

# Illusory Finger Stretching and Somatosensory Responses

## Abstract

Resizing illusions, delivered using augmented reality, resize a body part through either stretching or shrinking manipulations. These resizing illusions have been investigated in visuotactile, visual-only, and visuo-auditory presentations. However, the neural underpinnings of these resizing illusions remain undefined. This study sought to understand the neural mechanisms behind these illusions ( $n = 46$ ), by using somatosensory steady state evoked potentials in addition to subjective self-report questionnaires, to enhance knowledge of what drives the subjective embodiment during resizing illusions. Since these Illusions have been shown to provide analgesic effects for individuals with chronic pain conditions, this study also aimed to provide an empirical basis for future investigations in chronic pain samples undergoing resizing illusions. Confirmatory results demonstrated significant differences in subjective experience between non-illusion and multisensory illusion conditions, whilst EEG data measuring somatosensory response across electrodes of interest (F1 & FC1) to 26Hz stimulation to the resized digit showed no significant effects of condition. Exploratory non-parametric SSEP analyses showed a significant effect of condition, with reduced amplitudes in illusory conditions compared to non-illusion conditions, however no significant differences arose in exploratory post hoc tests. These findings demonstrate no clear effect of resizing illusions on SSEP amplitudes for participants without chronic pain, but exploratory findings could hint to a potential sharpening of neural representations as a result of illusory stretching, thereby providing a basis for investigations of comparable subjective and steady state illusion responses in a chronic pain population, who are thought to have more diffuse neural representations of their affected body parts.

## Introduction

Illusory finger stretching is a form of multisensory illusion, specifically a resizing illusion, which alters the subjective perceptual experience of the size of one's finger. Resizing illusions, through changing the way in which a body part is perceived, exploit principles of multisensory

integration to elicit modulations in the perceived size and shape of the body (Preston and Newport 2011; Preston, Gilpin, and Newport 2020; Stanton et al. 2018). Resizing illusions are based on the rubber hand illusion, in which touch is delivered to a visible fake hand at the same time and in the same place that touch is delivered to the hidden real hand. This manipulation elicits feelings of ownership over the fake hand through the integration of multisensory (tactile and visual) inputs highlighting the apparent malleability of bodily self (Botvinick and Cohen 1998). Multisensory resizing illusions typically involve both tactile and visual inputs to the participant and can be delivered via an augmented reality system or through magnifying optics. Recent studies have also shown resizing illusions to be effectively administered through visual only, and visuo-auditory manipulations (Schaefer et al. 2007; Tajadura-Jiménez et al. 2017). However, multisensory visuotactile manipulations are reported as the most effective at inducing a strong experience of the illusion within an augmented reality system (K. J. Hansford et al. 2023).

The augmented reality system used to deliver these resizing illusions presents real-time video capture of the hand, from the same position and perspective as if the hand were being viewed directly (Preston and Newport 2011). This allows the experimenter to deliver tactile manipulations, such as gently pulling or pushing the hand/finger, whilst the participant views their hand/finger either stretching or shrinking in the augmented image. Newport, Pearce and Preston (2010) found strong embodiment using a synchronous multisensory visuotactile illusion, which was replicated in our pilot data using the same experimental set up as the current study. The pilot data showed, although not significant, a numerically greater illusory experience during synchronous visuotactile manipulations compared to asynchronous (mismatching visuotactile manipulation) control conditions (Appendix A) for illusory finger resizing. When comparing multisensory visuotactile resizing illusions to unimodal visual resizing illusions, our recent work (K. J. Hansford et al. 2023) shows that multisensory illusions elicit significantly greater illusory experience compared to non-illusion and unimodal visual illusion conditions in healthy participants. We also showed, in exploratory analysis, that a subset of participants who experienced an illusion in the unimodal visual condition reported a stronger illusory experience in this condition than in an incongruent (mismatching visual and tactile inputs) control condition. This subset analysis, however, was of a small sample size, and was selected based on one of the measures analysed thus should be taken with caution, meaning further replication of the findings are needed. Furthermore, we have demonstrated that a visuo-auditory presentation of the finger resizing illusion, using non-naturalistic auditory input, provides a stronger illusory experience than a visual only presentation, but this does not surpass the illusion strength given by a visuo-tactile illusion (K. Hansford et al. 2023).

Neuroimaging has previously been used in healthy populations experiencing resizing illusions, whereby modulation of the primary somatosensory cortex has been found using neuromagnetic source imaging during visual only resizing illusions of the arm (Schaefer et al. 2007). Briefly, the more the subjects felt the subjective experience of an elongated arm, the more the cortical distance between the first and fifth digit decreased, showing the topographical representation of the somatosensory cortex being modulated by perceived location of a stimulus. Specifically looking at stretching multisensory visuotactile illusions, which as mentioned are those that

elicit the greatest illusion strength in a majority of participants, recent research suggests that these illusions impact the neural representations of the body and reflect early-stage multi-modal stimulus integration through modulation of gamma band activity (Kanayama, Hara, and Kimura 2021). We have recently also investigated this illusion in healthy participants using electroencephalography (EEG) and have found support for this previous research, finding significant increases in gamma band power, likely reflecting multimodal stimulus integration, in multisensory visuotactile compared to unimodal visual conditions during illusory resizing of a finger (K. J. Hansford et al. 2023). Previous research using rubber hand illusions found this multisensory integration effect in early-stage gamma band increases (Kanayama, Hara, and Kimura 2021), whilst our recent findings show a later stage of multimodal stimulus integration when using illusory finger resizing manipulations (K. J. Hansford et al. 2023).

Looking specifically at research into somatosensory cortex modulation using steady-state evoked potentials (hereafter referred to as SSEPs), low-level somatosensory responses have been induced directly using vibrations of a known frequency applied to a body part. These generate a frequency-locked SSEP detectable at the scalp using EEG (Snyder 1992; Tobimatsu, Zhang, and Kato 1999), and are an index of the cortical response to a stimulus. This paradigm has been used with other sensory modalities to better understand the neural mechanisms underlying multisensory integration, with findings showing that presentation of temporally congruent auditory and visual stimuli significantly enhances the magnitude and inter-trial phase coherence of auditory and visual steady-state responses (Nozaradan, Peretz, and Mouraux 2012). Research has also found evidence of enhanced steady-state responses for within-modality stimulation of auditory and visual stimuli in isolation (Giani et al. 2012), complementing Nozaradan et al.’s findings regarding visuo-auditory combination. Research using vibrotactile stimulation has found increases in steady-state response magnitude corresponding with the amplitude modulation rate of stimulation (Colon, Legrain, and Mouraux 2012; Rees, Green, and Kay 1986) suggesting an entrainment of oscillatory activity to temporal features of sensory stimulation (Timora and Budd 2018). Given these findings, we anticipate that SSEPs might change during finger resizing illusions, due to the multisensory manipulations present, to give a potential index of changes in neural representations during the illusion.

Several studies have investigated the analgesic effect of these resizing illusions, as they have been shown to reduce chronic pain in conditions such as osteoarthritis (Preston and Newport 2011; Preston, Gilpin, and Newport 2020; Stanton et al. 2018), chronic back pain (Diers et al. 2013), and complex regional pain syndrome (Moseley, Parsons, and Spence 2008). However, the understanding of how these illusions reduce pain is still undetermined. It has been suggested chronic pain involves cortical misrepresentations of the size of the affected body part (Boesch et al. 2016), however, it is unknown if resizing illusions affect this cortical misrepresentation, and if this is therefore what causes the reduction in pain. No study has yet used neuroimaging with a chronic pain population to determine the cortical activity correlated with this illusory analgesia. However, importantly, there has also not been research conducted using SSEPs in participants without chronic pain, to understand what the cortical representations of these resizing illusions are like without the impact of a chronic pain condition. Therefore, the aim of this study is to examine potential changes in the somatosensory cortex during illusory finger

resizing in participants without chronic pain, using vibrotactile SSEPs, to use as a basis for later investigations in a sample of chronic pain participants. If we can identify a link between illusory resizing and somatosensory cortex changes, this will enhance our understanding of what is happening in the brain during these illusions and will act as a reference for comparison with neural representations in individuals with chronic pain conditions.

