Does restoration of sagittal cervical alignment improve cervicogenic headache pain and disability: A 2-year pilot randomized controlled trial

Author links open overlay panellbrahim M.Moustafa^{ab}AliaaDiab^bTamerShousha^{ab}Deed E.Harrison^c <u>https://doi.org/10.1016/j.heliyon.2021.e06467Get rights and content</u> Under a Creative Commons license

Abstract

Background

To investigate the <u>feasibility</u> and effect of a multimodal program for improving chronic cervicogenic headache (CGH) via the addition of sagittal <u>cervical</u> <u>spine</u> alignment correction.

Design

Pilot, parallel-group, randomized controlled trial.

Participants

60 patients with CGH, straightening of the cervical <u>lordosis</u>, and <u>forward head</u> <u>posture</u> (FHP) were randomly assigned using permuted-block randomization either to a control (n = 30) or an experimental group (n = 30).

Interventions

Subjects in both groups received a multimodal program where the denneroll cervical spine extension <u>traction orthotic</u> was added to the experimental group only. Feasibility was assessed through recruitment rate, compliance rate, adherence rate, safety, and global satisfaction in addition to clinical outcome measures: FHP distance, cervical lordosis, headache frequency, headache disability inventory (HDI), headache impact

test-6 (HIT-6), and <u>daily defined dose</u> (DDD). Evaluations were performed at: baseline, 10 weeks, 1 year follow up, and 2-year follow up. The <u>assessor</u> was blind to group allocation for all measured outcomes.

Results

The recruitment rate was 60%, 78 % out of them completed the entire study. The recruited participants complied with 98% of the required visits. No <u>adverse</u> <u>events</u> were recorded and greater overall satisfaction with the interventions was reported. Greater improvements were found for the experimental group's cervical lordosis (f = 259.9, P < < .001) and FHP (f = 142.5, P < < .001). <u>At 10</u> weeks, both groups showed equal improvements in CGH outcomes: headache frequency (P = 0.07), HDI (P = 0.07), HIT-6 (P = .2), and DDD (P = .3). In contrast, at the 1-year and 2-year follow up, between group differences were found for all CGH outcomes, P < .00, indicating greater improvement in the experimental group.

Conclusion

The results indicated feasibility for recruitment rate, compliance rate, exercise session adherence, safety, and global satisfaction. At 1-year and 2-year follow-up, the addition of the denneroll orthotic device revealed positive influence on CGH management outcomes.