 Bacon/Pearson Education; 2007. pp. 78.
- 12 [60] Vardeh D, Mannion RJ, Woolf CJ. Toward a mechanism-based approach to pain  
13 diagnosis. *J Pain* 2016;17:T50-69.
- 14 [61] Vernon H, Mior S. The Neck Disability Index: a study of reliability and validity. *J*  
15 *Manipulative Physiol Ther* 1991;14:409-15.
- 16 [62] Village J, Trask C. Ergonomic analysis of postural and muscular loads to diagnostic  
17 sonographers. *Int J Ind Ergon* 2007;37:781-9.
- 18 [63] Waddell G, Newton M, Henderson I, Somerville D, Main CJ. A Fear-Avoidance Beliefs  
19 Questionnaire (FABQ) and the role of fear-avoidance beliefs in chronic low back pain  
20 and disability. *Pain* 1993;52.
- 21 [64] Walton DM, Carroll LJ, Kasch H, Sterling M, Verhagen AP, Macdermid JC, Gross A,  
22 Santaguida PL, Carlesso L. An overview of systematic reviews on prognostic factors in  
23 neck pain: results from the International Collaboration on Neck Pain (ICON) Project.  
24 *Open Orthop J* 2013;7:494-505.

- 1 [65] Walton DM, Macdermid JC, Nielson W, Teasell RW, Chiasson M, Brown L. Reliability,  
2 standard error, and minimum detectable change of clinical pressure pain threshold testing  
3 in people with and without acute neck pain. *J Orthop Sports Phys Ther* 2011;41:644-50.
- 4 [66] Wells R. Why have we not solved the MSD problem? *Work* 2009;34:117-21.
- 5 [67] Xie Y, Coombes BK, Thomas L, Johnston V. Musculoskeletal pain and disability in  
6 sonographers: more than an ergonomic issue. *J Am Soc Echocardiogr* 2020.
- 7 [68] Xie Y, Jun D, Thomas L, Coombes BK, Johnston V. Comparing central pain processing  
8 in individuals with non-traumatic neck pain and healthy individuals: a systematic review  
9 and meta-analysis. *J Pain* 2020.
- 10 [69] Yarnitsky D, Arendt-Nielsen L, Bouhassira D, Edwards RR, Fillingim RB, Granot M,  
11 Hansson P, Lautenbacher S, Marchand S, Wilder-Smith O. Recommendations on  
12 terminology and practice of psychophysical DNIC testing. *Eur J Pain* 2010;14:339.
- 13 [70] Zebis MK, Andersen LL, Pedersen MT, Mortensen P, Andersen CH, Pedersen MM,  
14 Boysen M, Roessler KK, Hannerz H, Mortensen OS, Sjøgaard G. Implementation of  
15 neck/shoulder exercises for pain relief among industrial workers: a randomized  
16 controlled trial. *BMC Musculoskelet Disord* 2011;12:205.
- 17 [71] Zusman M. Forebrain-mediated sensitization of central pain pathways: 'non-specific'  
18 pain and a new image for MT. *Man Ther* 2002;7:80-8.

## 19 **Figure legends**

20 **Figure 1.** Study participant flow. NDI, neck disability index.

21 **Figure 2.** Pain frequency maps generated separately for no, mild and moderate/severe  
22 disability groups by superimposing the pain drawings of all participants within the group. The  
23 color grid indicates both the number and the percentage of participants who reported pain in



1 that specific area. Female and male participants shaded their pain areas on a female and male  
2 body chart, respectively. For illustrative purposes, all data is presented on a female body chart.

3 **Figure 3.** Mean intensity of worst pain experienced in different body regions in the previous  
4 week for the no, mild and moderate/severe disability groups. Error bars depict standard  
5 deviations. \* $P < 0.05$ ; \*\* $P < 0.001$ .

6 **Figure 4.** Mean cold (A), heat (B) and pressure (C) pain thresholds at local (neck, upper  
7 trapezius and deltoid) and remote (tibialis anterior) sites for the no, mild, and moderate/severe  
8 disability groups. A higher score indicates worse (hyperalgesia) for cold pain threshold while  
9 a lower score is worse for heat and pressure pain thresholds. Error bars depict standard  
10 deviations. \* $P < 0.05$ ; \*\* $P < 0.001$

11 **Figure 5.** Individual data plots for temporal summation of pain (A) and conditioned pain  
12 modulation (B) for the no, mild, and moderate/severe disability groups. (A) Changes in pain  
13 rating from a single to 10 repeated pinprick stimuli, with higher values representing greater  
14 temporal summation of pain. (B) Changes in pressure pain threshold from before to during the  
15 conditioning stimulus, with positive changes reflecting endogenous pain inhibition and higher  
16 values representing greater pain inhibition. Black diamonds and lines depict group means and  
17 standard deviations. \* $P < 0.05$

18 **Figure 6.** Mean changes of pressure pain threshold (PPT) at local (neck, upper trapezius,  
19 deltoid) and remote (tibialis anterior) sites in response to an isometric exercise for the no, mild,  
20 and moderate/severe disability groups. A positive change reflects exercise-induced analgesia  
21 and higher values represent greater efficacy of analgesia. Error bars depict standard deviations.

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