Upper cervical two-point discrimination thresholds in migraine patients and headache-free controls

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Abstract

Background: Chronic pain including migraine is associated with structural and functional changes in the somatosensory cortex. Previous reports proposed two-point discrimination (TPD) as a measurement for cortical alterations. Limited evidence exists for tactile acuity in the neck and no data is available for migraine.

Methods: To introduce a standardized protocol for the measurement of TPD in the upper cervical spine, 51 healthy participants were investigated with a newly developed paradigm which was evaluated for intra-rater reliability. The same protocol was applied by two further examiners to 28 migraine patients and 21 age-, and gender-matched healthy controls to investigate inter-rater reliability and between group differences.

Results: Results indicated excellent intra-rater (right ICC(2,4) = 0.82, left ICC(2,4) = 0.83) and good inter-rater reliability (right ICC(2,4) = 0.70, left ICC(2,4) = 0.75). Migraine patients had larger TPD thresholds (26.86 ± 7.21) than healthy controls (23.30 ± 6.17) but these became only statistically significant for the right side of the neck (p = 0.02). There was a significant, moderate association with age for the right side (r = 0.42, p = 0.002, n = 51), and less strong association for the left side (r = 0.34, p = 0.14) in healthy individuals. TPD did not correlate with headache days per month or the dominant headache side in migraine patients.

Conclusions: Surprisingly, migraine patients showed increased TPD thresholds in the upper cervical spine interictally. Although a body of evidence supports that hypersensitivity is part of the migraine attack, the current report indicates that interictally, migraine patients showed worse tactile acuity similar to other chronic pain populations. This has been hypothesized to indicate structural and functional re-organisation of the somatosensory cortex.

Keywords: Tactile acuity, Hypersensitivity, Migraine, Headache, Two-point discrimination

Background

Current knowledge on functional and structural cortical alterations following longstanding pain, is mainly based on chronic low back pain populations [1–6] with very limited data for the upper spinal region [7]. However, it has been suggested, that at least structural brain changes are similar for any chronic pain condition including headaches [8] and that the somatosensory cortex is one of the structures typically affected [5, 9, 10].

Two-point discrimination threshold (TPD) assessment is a routine procedure during the neurological examination that has been shown to be associated with such structural brain changes [9, 10]. TPD thresholds indicate the smallest distance between two points of sliding mechanical callipers, that can still be perceived as two distinct points. This paradigm has been used in research to investigate tactile acuity in chronic pain patients and a recent systematic review reported a general tendency to increased TPD thresholds in patients with longstanding pain compared to pain-free controls [11, 12].
Six studies have previously investigated TPD in the neck: Song et al. investigated patients with spinal cord injuries and found multiple sites of reduced sensory discrimination compared to healthy controls that were reported to be particularly distinct in the neck area [13]. Moreira et al. found increased TPD thresholds in patients with idiopathic neck pain but although there was a mean difference between groups of almost 5 mm, values failed to reach statistical significance [14]. Nolan et al. investigated TPD only in healthy participants; the mean TPD at the level of C7 was 55.4 mm [15]. All three studies applied their measurements to the lower cervical area. The only study that included a measurement in the upper cervical spine in patients with mild recurrent neck pain, while observing slightly larger TPD values in the patient group, failed to show any statistically significant difference between patients and controls at C2 and at C7 [16]. The mean two-point discrimination threshold of the included 30 neck pain patients was 29.75 mm at C2 and 32.5 mm at C7. Catley et al. and Harvie et al. conducted reliability studies in healthy participants. The authors examined TPD threshold only at the C7 spinal level and reported a mean of 45.9 mm and of 35 mm, respectively [17, 18].

All six studies showed that tactile acuity can feasibly be measured in the cervical area, two studies confirmed the intra-rater and inter-rater reliability (ICC > 0.75) of the measurement [17, 18]. All studies evaluating patients showed that the patient population had larger TPD values compared to the control group. No study has previously applied TPD to a headache population. However, the test procedures used in the six studies were heterogeneous and reported a wide range of mean threshold in a variety of populations. Only one study investigated the upper cervical region and no data were available for migraine patients.