Using different sensory manipulations of finger resizing illusions, in addition to using an electromagnetic solenoid stimulator, this study aimed to investigate subjective illusory experience and somatosensory SSEP responses in participants without chronic pain, to better understand the experience of body ownership illusions from subjective experience and cortical representation perspectives. To test this, different finger resizing illusions consisting of multisensory (visuotactile) stretching (MS), unimodal-visual stretching (UV), a non-illusion control condition without tactile input (NI), and a non-illusion control condition with tactile input (NIT) were used to assess alternate aspects of illusory resizing manipulations and their related effects on somatosensory SSEP response. The inclusion of two control conditions (NI, NIT) was to assess whether localisation of cortical representations arise from resizing manipulations to the finger, or from tactile input given to the finger. The first hypothesis, acting as a positive control (1), was that there will be a greater illusory experience, measured via a subjective illusory experience questionnaire, in the (1a) MS condition compared to the NI condition and in the (1b) MS condition compared to the NIT condition. The main experimental hypothesis for this study was that (2) there will be a significant difference in somatosensory SSEP response across the electrodes of interest (F1 & FC1, see Appendix A Pilot Data) when comparing across all conditions. Subsequent hypotheses were that there would be significant differences in SSEP response when comparing (2a) the MS condition to the NI condition, when comparing (2b) the UV condition to the NI condition, but (2c) that there would be no significant difference when comparing the NIT condition to the NI condition.

## Methods

### Sample Size

Overall, based on the power analyses in section 2.5, a total sample size of 46 participants were tested. This sample size adheres to the higher end of sample size estimates (Hypothesis 2 (2.5.2) showing 46 participants needed for post hoc tests 2a – 2c).

### Participants

Ethical approval for this research was gained from the Department of Psychology, University of York (ethics application code 950), in line with the Declaration of Helsinki. All participants gave informed consent prior to the start of any experimental set up, and participants were instructed that they could withdraw their participation at any time during or after completion

of the experiment. 46 participants were tested, with the participant's manipulated finger being randomly split between use of either the index or middle finger. However, 2 participant's data needed removal (>50% of electrodes requiring removal), and therefore 2 additional participants were tested to account for this missing data, both using the index finger as the manipulated digit, resulting in a final sample size of 46 participants (37 Female, 8 Male, 1 Prefer not say; Mean age = 20.3 years, age range = 18.3 – 32.7 years; 32 White, 11 Asian or Asian British, 3 Mixed or Multiple Ethnic Groups; Sample population = students at the University of York). 23 participants were tested using their index finger, the other half using their middle finger.

Sample inclusion / exclusion criteria:

Inclusion and exclusion criteria were determined using self-report responses relating to each item listed below:

- Inclusion Criteria: Right-handed, 18 years of age or over, no older than 75 years of age (include those aged 75 years).
- Exclusion Criteria: Prior knowledge or expectations about the research, a history of developmental, neurological or psychiatric disorders, history of drug or alcohol abuse, history of sleep disorders, history of epilepsy, having visual abnormalities that cannot be corrected optically (i.e. with glasses), or being under 18 years of age, or over 75 years of age. A history of chronic pain conditions, operations or procedures that could damage peripheral nerve pathways in the hands, current experiences of pain or more than 4 hours of consistent pain experienced in the preceding week.

Raw data exclusion criteria:

- Less than 100% of the experiment completed by a participant, more than 50% of electrodes for a single participant requiring removal from EEG data, or if both electrodes F1 and FC1 (electrodes of interest) required removal. More information about data removal can be found in section 2.4.1 Preprocessing Steps.

## **Experimental Procedure**

All participants completed a demographic survey, asking their age, ethnicity, and sex, and were asked to complete the revised Waterloo Handedness Questionnaire (WHQr) (Elias, Bryden, and Bulman-Fleming 1998). The WHQr self-reported handedness questionnaire consists of 36 questions. The questions are answered on a 5-level Likert scale to determine the degree of preferred hand use, with right always being +2, right usually being +1, equal use being 0, left usually being -1, and left always being -2. The sum of the total WHQr score was then used to categorise respondents as left-handed (score of -24 or lower), mixed handed (score of -23 to +23), or right-handed (score of +24 or higher). Only participants who were categorised as right-handed continued participation. Mean handedness score across participants was +57.91 (range = +29 to +71).

Participants were then set up with an appropriately sized 64-channel EEG cap with electrodes arranged according to the 10/20 system. The experimenter used conductive gel to make a conductive bridge between the electrodes and the scalp to attempt to obtain impedance levels of  $<10k\Omega$  per electrode. Data were collected using an ANT Neuroscan system, sampling at 1kHz. The whole head average was used as a reference.

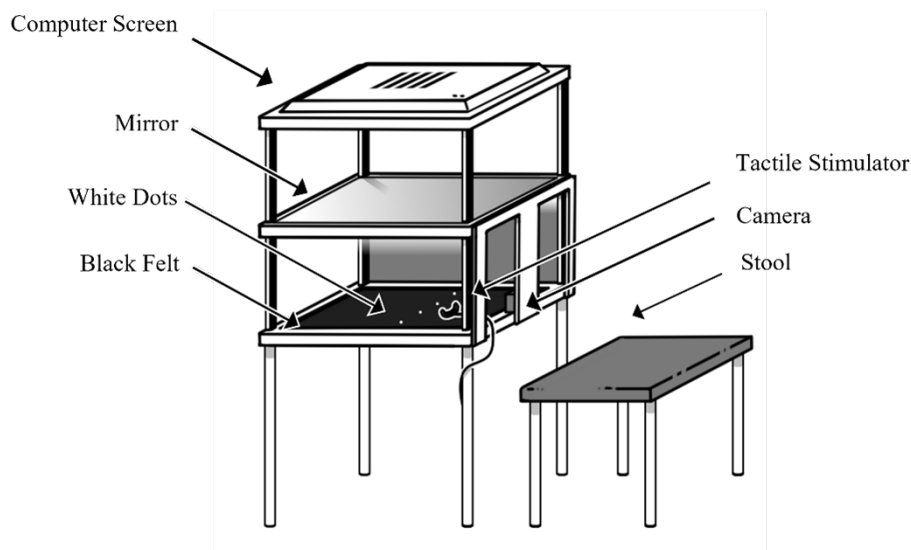


Figure 1: Schematic of Augmented Reality System with Tactile Stimulator.

Participants were then seated behind the augmented reality system (Figure 1) and instructed to place their hand onto the black felt fabric within the augmented reality system. Within the self-built system there was a 1920 x 1080-pixel Spedal Webcam Wide Angle Camera at the edge of the black felt on the side the participant sits, away from the participant's view. 26cms above the felt base, there was a mirror, which was placed 26cms below a screen with a resolution of 1920 x 1200 pixels, with a width of 52cms and a height of 32cms. The thickness of section on which the mirror sat was 2cms. This screen was 54cms from the base of the system, and the base of the system was 82cms from the ground. Participants were instructed to place either their right index or middle finger outstretched onto the felt. The decision of whether the participant used their index or middle finger was pseudo randomised (to give equal representation of each finger) via MATLAB prior to any participants taking part. There were two white dots for each hand on the felt and participants were instructed to place their hand between these two dots. Participants were instructed to view their hand's image in the mirror (whilst the real hand was hidden from view) throughout the experiment. The camera placed underneath the mirror on the felt base was used to deliver a live feed video of the participant's hands to the computer screen at the top of the augmented reality system, which showed in the mirror reflection to the participants. There was a delay of 170ms in the video processing pipeline from the camera image to the augmented video image.

Participants underwent 4 conditions: multisensory stretching (MS), unimodal-visual stretching (UV), a non-illusion control condition without tactile input (NI), and a non-illusion control condition with tactile input (NIT). There was vibrotactile stimulation to the finger in all conditions, but only tactile input of the researcher touching the participants finger in the MS and NIT conditions. Each trial lasted 2.4 seconds for the manipulation phase, where the finger was stretched by 60 pixels (2.1 centimetres) in UV and MS conditions, followed by a further 2.4 second habituation phase in which participants could view and move their (augmented) finger, whilst they keep the rest of their hand still, before the screen went dark, indicating that the next trial could start. The MS condition consisted of the researcher touching and pulling the participant's finger as the participant viewed their finger stretching in a congruent manner. The UV condition consisted of the participants viewing their finger stretch without any experimenter manipulation. The NI condition provided no visual or touching tactile manipulations to the finger, the image of their finger was visible and unchanged throughout. The NIT control condition involved no visual input of the finger stretching, again the image of their finger was visible but unchanged. Additionally, this condition included tactile input of the experimenter's hand touching the participant's finger, but without pulling. Visualisation of all conditions can be seen in Figure 2.