The aim of the current project was therefore to investigate TPD thresholds in the upper cervical spine in patients with migraine in comparison to healthy controls. Since only one study has investigated this region previously, this current project includes the development and both, intra- and inter-reliability testing of the procedure.

Methods
Study overview
The study was designed in two stages: the initial stage was the development of the test protocol and determination of intra-rater reliability using a test-retest design. The second stage for the inter-rater reliability and the difference between migraine patients and controls was a repeated-measures case-control design with blinded examiners (Fig. 1).

The study was approved by the local ethics committee of the University of Hamburg (AZ 2017_86). It was registered a priori in the German clinical trials register (DRKS00011795). The data collection for the intra-rater stage was conducted between November and December 2016 and for the inter-rater and case control stage between May and June 2017.

Participants
Healthy control participants for the intra-rater reliability phase as well as those serving as controls for the case-control study had to be adults (18+ years old), age and gender matched to the patient group (only for the case-control phase), and right-handed. Patients for the case-control phase were: adults, diagnosed with migraine according to IHS 3 criteria [19], had a minimum of 4 migraine days on average per month, took pain medication on < 10 days per month to exclude medication overuse headache, had no other relevant headache diagnosis (episodic tension-type headache was accepted), had to be headache-free 48 h prior to the assessment. Patients and controls were excluded when they were older than 65 years; to reduce bias and to differentiate between the influence of headache and any other pain condition, they were also excluded when they had acute pain including toothache on the day of testing, had any chronic pain condition, had neck pain in the past 3 months, had a history of cervical pathology or trauma including whiplash associated disorder, disc disease or others requiring medical intervention, suffered from any psychological, neurological or other disease potentially influencing the sensory system, or a skin condition in the area over the upper cervical spine.

Patients were recruited from the University Headache Clinic at the University Medical Center Hamburg-Eppendorf and from local neurology clinics. Control participants were selected from social media and digital network postings and personal contacts.

Measurement procedure
An initial test protocol was developed and piloted in a small convenience sample of healthy participants before it was subsequently applied to the study population. It was based on the most widely used procedure for the lumbar spine of Moberg [20] who introduced a protocol with increasing and decreasing distances between two tips of a mechanical sliding calliper [20] similar to the method of limits used in the quantitative sensory testing protocol [21]. Previous publications suggested that the discrimination threshold in the neck will be between 10 and 45 mm [11, 15, 18] and it was reported as 29.75 mm at C2 in patients with neck pain [16]. Therefore, the TPD test started at a distance of 50 mm, a distance that was perceived by all participants as two distinct points. The distance was subsequently decreased by 5 mm until only 1 point was perceived and increased in steps of
1 mm until two points were felt. This was followed by decreasing the distance in 1 mm steps until only one point was perceived. This was repeated until 4 values were documented, two from increasing and two from decreasing the distance. The mean of these 4 data points was the TPD threshold for one measurement. For the intra-rater evaluation, the procedure was repeated by the same examiner after a five-minute break. All participants were tested on both sides of the neck in a randomized order.

Participants were positioned on a stool in front of a table with the forehead resting on a folded towel. The third vertebra was located manually and marked with a felt tip pen. The medial tip of the callipers was always within a radius of 2 cm from the midpoint of the 3rd vertebrae. The TPD was always measured horizontally at the level of C3. To standardize the pressure, calliper tips were placed on the test location until the first blanching of the skin. This standardization has been widely applied in previous studies assessing tactile acuity [17, 22].