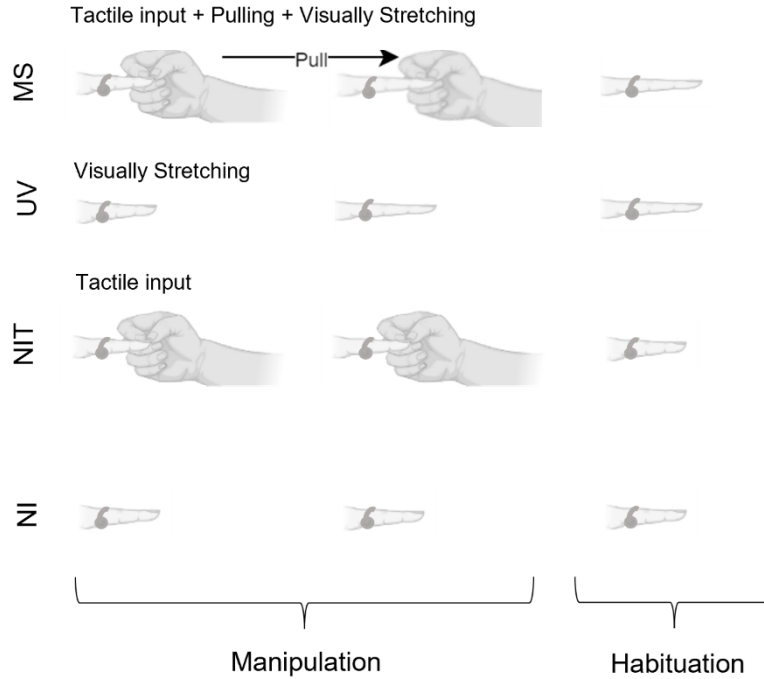


Figure 2: Infographic of Experimental Conditions. MS = Multisensory Stretching, UV = Unimodal Visual Stretching, NIT = Non-Illusion Tactile, NI = Non-Illusion. During the manipulation phase (2.4 seconds) the visual image of the finger is stretched in the MS and UV conditions, and/or the experimenter provides tactile input (touch) in the MS and NIT conditions. The tactile input in the MS condition is accompanied by pulling. During the habituation phase (2.4 seconds) participants are free to move their finger. The arrow denotes the direction of the experimenter’s action. The vibrotactile stimulator is depicted on the finger in each phase of the experiment as vibrations are presented throughout.

The experimenter was seated opposite the participant, the other side of the augmented reality machine and touched the digit during MS and NIT conditions by holding onto the distal interphalangeal joint and gently touching (NIT) or pulling (MS) the finger whilst the participant kept their hand in place. Conditions were delivered across 4 blocks, with each block consisting of 24 trials of the same experimental condition, totalling 96 trials over all 4 blocks. The ordering of the blocks was randomised for each participant to prevent ordering effects. The experiment was programmed in, and the conditions randomised using MATLAB R2017a and the experimenter was informed of whether to pull the finger or to touch the finger via an indicative box displayed on the screen out of the participant’s view. If the box was blue, this



indicated a need to pull the finger, if it was white it indicated a need to touch the finger, if there was no box displayed then this indicated no tactile manipulation from the experimenter. The researcher used a button press to trigger the start of the manipulation, and started pulling the finger, when needed, synchronously within the 2.4 second manipulation phase. If the experimenter were to forget to pull the finger on a multisensory condition, or mistakenly pulled the finger in a control trial, then this would be noted during the experiment, and that trial would be removed from analysis. Fortunately, no trials needed removal due to experimenter error. Vibrations were delivered to the participant’s finger in all conditions using a miniature electromagnetic solenoid stimulator (Dancer Design Tactor; diameter 1.8mm) emitting vibrations produced by sending amplified 26Hz sine wave sound files, with stimulus intensity controlled by an amplifier (Dancer Design TactAmp). The tactor was driven at 50% of the maximum (i.e. a peak input voltage of 3V) using a 26Hz sine-wave, and delivered a peak force of 0.18N. The electromagnetic solenoid stimulator was attached to the participant’s finger that was outstretched to receive the manipulations, between the knuckle and the first finger joint, using clear medical tape and gave continuous stimulation for the duration of each trial. Participants were encouraged to take a break between each of the blocks to stretch their hand. EEG was recorded throughout as a continuous recording with conditions denoted by numbered 8-bit digital at the start of each trial (USB-TTL Module, Black Box Toolkit Ltd.).

Finally, at the end of each block, the participant was asked to complete the subjective illusory experience questionnaire regarding a condition presented in a given block using a Samsung Galaxy Tab A6 tablet via a questionnaire on Qualtrics (Qualtrics, Provo, UT). The questionnaire consisted of six questions relating to the trials the participant had just experienced. Two statements related to illusory experience: “It felt like my finger was really stretching” / “It felt like the finger I saw was part of my body”, two related to disownership: “It felt like the finger I saw no longer belonged to me” / “It felt like the finger I saw was no longer part of my body”, and two were control questions: “It felt as if my finger had disappeared” / “It felt as if I might have had an extra finger” (all questions were directed towards the participants manipulated finger). Control questions were included to create an index for the illusion and disownership questions (more detail can be found in section 2.4.1 - Preprocessing steps), whilst disownership questions were included to assess if the potential experience from the illusions resulted from a disownership of the body part, or from subjective embodiment of the body part (McCabe 2011). A visual analogue scale from 0 – 100 was used for each statement, with 0 being strongly disagree, 50 being neutral and 100 being strongly agree.

Data collection was terminated when the full sample of participants had been tested. If a participant completed <100% of the experiment or if over 50% of electrodes needed removal, or if both electrode F1 and FC1 needed removal, then their data was not included, and additional participants were recruited to replace any lost data.

## Analysis Pipeline

### Preprocessing Steps

EEG data were first converted using MATLAB and EEGLab from the ANT EEProbe .cnt format to EEGLab .set format. All subsequent analysis was then be conducted using the MNE-Python toolbox (Gramfort et al., 2013). A 50Hz notch filter was first applied to the raw EEG data for all electrodes, followed by calculation of the standard error across time for each electrode for each participant (Luck et al. 2021). Across the standard errors for all participants, the 5% of electrodes which showed the largest standard errors were used to create a standard error threshold. Any electrode with a standard error above this threshold, or with a value of 0, was removed from analysis. Where a participant had over 50% of their electrodes over the standard error threshold or with a value of 0, or if the electrodes requiring removal included either electrodes F1 or FC1 (electrodes of interest), then their data was removed. Primary analysis of the remaining EEG data then involved averaging the signal across the electrodes of interest (or using just electrode F1 or FC1 in case of electrode removal), and calculating the Fourier transform for each trial per participant. These amplitudes were then averaged across trials per condition to give overall results for each participant per condition. Statistical comparisons were then performed on the Fourier amplitudes at the stimulation frequency (26Hz), across conditions and participants. No additional filtering or denoising steps were applied to the EEG data, in line with Figueira et al.'s (2022) report that only a Fourier transform is typically needed for this type of EEG data.

Regarding questionnaire data, scores for both illusion experience questions were combined to give median scores, along with both disownership questions and both control questions, resulting in 3 median scores per condition per participant. The median control scores were used to create an index of the illusion and disownership scores by subtracting the median control score from the median illusion and median disownership scores, in line with previous research doing similarly (Matsumiya 2021; Kiltner and Ehrsson 2017; Kalckert and Ehrsson 2012). The normalised (control indexed) data were used for analyses, with a new scale from -100 to +100 with 100 indicating strongly agree, 50 indicating a neutral opinion, and scores below 0 indicating strongly disagree with the statements on the questionnaire. 50 is maintained as a neutral opinion so that the normalised data still adhered to the thresholds that the participants were presented with during the experiment.

### Planned Analyses

#### Hypothesis 1 (Positive Control)

*(1 – Positive Control) There will be a greater illusory experience, measured via a subjective illusory experience questionnaire, in the (1a) MS condition compared to the NI condition and in the (1b) MS condition compared to the NIT condition.*

The subjective illusory experience questionnaire was used as a positive control for the current study. Previous research has shown significantly greater illusion strength for MS conditions compared to non-illusion conditions (Carey et al., 2019; Hansford et al., 2023a), which we attempted to replicate. Questionnaire data were analysed using R (R Core Team, 2021). A one-way ANOVA would be run to compare the dependent variable of normalised (control indexed) illusion score from each independent condition. Given significant findings, post-hoc tests would be run, with Bonferroni correction for 2 comparisons (MS Vs NI, MS Vs NIT) at an initial alpha of 0.05.

Interpretations for hypothesis 1 can be found in the design table (Table 1).

## Hypothesis 2

*(2) There will be a significant difference in somatosensory SSEP response across the electrodes of interest (F1 & FC1). Subsequent hypotheses are that there will be a significant difference in SSEP response in (2a) the MS condition compared to the NI condition, and (2b) the UV condition compared to the NI condition, but (2c) that there will be no significant difference when comparing the NIT condition to the NI condition.*

As mentioned in the EEG pre-processing steps in section 2.4.1, analysis of EEG data involved taking a Fourier transform for each waveform averaged across the electrodes of interest, to obtain the amplitude for each trial at the vibration frequency (26Hz). These amplitudes would then be averaged across trials to give overall results for each participant, before running a repeated measures one way ANOVA comparing somatosensory SSEP response from each experimental condition. The dependent variable would be SSEP amplitude in  $\mu\text{V}$ , whilst the independent variable would be the different manipulations given in each comparison condition. Given significant findings in the ANOVA, post hoc comparisons would be conducted at a new alpha of .016 (corrected for 3 comparisons (MS Vs NI, UV Vs NI, NIT Vs NI)). Based on the pilot data in Appendix A, we expected to see activation most pronounced over mid-frontal distributions, covering F1 and FC1 electrodes and therefore these electrodes were selected as the electrodes of interest.