For the inter-rater reliability and case-control stage, patients and controls were randomly allocated to a first blinded examiner who performed the TPD testing as described above (two measurements on each side of the neck, 1 measurement at the hand). After a resting period of 10 min they changed to the second blinded examiner. During this stage of the study, the lateral border of the dorsal hand was additionally tested as a control region in both groups, healthy controls and migraines. This allowed an indication whether tactile acuity changes are limited to the painful region or can also be identified in remote regions, thereby potentially indicating central sensitization.

Blinding of the examiners was ensured by a third researcher, who contacted potential participants, performed the randomization to the first blinded examiner, distributed the questionnaires and instructed participants to not reveal their status as a patient or a control person during the test procedure. Both examiners were experienced in the physical examination of patients and trained during repeated sessions to perform the test in a standardized manner. In addition to the TPD thresholds, the following parameters were recorded during the test days by a piloted questionnaire: gender, age, number of headache days per month, years since diagnosis and dominant headache side.
Proposed sample size and statistical analysis

The required sample size for the case-control phase was based on data for patients with chronic low back pain [13, 22] pain using 80% power and an alpha value of 0.05. The minimal required sample size per group was minimum of 19, the calculation was conducted in G*Power [23].

Reliability was calculated using the two-way random average measures intraclass-correlation coefficient for absolute agreement (ICC\(_{2,4}\)). Reliability was interpreted as excellent if the ICC was > 0.75 [24] and good if the ICC was between 0.74–0.50. The variance components (\(\sigma^2_p\) patient, \(\sigma^2_o\) observer and \(\sigma^2_r\) residual) were calculated with VARCOMP-Analysis (Method: ANOVA Type III Sum of Squares) to determine Standard Error of Measurement (SEM) and the smallest detectable difference (SDD) [25]. Differences across groups were calculated using two-sided t-tests for independent samples for each side of the neck and for the reference area (hand) individually. For these calculations, the mean ratings across the two examiners were used.

Since previous research has indicated that age and gender may play a role in the TPD measurement [26, 27], secondary analyses evaluated whether TPD thresholds were significantly different between male and female patients and in younger and older patients. For the analysis of age, a median split was used and significance was tested using t-tests at an alpha level of 0.05. Additionally, Pearson’s correlation coefficient was calculated to assess association between age and TPD threshold. Because TPD in the neck region has recently been shown to be associated with the severity of a disease [28], TPD thresholds were correlated in the headache group with headache days per month and with the dominant headache side using Pearson’s correlation. All analyses were calculated using the software packages SPSS 23 (IBM, Illinois, USA) and Stata 15 (StataCorp LLC, College Station, Texas, USA).

Results

Intra-rater reliability

Fifty-one healthy participants (26 female) aged between 20 and 65 years (mean 37.8 ± 13 years) were recruited. TPD thresholds ranged between 7.5 and 41 mm (mean 26.7 ± 7.1 mm). Values for male and female as well as older and younger participants are presented in Table 1. The intra-rater reliability was excellent with an ICC\(_{2,4}\) of 0.83 (95% CI, 0.72–0.90) for the left side of the neck and an ICC\(_{2,4}\) of 0.82 (95% CI, 0.70–0.89) for the right side of the neck. The SEM was 3.1 mm for both sides with corresponding SDDs of 8.5 (left side) and 8.6 (right side). There was a significant, moderate correlation between age and TPD (left side \(r = 0.342\); right side \(r = 0.421\)).

Inter-rater reliability and case-control

For the inter-rater and case-control phase of the study, 49 participants (28 migraine patients) were recruited. Characteristics for patients and control participants are reported in Table 2. The inter-rater reliability for all measurements on the left (ICC\(_{2,4}\) 0.75; 95% CI 0.55–0.86; SDD 12.02; SEM 4.34) and the right side (ICC\(_{2,4}\) 0.70; 95% CI 0.46–0.83; SDD 12.36; SEM 4.46) of the neck and for the measurements on the hand (ICC\(_{2,4}\) 0.70; 95% CI 0.48–0.83; SDD 6.01; SEM 2.1) was good. Migraine patients had larger TPD thresholds at all test locations including the hand. Statistical significance was only reached for the right side of the neck (\(p = 0.02\)) (Table 3). There was no correlation with headache days per month for left (\(r = -0.09\)), or right side (\(r = -0.11\)) of the neck, and the hand (\(r = -0.15\)). Neither was a correlation with the dominant headache side revealed for the left (\(r = -0.19\)) or the right side (\(r = -0.20\)) of the neck.