Interpretations for hypothesis 2 can be found in the design table (Table 1).

## Power analysis

### Hypothesis 1 (Positive Control)

Effect sizes were determined by research from Hansford et al. (2023a) using the subjective illusory experience questionnaire and comparing MS, UV, and incongruent finger-based resizing illusions to control conditions with no illusory resizing, using the same finger stretching illusions and the same equipment ( $n = 48$ ), which show an effect size of  $\eta^2 = .33$  (converted to Cohen's  $f = .70$  and Cohen's  $d = 1.4$ ). Additional effect size information comes from a

visual capture study ( $n = 80$ ) using a subjective embodiment questionnaire and visual and tactile manipulations to a mannequin body (Carey et al., 2019), showing an effect size of  $r = .64$  (converted to Cohen's  $f = .83$ ) when comparing embodiment scores from the questionnaire against control scores. An effect size of  $f = .70$  was used for hypothesis 1 to adhere to the lower end of previous effect sizes.

Hypothesis 1: A priori power analysis using G\*Power for the smallest effect size of interest ( $f = .70$ ) showed that for a repeated measures, within factors one way ANOVA, with an effect size ( $f$ ) of 0.70, alpha level of 0.05, power at 80% and 1 group with four measurements, 5 participants were needed.

Hypotheses 1a and 1b: A priori power analysis using G\*Power shows that for a one-tailed difference between 2 means (pairwise) t test, with an effect size of  $d_z = 1.4$ , alpha of 0.025, power at 80%, a total sample size of 7 participants was required.

## Hypothesis 2

This was the first study to investigate illusory finger stretching using SSEPs, so appropriate effect size estimates were not available. We therefore conducted power calculations based on a smallest effect size of interest, in line with the recommendation of Lakens (2014). Here, we have chosen an effect size of  $d = 0.5$  (a medium effect, see Cohen, 1988), since this is the smallest effect size we were interested in detecting, which we converted to a Cohen's  $f$  of 0.25 for Hypothesis 2's power analysis, and have maintained at 0.5 for the subsequent post hoc power analyses.

Hypothesis 2: A priori power analysis using G\*Power showed that for a repeated measures, within factors one way ANOVA, with an effect size ( $f$ ) of 0.25, alpha of 0.05, power at 80%, and 1 group with four measurements, a total sample size of 24 participants was needed.

Hypotheses 2a – 2c: A priori power analysis using G\*Power shows that for a two-tailed difference between 2 means (pairwise) t test, with an effect size of  $d_z = .5$ , alpha of 0.016 (corrected for multiple comparisons), power at 80%, a total sample size of 46 participants was needed.

The stage 1 report for this study, with details regarding planned analyses and pilot data, can be seen at the following OSF page: <https://osf.io/pfksu/>. All data and code to reproduce the analyses can be found at the following OSF page: <https://osf.io/yhz6j/>. A design planner encompassing research questions, hypotheses, sampling and analysis plans and their resulting interpretations and observed outcomes can be seen in Figure 3.

Question	Hypothesis	Sampling plan (e.g., power analysis)	Analysis plan	Interpretation given different outcomes	Theory that could be proved wrong given outcomes	Observed Outcome
Does the finding of greater subjective illusory experience in multisensory compared to non-illusion conditions replicate in this study?	(1 - Positive Control) There will be a greater illusory experience, measured via a subjective illusory experience questionnaire, in the MS condition compared to the NI and NIT conditions.	A priori power analysis using G*Power shows that for a repeated measures, within factors ANOVA, with an effect size (f) of 0.70, alpha of 0.05, power at 80% and 1 group with four measurements, 5 participants are needed.	An ANOVA will be run to compare mean scores from each condition.	If Hypothesis 1 is supported: Indicates that the augmented reality manipulations are inducing effective illusions, and shows success of positive control, giving weight to the subsequent EEG findings.  If Hypothesis 1 is unsupported: Indicates that the augmented reality manipulations are not inducing effective illusions, and therefore the findings regarding hypotheses 2 will be called into question.	The theory that adding vibrotactile stimulation will not influence the subjective illusion experience of resisting illusions would be proved wrong within this sample if hypothesis is unsupported.	Hypothesis supported.
Does the finding of greater subjective illusory experience in the MS condition compared to the NI condition replicate in this study?	(1a) There will be a greater illusory experience, measured via a subjective illusory experience questionnaire, in the MS condition compared to the NI condition	A priori power analysis using G*Power shows that for a one-tailed difference between 2 means (pairwise) t test, with an effect size of $d_z = 1.4$ , alpha of 0.025, power at 80%, a total sample size of 7 participants is needed	Given significant findings from the ANOVA (Hypothesis 1), post-hoc tests will be run, with Bonferroni correction for 2 comparisons at an initial alpha of 0.05 (adjusted alpha = .025).	If Hypothesis 1a is supported: Indicated that the MS condition is giving a significantly stronger illusion than the NI condition.  If Hypothesis 1a is unsupported: Indicates that tactile input could be needed for an effective non illusion condition, calling NI EEG analyses into question.	The theory that there is significantly greater subjective illusory experience in the MS condition compared to the NI condition would be proved wrong within this sample if the hypothesis is unsupported.	Hypothesis supported.
Does the NIT condition act as an appropriate control condition compared to the MS condition?	(1b) There will be a greater illusory experience, measured via a subjective illusory experience questionnaire, in the MS condition compared to the NIT condition	A priori power analysis using G*Power shows that for a one-tailed difference between 2 means (pairwise) t test, with an effect size of $d_z = 1.4$ , alpha of 0.025, power at 80%, a total sample size of 7 participants is needed	Given significant findings from the ANOVA (Hypothesis 1), post-hoc tests will be run, with Bonferroni correction for 2 comparisons at an initial alpha of 0.05 (adjusted alpha = .025).	If Hypothesis 1b is supported: Indicated that the MS condition is giving a significantly stronger illusion than the NIT condition.  If Hypothesis 1b is unsupported: Indicates that tactile input could remove effect of NIT condition, calling EEG analyses with NIT condition into question.	The theory that tactile input has no effect on the non-illusion control condition would be proved wrong if this hypothesis is unsupported.	Hypothesis supported.
Are there significant changes in the somatosensory response when comparing different conditions healthy participants?	(2) There will be a significant difference in somatosensory SSEP response across the electrodes of interest (F1 & FC1) when comparing across all conditions	A priori power analysis using G*Power shows that for a repeated measures, within factors one way ANOVA, with an effect size of $f = .25$ , alpha of 0.05, power at 80%, and 1 group with four measurements, a total sample size of 24 participants is needed.	A repeated measures one way ANOVA will be run comparing all experimental conditions. The dependant variable will be somatosensory SSEP amplitude in $\mu V$ .	If Hypothesis 2 is supported: Indicates that there are significant differences in somatosensory SSEP response when comparing across all conditions in healthy participants.  If Hypothesis 2 is unsupported: Indicates that there is no evidence of a significant difference in somatosensory SSEP response when comparing across all conditions in healthy participants. This will result in hypotheses 2a - 2c being unsupported.	The theory regarding cortical changes in somatosensory representation during illusory finger stretching would be proved wrong if hypothesis 2 is unsupported.	Hypothesis supported.
Are there significant changes in the somatosensory response when comparing the multisensory vibrotactile to the non-illusion condition in healthy participants?	(2a) There will be a significant difference in somatosensory SSEP response across the electrodes of interest (F1 & FC1) when comparing the MS illusory resisting condition to the NI condition.	A priori power analysis using G*Power shows that for a two-tailed difference between 2 means (pairwise) t test, with an effect size of $d_z = .5$ , alpha of 0.015, power at 80%, a total sample size of 46 participants is needed.	A post hoc comparison will be run after the repeated measures one way ANOVA to compare the MS condition to the NI condition. The dependant variable will be somatosensory SSEP amplitude in $\mu V$ .	If Hypothesis 2a is supported: Indicates that there are significant differences between MS and the NI condition in healthy participants.  If Hypothesis 2a is unsupported: Indicates that there is no evidence of a difference between MS and the NI condition in a healthy population.	The theory regarding cortical changes in somatosensory representation during multisensory illusory finger stretching would be proved wrong if hypothesis 2a is unsupported.	Hypothesis unsupported.
Are there significant changes in the somatosensory cortex when comparing the unimodal visual condition to the non-illusion condition in healthy participants?	(2b) There will be a significant difference in somatosensory SSEP response across the electrodes of interest (F1 & FC1) when comparing the UV illusory resisting condition to the NI condition.	A priori power analysis using G*Power shows that for a two-tailed difference between 2 means (pairwise) t test, with an effect size of $d_z = .5$ , alpha of 0.015, power at 80%, a total sample size of 46 participants is needed.	A post hoc comparison will be run after the repeated measures one way ANOVA to compare the UV condition to the NI condition. The dependant variable will be somatosensory SSEP amplitude in $\mu V$ .	If Hypothesis 2b is supported: Indicates that there are significant differences between UV and the NI condition in healthy participants.  If Hypothesis 2b is unsupported: Indicates that there is no evidence of a difference between UV and the NI condition in a healthy population.	The theory regarding cortical changes in somatosensory representation during unimodal visual illusory finger stretching would be proved wrong if hypothesis 2b is unsupported.	Hypothesis unsupported.
Are there significant changes in the somatosensory cortex when comparing the non-illusion tactile condition to the non-illusion condition in healthy participants?	(2c) There will be no significant difference in somatosensory SSEP response across the electrodes of interest (F1 & FC1) when comparing the NIT condition to the NI condition.	A priori power analysis using G*Power shows that for a two-tailed difference between 2 means (pairwise) t test, with an effect size of $d_z = .5$ , alpha of 0.015, power at 80%, a total sample size of 46 participants is needed.	A post hoc comparison will be run after the repeated measures one way ANOVA to compare the NIT condition to the NI condition. The dependant variable will be somatosensory SSEP amplitude in $\mu V$ .	If Hypothesis 2c is supported: Indicates that there are no significant differences between NIT and NI conditions in healthy participants.  If Hypothesis 2c is unsupported: Indicates that there is evidence of a difference between NIT and NI conditions in a healthy population.	The theory that tactile input alone will not give significant differences in cortical changes in somatosensory representation during non-illusion conditions would be proved wrong if hypothesis 2c is unsupported.	Hypothesis supported.

Figure 3: Design Planner

## Results

Positive control analyses of the subjective illusion data can be seen in Figure 4. A one-way ANOVA found a significant overall effect of condition with a large effect size ( $F(3,135)$ ,  $p.adj = <0.001$ ,  $\eta^2 = 0.229$ ). Post hoc t tests with Bonferroni corrections found significantly greater combined illusion score in the MS condition ( $M = 61.79$ ,  $SD = 28.31$ ) compared to the Non-Illusion (NI;  $M = 31.2$ ,  $SD = 26.08$ ,  $t = -5.67$ ,  $p.adj = <0.001$ , Cohen's  $d = -31$ ) and Non Illusion Tactile (NIT;  $M = 37.41$ ,  $SD = 20.59$ ,  $t = -5.61$ ,  $p.adj = <0.001$ , Cohen's  $d = -24$ ) conditions, thereby supporting hypotheses 1, 1a, and 1b and fulfilling the positive control checks.

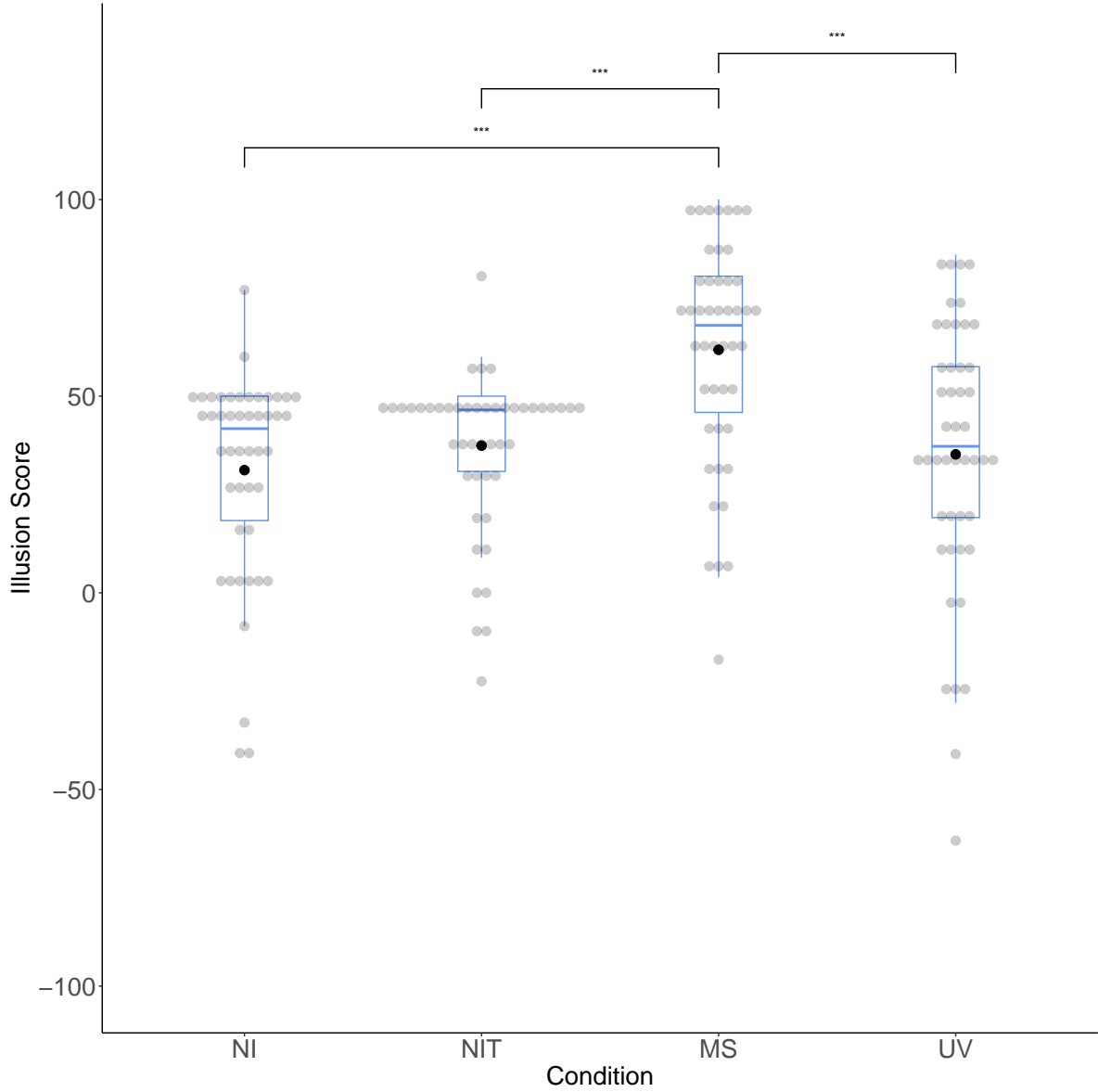


Figure 4: Combined Illusion Score Index Across Conditions (NI: Non-Illusion; NIT: Non-Illusion Tactile; MS: Multisensory; UV: Unimodal Visual). Scores below 50 indicate disagreement with experience of illusion statements, whilst scores above 50 indicate agreement. A continuous visual analogue scale was used in data collection, with agreement and disagreement statements located at each end of the scale. Box plots show means, medians and inter-quartile ranges of data. Medians are indicated with a horizontal line whilst means are indicated by a black dot. Data points are shown in grey jitter binned along the y-axis, grouped by condition.

Analyses of SSEP data can be seen in Figure 5b. The left panel confirms the presence of a clear steady-state signal at 26Hz, which was strongest over the fronto-central electrodes. A one-way ANOVA found no significant effect of condition with a small effect size ( $F(3,135)$ ,  $p = 0.209$ ,  $\eta p^2 = 0.033$ ), opposing Hypothesis 2. Post hoc t tests with Bonferroni corrections found no significant differences between SSEP amplitude when comparing the NI condition ( $M = 0.49$ ,  $SD = 0.76$ ) to the MS condition ( $M = 0.31$ ,  $SD = 0.57$ ,  $t = 1.7$ ,  $p.adj = 0.571$ , Cohen's  $d = 0.18$ ), or UV condition ( $M = 0.36$ ,  $SD = 0.83$ ,  $t = 1.15$ ,  $p.adj = 1.000$ , Cohen's  $d = 0.13$ ), meaning Hypotheses 2a and 2b were unsupported. There was no significant difference found when comparing the NI condition to the NIT condition ( $M = 0.29$ ,  $SD = 0.35$ ,  $t = 2.02$ ,  $p.adj = 0.298$ , Cohen's  $d = 0.19$ ), supporting Hypothesis 2c.