Discussion

This paper aimed to investigate tactile acuity in migraine patients compared to headache-free controls and to develop a standardised test protocol with sufficient intra- and inter-rater reliability. The mean TPD thresholds in healthy participants in this study are in line with those in the only other study on the upper cervical spine [16]. Intra- and inter-rater reliability coefficients were similarly high as those reported by Harvie et al. [28] and Catley et al. [17] for the lower part of the neck, thereby supporting TPD testing as a clinically reliable tool. Also similar to the results reported by Harvie et al. [28], who

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Table 1  TPD thresholds for male and female as well as younger and older participants (intra-rater reliability sample)

| Between group differences | Male (n = 25) | Female (n = 26) | \(r\) - test |
|---------------------------|--------------|----------------|-------------|
| Left neck (mean ± SD)     | 2.75 (0.71) cm | 2.6 (0.72) cm | \(p = 0.458\) |
| Right neck (mean ± SD)    | 2.9 (0.73) cm | 2.5 (0.53) cm | \(p = 0.031\) |

| Between group differences | Age 36 ≤ (n = 26) | Age > 35 (n = 25) | \(r\) - test |
|---------------------------|------------------|------------------|-------------|
| Left neck (mean ± SD)     | 2.42 (0.64) cm   | 2.93 (0.7) cm    | \(p = 0.008\) |
| Right neck (mean ± SD)    | 2.40 (0.54) cm   | 3.0 (0.65) cm    | \(p = 0.001\) |

SD Standard deviations
assessed patients with chronic neck pain, migraine patients scored worse, i.e. had larger TPD thresholds than control participants, although the results in this current study were only statistically significant for the right side of the neck. TPD thresholds for the left side of the neck were also larger than in healthy controls but not statistically significant. Furthermore, results did not overcome the SDD and were very close to the SEM and might therefore be in line with Elsig et al. who found no difference between patients with recurrent neck pain and healthy controls [16]. The consistency of the direction of change for all tested body regions and the overwhelming majority of publications in this field showing larger TPD thresholds in pain patients compared to healthy populations, increase the confidence in the current results. Interestingly, although not statistically significant was, that TPD thresholds were also increased when measured outside the area of clinical pain at the right hand, pointing towards central rather than peripheral or region-specific changes. This result is also similar to that reported by Harvie et al. [28]. Older and younger healthy participants (median split at 35 years) showed a statistically significant difference and younger females showed the highest tactile acuity, i.e. smallest TPD thresholds.

Although significance was only reached for the right side of the neck, the generally increased TPD thresholds in migraine patients compared to controls might support the frequently stated hypothesis and that repeated migraine attacks result in similar cortical changes as observed in patients with chronic pain conditions [11]. While not surprising in the context of chronic pain research, the results of the current study are somewhat puzzling considering the prevalence of hypersensitivity and/or alldynia in patients with migraine. Altered sensitivity to external stimuli such as light, noise, smell or touch is one of the trait symptoms and diagnostic criteria for migraine [19]. Numerous studies focused on this phenomenon and reported hypersensitivity and/or alldynia in patients with migraine [29–37]. Smaller TPD thresholds would have been the more intuitive result representing an increased tactile sense and thereby a phenomenon which could be interpreted as hypersensitivity. One explanation for this controversy could be that hypersensitivity and/or alldynia are indeed only present during the pre-ictal and ictal phase of migraine [33], while reports on interictal hypersensitivity are conflicting [38]. A further explanation could be that alldynia and/or hypersensitivity is restricted to the reference area of the trigeminal nerve. Very few studies, additionally to the typically measured supraorbital region, measured hypersensitivity in the neck [34, 39, 40]. This limited evidence is surprising, since neck pain is more common in migraine patients than other typically associated symptoms - such as nausea - and occurs in more than 70% of migraine patients [41]. The 12-item Allodynia Checklist contains items such as “wearing a neckless” which was found to be the second most discriminative item between migraine patients with and without cutaneous alldynia [36]. Based on the trigeminocervical convergence theory, stating that afferent fibres from the trigeminal and the cervical systems connect to the same nuclei in the brainstem, there is a strong anatomical and physiological explanation for a reciprocal influence of the trigeminal system on cervical reference areas and vice versa [42–44]. Furthermore, a study using a neural staining technique in rodents reported a connection between the dura mater and peripheral muscles through the cranial fissures and thereby another possible pathway explaining neck pain or and sensitivity when the dura mater is sensitized [45].

A final explanation is that tactile acuity and hypersensitivity/alldynia, while fulfilling the same biological role of an increased awareness to potentially damaging stimuli to a diseased or painful body region, are unrelated entities. The most common paradigm to quantify hypersensitivity and/or alldynia is the quantitative sensory testing (QST) protocol where different noxious and non-noxious modalities are applied to the affected body part and perception thresholds, pain thresholds and pain responses are recorded [21]. However, QST requires

### Table 2 Characteristics of groups

| Variable                        | Migraine n = 28 | Control participants n = 21 |
|---------------------------------|-----------------|-----------------------------|
| Female: number (%)             | 27 (93.1%)      | 17 (85%)                    |
| Male: number (%)               | 2 (6.9%)        | 3 (15%)                     |
| Age (years)                    | 34.4 (11.9)     | 39.8 (13.6)                 |
| Years since diagnosis          | 19.7 (14.1)     |                             |
| Headache days per month        | 9.4 (9.1)       |                             |
| Dominant headache side         |                 |                             |
| - Left number (%)              | 7 (24.1)        |                             |
| - Right number (%)             | 7 (24.1)        |                             |
| - Bilateral number (%)         | 15 (51.7)       |                             |

### Table 3 Mean and standard deviations of two-point discrimination thresholds

| Side / location of testing | Migraine patients (n = 28) | Healthy control (n = 21) | t - test |
|----------------------------|-----------------------------|--------------------------|---------|
| Left                       | 27.4 (8.2) mm               | 24.9 (7.1) mm            | p = 0.27|
| Right                      | 26.3 (7.3) mm               | 21.7 (6.2) mm            | p = 0.02|
| Hand                       | 8.3 (4.0) mm                | 6.4 (2.2) mm             | p = 0.05|
equipment not always available to clinicians and the full protocol is extremely time consuming. Whether TPD thresholds can be used as an alternative test paradigm feasible for the use in the daily clinical practice remains to be evaluated in future studies.

A limitation of this study is, that only the neck and the hand but not the reference area of the trigeminal nerve was assessed. Although this was the aim of this project, retrospectively, an additional assessment of the supraorbital area on the side most affected by migraine would have been helpful to at least partially disentangle whether the test location influences results. Furthermore, a second modality, such as thermal thresholds or von Frey hairs would have been interesting to compare the current results with previous publications and to investigate whether TPD thresholds are related to hypersensitivity and/or allodynia. To disentangle responses to tactile stimuli and thereby better understand what the test modalities actually measure would be valuable future projects.

Conclusion
Despite the limited available space in the upper cervical region, TPD thresholds can be measured reliably in healthy participants and in patients with migraine. Younger and female patients have smaller TPD thresholds than older and male patients, future research should take this into account. Patients with migraine show larger TPD thresholds in the upper cervical region, but significance was only reached for the right side of the neck. Remote body parts also showed increased values (but did not reach statistical significance), potentially pointing towards cortical changes similar to those reported for chronic pain conditions.