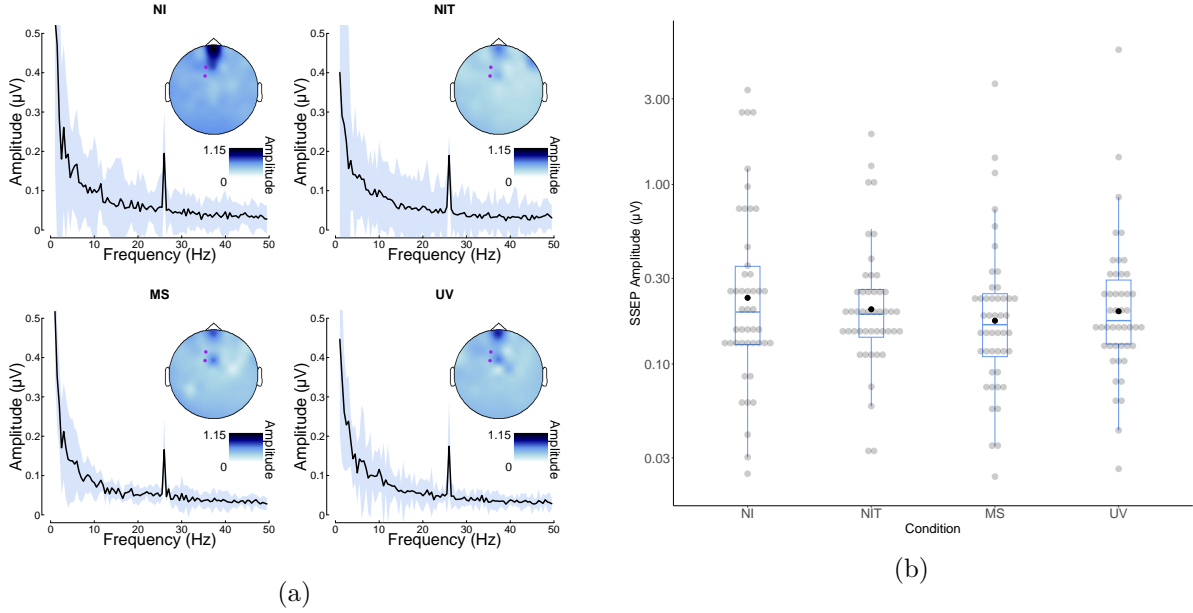


Figure 5: (a): SSEP Amplitude Spectra Across Conditions (NI: Non-Illusion; NIT: Non-Illusion Tactile; MS: Multisensory; UV: Unimodal Visual) for electrodes of interest (F1 & FC1). Shading shows  $\pm 1$  standard error across participants ( $n=46$ ). (b): SSEP Amplitudes Across Conditions. Box plots show means, medians and interquartile ranges of data. Medians are indicated with a horizontal line whilst means are indicated by a black dot. Data points are shown in grey jitter binned along the y-axis, grouped by condition.

## Exploratory Analyses

Since illusion data violated assumptions for parametric tests, an exploratory Friedman test was run and found a significant overall effect of condition with a moderate effect size ( $\chi^2(3) = 42.05$ ,  $p = <0.001$ , Kendall's  $W = 0.3$ ) and post hoc Wilcoxon tests with Holm corrections found

significantly greater combined illusion score in the Multisensory Stretching (MS) condition ( $M = 68$ ,  $SD = 28.31$ ) compared to the Non-Illusion (NI;  $M = 41.75$ ,  $SD = 26.08$ ,  $z = 103$ ,  $p.adj = <0.001$ ,  $r = -0.29$ ), Non Illusion Tactile (NIT;  $M = 46.5$ ,  $SD = 20.59$ ,  $z = 118$ ,  $p.adj = <0.001$ ,  $r = -0.23$ ) and UV conditions (UV;  $M = 37.25$ ,  $SD = 34.37$ ,  $z = 903.5$ ,  $p.adj = <0.001$ ,  $r = 0.27$ ).

In addition to illusion data, disownership and control data were collected and therefore analyses on these datasets have also been run. Exploratory analysis of subjective disownership and control data can be seen in Appendix B. A significant increase in disownership scores were found in the UV condition compared to all other conditions, and there were no significant comparisons found for control data. All statistical reporting can be seen in Appendix B.

EEG data also violated assumptions for parametric tests and therefore a Friedman test was also run and found a significant overall effect of condition with a small effect size ( $\chi^2(3) = 8.17$ ,  $p = 0.043$ , Kendall's  $W = 0.06$ ), however, post hoc Wilcoxon tests with Holm corrections found no significant differences between SSEP amplitude when comparing the NI condition ( $M = 0.19$ ,  $SD = 0.76$ ) to the MS condition ( $M = 0.17$ ,  $SD = 0.57$ ,  $z = 725$ ,  $p.adj = 0.131$ ,  $r = 0.05$ ), or UV condition ( $M = 0.17$ ,  $SD = 0.83$ ,  $z = 719$ ,  $p.adj = 0.131$ ,  $r = 0.03$ ). There was no significant difference found when comparing the NI condition to the NIT condition ( $M = 0.19$ ,  $SD = 0.35$ ,  $z = 686$ ,  $p.adj = 0.131$ ,  $r = 0.03$ ).

Exploratory correlational analyses were also conducted to assess the correlation between participant's subjective illusion score and their SSEP amplitude across electrodes of interest (F1 & FC1) for each condition to see if those who experienced a stronger feeling of the illusion had more reduced SSEP amplitudes, results showed no significant correlations and can be seen in Appendix C.

## Discussion

This study sought to understand both subjective and neural responses to resizing illusions in participants without chronic pain, to provide not only a greater understanding of how bodily illusions affect cortical representations, but also a basis for investigating differences in cortical representations between participants with and without chronic pain conditions when using resizing illusions for analgesic treatment. Subjective data replicated previous findings of greater subjective illusory experience in multisensory compared to non-illusion conditions, showing that the addition of vibrotactile stimulation does not appear to impact subjective experience of resizing illusions using augmented reality, since these effects replicate ones found previously without vibrotactile stimulation. EEG data showed no significant effect of condition when assessing SSEP amplitudes across the electrodes of interest (F1 & FC1) at 26Hz. Exploratory non-parametric analyses of SSEP amplitudes showed a significant effect of condition however, with a decreased median amplitude in the multisensory and unimodal visual conditions compared to the non-illusion condition. However, these differences did not reach statistical significance with exploratory post hoc tests. These findings, therefore, demonstrate



that within exploratory analyses illusory resizing can lead to reductions in SSEP amplitude, but this finding would need to be replicated as a confirmatory analysis and a larger sample size would likely be needed to detect any significant differences in post hoc comparisons due to the small effect sizes found.

Whilst the subjective illusory experience data supported the positive control hypothesis of the multisensory condition providing greater illusory experience than either of the non-illusion conditions, exploratory analyses found that the unimodal-visual condition demonstrated a significantly reduced experience of the illusion compared to the multisensory condition. This reduction in illusory experience for the unimodal visual condition compared to the multisensory condition was also found in our previous work (K. J. Hansford et al. 2023), and similarly shows a more diverse range of responses compared to the multisensory condition. These findings reinforce the idea that not everyone experiences resizing illusions with only visual stimuli, and this should be considered when assessing the application of resizing illusions to chronic pain samples, as if subjective experience of the illusion is required for analgesic effects, then it is possible that not everyone will experience this from a unimodal visual presentation. Exploratory data looking at disownership of the digit during illusory resizing found significantly greater experiences of disownership in the unimodal visual condition compared to the multisensory, non-illusion, and non-illusion tactile conditions. This heightened disownership might explain the reduced illusory experience in the unimodal visual condition, as it could be that the presence of tactile input is needed during illusory resizing to ground the digit within the augmented reality system, otherwise feelings of disownership can arise.

Regarding SSEP data, the reduced amplitudes seen in the exploratory analyses could be explained through the somatosensory blurring / sharpening hypothesis (Haggard, Iannetti, and Longo 2013). This theory proposes that the somatosensory representation of a body part can be sharpened through improved tactile discrimination and acuity training. This sharpening is thought to represent increased organisation of the somatosensory area responding to the stimuli (Haggard, Iannetti, and Longo 2013). Tactile acuity can be increased through simply viewing an enlarged body part (Kennett, Taylor-Clarke, and Haggard 2001). Therefore, it is likely that the enlarged digits created through illusory resizing are sharpening the somatosensory representations of the digits. The reduced amplitudes found during exploratory analyses in the illusory conditions compared to the non-illusion conditions therefore could demonstrate a neural representation of this somatosensory sharpening. However, since these differences were not found to be significant through confirmatory analyses or exploratory post hoc tests, it is likely that for people without chronic pain there are not clear changes in SSEP response during illusory resizing.