Abbreviations
ICC: Intraclass correlation coefficient; QTS: Quantitative sensory testing; SD: Standard deviation; SEM: Standard error of measurement; TPD: Two-point discrimination threshold

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Availability of data and materials
Data available upon individual request.

Authors’ contributions
KL planned and designed the study, supported the analysis and wrote the final manuscript. WA designed the two-point discrimination paradigm, supported the data analysis and corrected the final manuscript. KM, IM, LR, JS and CZ recruited participants, collected the data, analysed the data and prepared a first draft of the manuscript. AS supervised the intra-rater stage and analysed intra-rater data. TS supported the data analysis and helped prepare the final draft of the manuscript, BW supervised the inter-rater stage of the study and helped prepare the final manuscript. All authors read and approved the final manuscript.

Ethics approval and consent to participate
The study was approved by the local ethics committee of the University of Hamburg (AZ 2017_86).

Consent for publication
Not applicable.

Competing interests
The authors declare that they have no competing interests.

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References
1. Apkarian AV, Sosa Y, Sonty S et al (2004) Chronic back pain is associated with decreased prefrontal and thalamic gray matter density. J Neurosci 24:10410–10415. https://doi.org/10.1523/JNEUROSCI.2541-04.2004
2. Apkarian AV, Hashmi JA, Baliki MN (2010) Pain and the brain: specificity and plasticity of the brain in clinical chronic pain. Pain 152:549–564. https://doi.org/10.1016/j.pain.2010.11.010
3. Buckalew N, Haut MW, Morrow L, Weiner D (2008) Chronic pain is associated with brain volume loss in older adults: preliminary evidence. Pain Med Malden Mass 9:240–248. https://doi.org/10.1111/j.1526-4637.2008.00412.x
4. Schmidt-Wilcke T, Leinisch E, Gänßbauer S et al (2006) Affective components and intensity of pain correlate with structural differences in gray matter in chronic back pain patients. Pain 125:89–97. https://doi.org/10.1016/j.pain.2005.05.004
5. Flor H, Braun C, Elbert T, Birbaumer N (1997) Extensive reorganization of primary somatosensory cortex in chronic back pain patients. Neurosci Lett 224:5–8
6. Giesecke T, Gracey RH, Grant MAB et al (2004) Evidence of augmented central pain processing in idiopathic chronic low back pain. Arthritis Rheum 50:613–623. https://doi.org/10.1002/art.20063
7. Mao C, Wei L, Zhang Q et al (2013) Differences in brain structure in patients with distinct sites of chronic pain: a voxel-based morphometric analysis. Neural Regen Res 8:2081–2090. https://doi.org/10.3969/j.issn.1673-5374.2013.32.001
8. May A (2011) Structural brain imaging: a window into chronic pain. Neurosci Rev J Bringing Neurobiol Neurol Psychiatry 17:209–220. https://doi.org/10.1177/1073858410396220
9. Pleger B, Rager P, Schwenkres P et al (2006) Patterns of cortical reorganization parallel impaired tactile discrimination and pain intensity in complex regional pain syndrome. Neuroimage 32:503–510. https://doi.org/10.1016/j.neuroimage.2006.03.045
10. Schmidt-Wilcke T, Wulms N, Heba S et al (2018) Structural changes in brain morphology induced by brief periods of repetitive sensory stimulation. Neuroimage 165:148–157. https://doi.org/10.1016/j.neuroimage.2017.10.016
11. Catley MJ, O’Connell NE, Berrymen C et al (2014) Is tactile acuity altered in people with chronic pain? A systematic review and meta-analysis. J Pain Off J Am Pain Soc 15:985–1000. https://doi.org/10.1016/j.jpain.2014.06.009
12. Adamczyk W, Luedtke K, Sausitz E (2017) Lumbar tactile acuity in patients with low back pain and healthy controls: systematic review and meta-analysis. Clin J Pain. https://doi.org/10.1097/AJP.0000000000000499
13. Song ZK, Cohen MJ, Ament PA et al (1993) Two-point discrimination thresholds in spinal cord injured patients with dysesthetic pain. Paraplegia 31:425–493. https://doi.org/10.1034/jc.1993.79
14. Moreira C, Bassi AR, Brando MP, Silva AG (2017) Do patients with chronic neck pain have distorted body image and tactile dysfunction? Eur J Phys 19: 215–221. https://doi.org/10.1080/21679169.2017.1339418
15. Nolan MF (1985) Quantitative measure of cutaneous sensation. Two-point discrimination values for the face and trunk. Phys Ther 65:181–185
16. Elsig S, Luomajoki H, Sattelmayer M et al (2014) Sensormotor tests, such as movement control and laterality judgment accuracy, in persons with recurrent neck pain and controls. A case-control study. Man Ther 19:155–561. https://doi.org/10.1016/j.math.2014.05.014
17. Catley MJ, Tabor A, Ward BM, Mosley GL (2013) Assessing tactile acuity in rhenumatology and musculoskeletal medicine–how reliable are two-point discrimination tests at the neck, hand, back and foot? Rheumatol Oxf Engl 52:1454–1461. https://doi.org/10.1093/rheumatology/ket140
18. Harvie DS, Kelly J, Buckman H et al (2017) Tactile acuity testing at the neck: a comparison of methods. Musculoskelet Sci Pract 32:23–30. https://doi.org/10.3101/jmskp.2017.07.002
19. International Headache Society (2018) Headache classification Committee of the International Headache Society (IHS) the international classification of headache disorders, 3rd edition. Cephalalgia Int J Headache 38:1–211. https://doi.org/10.1177/0333102417773820
20. Moberg E (1990) Two-point discrimination test. A valuable part of hand surgical rehabilitation, e.g. in tetraplegia. Scand J Rehabil Med 22:127–134
21. Rolke R, Baron R, Moher C et al (2006) Quantitative sensory testing in the German research network on neuropathic pain (DFNS): standardized protocol and reference values. Pain 123:231–243. https://doi.org/10.1016/j.pain.2006.01.041
22. Adamczyk W, Tabor A, Ward BM, Mosley GL (2013) Assessing tactile acuity in rhenumatology and musculoskeletal medicine–how reliable are two-point discrimination tests at the neck, hand, back and foot? Rheumatol Oxf Engl 52:1454–1461. https://doi.org/10.1093/rheumatology/ket140
23. Rousson V (2013) Measurement in medicine, by H. C. W. de Vet, C. B. Terwee, L. B. Mokkink, and D. L. Knol. JBioPharmStat 23:277–279. https://doi.org/10.1080/21679169.2013.737220
24. Fleiss JL (2011) Design and analysis of clinical experiments. New Jersey: Wiley. https://doi.org/10.1002/9781118032923
25. Rousson V (2013) Measurement in medicine, by H. C. W. de Vet, C. B. Terwee, L. B. Mokkink, and D. L. Knol. JBioPharmStat 23:277–279. https://doi.org/10.1080/21679169.2013.737220
26. Kaneko A, Asai N, Kanda T (2005) The influence of age on pressure sensitivity over the cervical musculature associated with neck pain. J Hand Ther Off J Am Soc Hand Ther 18:421–424. https://doi.org/10.1177/033310241506.0042
27. Paul F, Erdfelder E, Lang A-G, Buchner A (2007) GPpower 3: a flexible statistical power analysis program for the social, behavioral, and biomedical sciences. Behav Res Methods 1:75–191