A possible explanation for the SSEP reductions found could be through the direction of attention to the digits in illusory conditions. However, previous research has found that attending to a specific vibrotactile stimulus can result in an increase, rather than a reduction, in SSEP amplitude (Giabbiconi et al. 2004). Furthermore, brain computer interfaces (BCIs) are used to intentionally modify a brain signal that can be detected by a computer to manipulate one's

environment, and these are often based on increasing SSEP response amplitudes through directing attention (Muller-Putz et al. 2006). Therefore, it is unlikely that the reduction in SSEP amplitudes seen here are due to increased attention in tactile and illusory manipulation conditions. It is possible though, that if somatosensory sharpening is occurring, the reduction in amplitude associated with this could be confounded by this attentional effect, with somatosensory sharpening reducing amplitudes whilst attention is increasing the amplitudes, resulting in the small effect sizes we see in this dataset. Due to these small effect sizes, replication of these effects in larger samples might be better able to assess somatosensory sharpening or attentional effects, as here the sample size was only powered to detect at least medium effects.

Further exploratory analyses assessed correlations between subjective illusory experience and SSEP amplitude across electrodes of interest and found no significant correlations for any condition. These findings could indicate that subjective experience of the illusion is not required for there to be changes in cortical responses, although without clear support for changes in SSEP amplitudes found within the confirmatory analyses, this suggestion cannot be empirically justified. It is possible that SSEP amplitudes are too noisy to show such somatosensory changes, or that the sample needed to detect these effects would have to be larger than the one in the present study. However, when considering resizing illusions as a non-pharmaceutical method for pain reduction, a lack of correlation between SSEP amplitude and illusory experience could mean that patients do not need to subjectively experience resizing illusions for there to be the potential of illusory analgesia. Future research is needed to consolidate both this hypothesis and the exploratory correlational findings from the present study.

One of the main aims of the present study was to provide a basis for investigating somatosensory representations of illusory resizing in samples with hand-based chronic pain. Illusory resizing has been found to provide analgesic effects for hand-based chronic pain (Preston and Newport 2011), however the neural underpinnings of this analgesia remain undefined. Since chronic pain is thought to create blurred somatosensory representations of the painful body part (Haggard, Iannetti, and Longo 2013), it is possible that when comparing the results seen here in participants without chronic pain to a sample of participants with chronic pain, the differences between amplitudes in the non-illusion and illusion conditions could be greater, due to more blurred initial representations of the painful digits. If somatosensory response changes are found in a sample with chronic pain, then these changes could underscore the analgesia experienced after illusory resizing, however if these changes are either not seen or do not align with pain reduction, then alternate mechanisms are likely behind illusory resizing analgesia. It is, however, possible that since there were no SSEP effects found through confirmatory analyses in the present study, that the impact of illusory resizing on SSEP responses could be too small to meaningfully detect in both population groups, especially since a patient group could have more varied and / or noisy data.

## Conclusions

The present study enhances our understanding of whether there are cortical changes associated with illusory resizing in people without chronic pain and provides an empirical basis for later investigations of somatosensory response changes in a sample with chronic pain. The subjective data suggest that vibrotactile stimulation does not affect experience of resizing illusions, and therefore highlights the suitability of this method for eliciting somatosensory steady state evoked potentials in future investigations. Confirmatory analyses of SSEP data showed no clear effect of illusory resizing on SSEP amplitudes, however, trends toward supporting the somatosensory blurring / sharpening hypothesis were found within exploratory analyses whereby reduced amplitudes were seen in both illusory conditions compared to the non-illusion conditions. If similar reductions are observed in a sample with chronic hand-based pain, then it would be possible to assume that these neural response changes could be driving illusory analgesia.

## Appendix

Pilot data regarding the experience of the illusion for healthy participants undergoing synchronous and asynchronous illusory resizing of the index finger can be seen in Figure 7. 9 participants had either synchronous or asynchronous multimodal manipulations delivered first in a random order, and were then given the other condition, after which all participants were given an illusion scale. Findings showed that across all participants, no significant difference in illusion experience between the synchronous and asynchronous conditions,  $t(8) = 1.877$ ,  $p = 0.097$ , however as can be seen in figure A1, despite the small sample size, illusion strength was seen to be greater in the synchronous condition compared to the asynchronous condition.

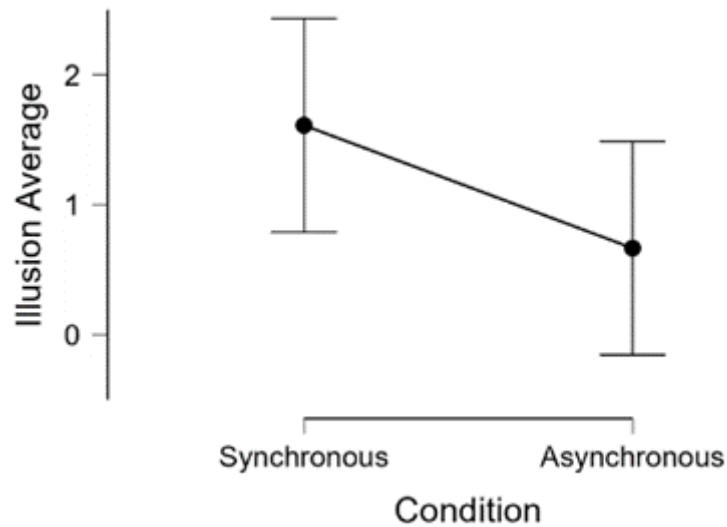


Figure 6: Pilot data from Healthy Participants Undergoing Synchronous and Asynchronous Illusory Finger Resizing.

Previous literature stated that the ideal vibration frequency to use to elicit somatosensory SSEPs is approximately 26Hz (Muller et al., 2001; Breitweiser et al., 2016; Pokorny et al., 2016; Snyder, 1992). Due to resizing illusions often manipulating the index finger, and previous studies using the index finger supporting around 26Hz as an optimal frequency (Muller et al., 2001; Breitweiser et al., 2016; Pokorny et al., 2016), it was hypothesised that 26Hz would elicit a dependable somatosensory SSEP response. Therefore, we ran a pilot study to check that our setup and equipment can reliably elicit and record a somatosensory SSEP at 26Hz, using the resizing illusion and EEG.

Pilot data were collected for 3 healthy participants. Participants underwent the same experimental protocol as mentioned in the “Experimental Procedure” section, minus the subjective illusory experience questionnaire. A Fourier transform was calculated for each waveform at each electrode for all conditions, and then averaged across repetition to obtain individual results. These were then averaged across all 3 participants to give the result seen in Figure 8.

As can be seen in Figure 8, there was a clear somatosensory SSEP response at 26Hz, which was strongest around electrodes F1 and FC1. Previous research using vibrotactile stimulation at 21Hz have also found the scalp topography of the activation to be most pronounced over mid-frontal distributions (Porcu et al., 2014; Timora & Budd, 2018), in line with the scalp topography seen here. Given these finding of a distinct 26Hz signal and mid-frontal scalp location, it appeared appropriate for 26Hz to be used as the vibration frequency in the current study.

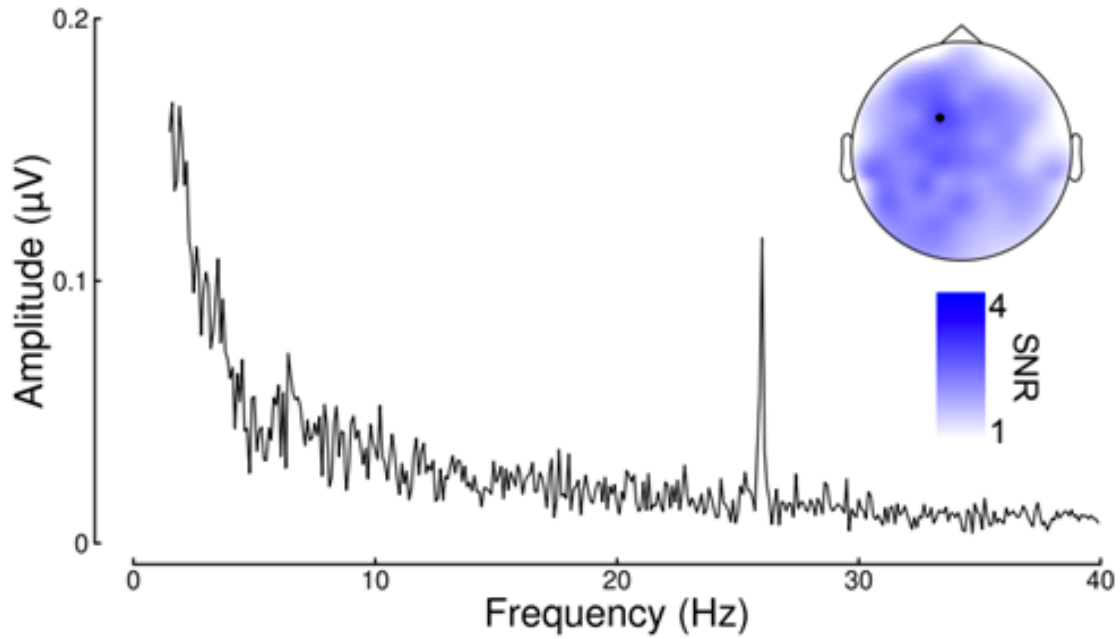


Figure 7: Averaged Pilot Data showing peak frequency at 26Hz, centred between electrodes F1 and FC1. The spectrum is derived from electrode FC1. Saturation bar represents signal to noise ratio (SNR). SNR is a measure of signal quality and describes the ratio of signal power (at 26Hz) to noise power (averaged across 10 adjacent frequency bins). SNR was used for the pilot figure because with a small sample (3 participants) we did not want a noisy electrode to influence the electrodes chosen as electrodes of interest.