28. Fleiss JL (2011) Design and analysis of clinical experiments. New Jersey: Wiley. https://doi.org/10.1002/9781118032923
29. Rousson V (2013) Measurement in medicine, by H. C. W. de Vet, C. B. Terwee, L. B. Mokkink, and D. L. Knol. JBioPharmStat 23:277–279. https://doi.org/10.1080/21679169.2013.737220
30. Moreira C, Bassi AR, Brando MP, Silva AG (2017) Do patients with chronic neck pain have distorted body image and tactile dysfunction? Eur J Phys 19: 215–221. https://doi.org/10.1080/21679169.2017.1339418
31. Nolan MF (1985) Quantitative measure of cutaneous sensation. Two-point discrimination values for the face and trunk. Phys Ther 65:181–185
32. Elsig S, Luomajoki H, Sattelmayer M et al (2014) Sensormotor tests, such as movement control and laterality judgment accuracy, in persons with recurrent neck pain and controls. A case-control study. Man Ther 19:155–561. https://doi.org/10.1016/j.math.2014.05.014
33. Ashkenazi A, Silberstein S, Jakubowski M, Burstein R (2007) Improved identification of allodynic migraine patients using a questionnaire. Cephalalgia Int J Headache 27:325–329. https://doi.org/10.1111/j.1468-2982.2007.01291.x
34. Ashkenazi A, Silberstein S, Jakubowski M, Burstein R (2007) Improved identification of allodynic migraine patients using a questionnaire. Cephalalgia Int J Headache 27:325–329. https://doi.org/10.1111/j.1468-2982.2007.01291.x
35. Landy S, Rice K, Lobo B (2004) Central sensitisation and cutaneous Allodynia in migraine. CNS Drugs 18:337–342. https://doi.org/10.2165/00023210-200418060-00001
36. Lipton RB, Bigal ME, Ashina S et al (2008) Cutaneous allodynia in the migraine population. Ann Neurol 63:148–158. https://doi.org/10.1002/ana.21211
37. Lovati C, D’Amico D, Rosa S et al (2007) Allodynia in different forms of migraine. Neurol Sci Off J Ital Neurol Soc Ital Soc Clin Neurophysiol 28(Suppl 2):5220–5221. https://doi.org/10.1007/s10072-007-0781-5
38. Nahman-Averbuch H, Shefi T, Schneider VJ et al (2018) Quantitative sensory testing in patients with migraine: a systemic review and meta-analysis. Pain. https://doi.org/10.1097/j.pain.2018.01.1231
39. Gonçalves MC, Chaves TC, Florencia LL et al (2015) Is pressure pain sensitivity over the cervical musculature associated with neck disability in individuals with migraine? J Bodyw Mov Ther 19:67–71. https://doi.org/10.1016/j.jbmt.2014.02.007
40. Florencia LL, Gantomasii MCM, Cevallo GF et al (2015) Generalized pressure pain hypersensitivity in the cervical muscles in women with migraine. Pain Med Malden Mass 16:1629–1634. https://doi.org/10.1111/pme.12767
41. Ashina S, Bendtsen L, Lyngberg AC et al (2015) Prevalence of neck pain in migraine and tension-type headache: a population study. Cephalalgia Int J Headache 35:211–219. https://doi.org/10.1177/0333102415535110
42. Bodguk N (1992) The anatomical basis for cervicogenic headache. J Manipul Physiol Ther 15:67–70
43. Bodguk N, Govind J (2009) Cervicogenic headache: an assessment of the evidence on clinical diagnosis, invasive tests, and treatment. Lancet Neurol 8939–968. https://doi.org/10.1016/S1474-4422(09)70209-1
44. Bartsch T, Goadsby PJ (2003) The trigeminocephalocervical complex and migraine: current concepts and synthesis. Curr Pain Headache Rep 7:371–376
45. Schueler M, Neuhuber WL, De Col R, Messlinger K (2014) Innervation of rat and human dura mater and pericranial tissues in the parieto-temporal region by meningeal afferents. Headache 54:996–1009. https://doi.org/10.1111/head.12371