Pilot data were also collected using the vibrotactile stimulator at 26Hz to make sure that the illusory experience is not affected by the addition of vibrotactile input. Pilot data were collected from 4 additional healthy participants, who underwent the same experimental protocol as mentioned in the “Experimental Procedure” section, but without EEG caps fitted. Illusory experience was calculated using the median of both illusion scores for each participant minus their median control scores, as per the preprocessing steps regarding the control index, and then the data were averaged over participants to give the results seen in Figure 9. As can be seen, there is a greater subjective experience of the resizing illusion, indexed by participant’s illusion score, in both experimental conditions (UV average = 64.25; MS average = 67.88) compared to both control conditions (NI average = 32.38; NIT average = 24.13). Scores below 50 are indicative of disagreement of experience of the illusion, whilst a score of 50 is a neutral option regarding the illusion experience, and scores above 50 are indicative of agreement of experiencing the illusion. This therefore shows that the experience of illusory resizing was maintained when vibrotactile stimulation was added to the procedure and can therefore be used in the current study to elicit somatosensory SSEPs without affecting the subjective illusory experience of the resizing illusion.

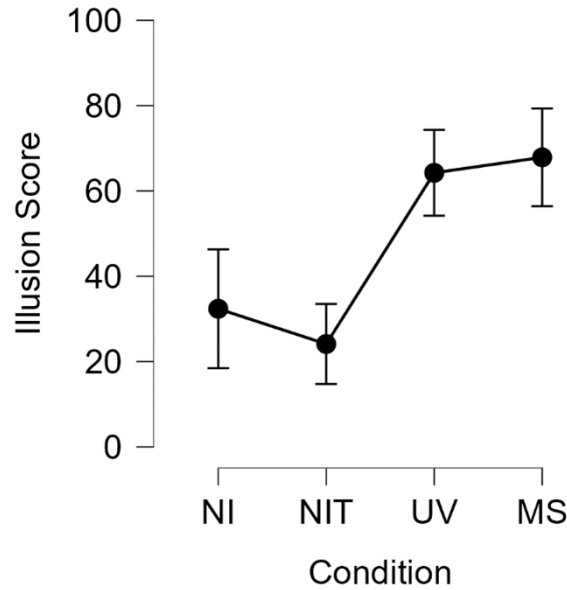


Figure 8: Averaged Illusion score for each condition. Error bars represent standard errors. NI represents the non-Illusion condition, NIT refers to the non-illusion tactile condition, UV refers to the unimodal-visual condition, and MS refers to the multisensory condition.

Exploratory analyses of the subjective disownership data using a Friedman test found a significant overall effect of condition with a small effect size ( $\chi^2(3) = 19.33$ ,  $p = <0.001$ , Kendall's  $W = 0.14$ ) and post hoc Wilcoxon tests with Holm corrections found significantly greater combined disownership score in the UV condition (Median = 23.25, SD = 27.6) compared to the NI (Median = 1.75, SD = 22.4,  $z = 211.5$ ,  $p.adj = 0.014$ ,  $r = -15.5$ ), NIT (M = 0, SD = 19.8,  $z = 132$ ,  $p.adj = <0.001$ ,  $r = -18.25$ ), and MS conditions, (Median = 7, SD = 25.11,  $z = 254$ ,  $p.adj = 0.033$ ,  $r = -10.25$ ).



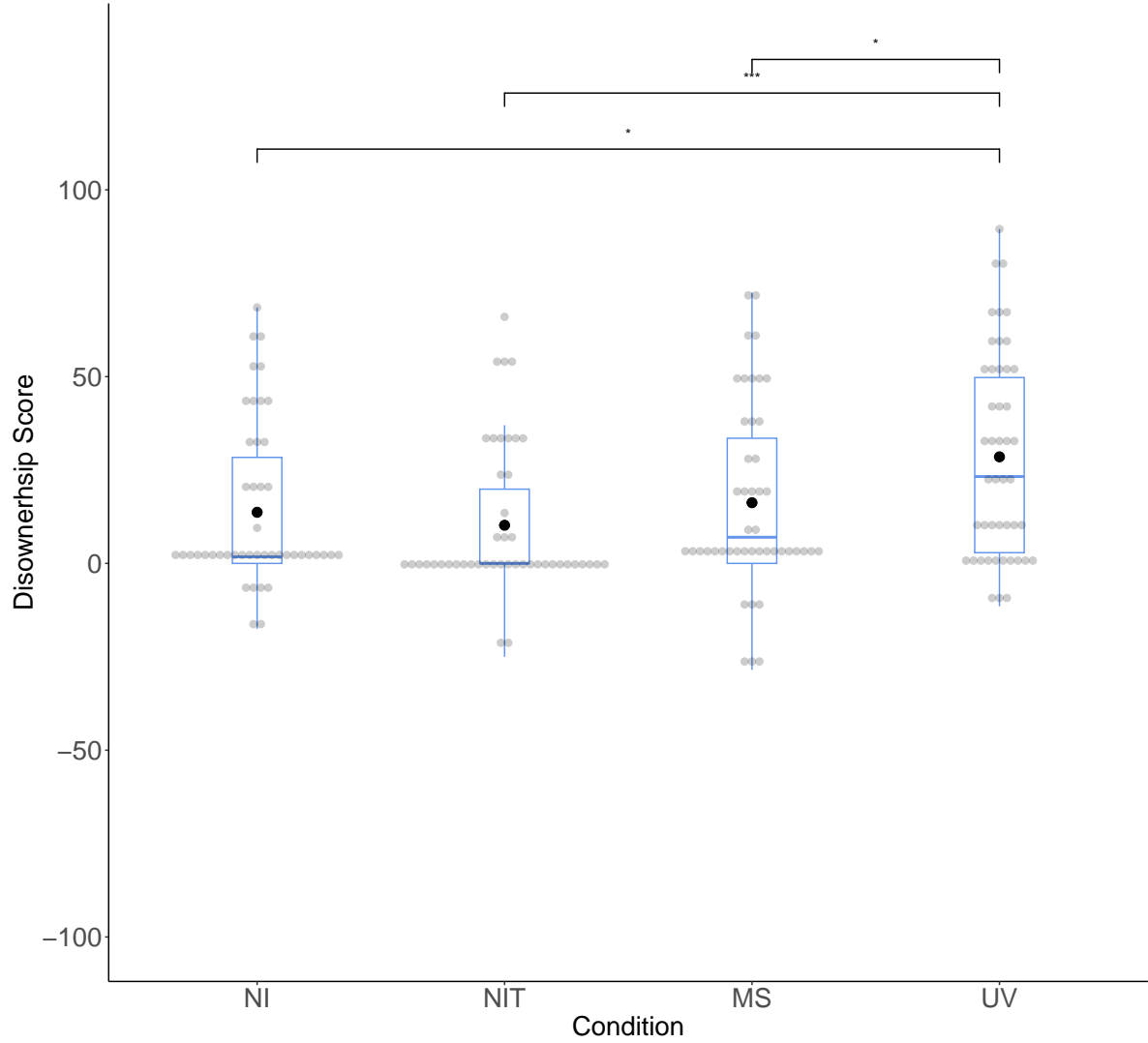


Figure 9: Combined Disownership Score Index Across Conditions (NI: Non-Illusion; NIT: Non-Illusion Tactile; MS: Multisensory; UV: Unimodal Visual). Scores below 50 indicate disagreement with experience of disownership statements, whilst scores above 50 indicate agreement. A continuous visual analogue scale was used in data collection, with agreement and disagreement statements located at each end of the scale. Box plots show means, medians and inter-quartile ranges of data. Medians are indicated with a horizontal line whilst means are indicated by a black dot. Data points are shown in grey jitter binned along the y-axis, grouped by condition.

Exploratory analyses of the subjective control data using a Kruskal-Wallis test (due to multiple ties) found no significant overall effect of condition ( $\chi^2(3) = 4.53$ ,  $p = 0.210$ ) and post hoc Dunn's tests with Bonferroni corrections found no significant differences between combined control scores when comparing the NI condition (Median = 0, SD = 15.58) to the NIT condition (Median = 0, SD = 13.24,  $z = 1$ ,  $p.adj = 1.000$ ), or the MS (Median = 0, SD = 18.15,  $z = 0.02$ ,  $p.adj = 1.000$ ), or UV conditions (Median = 1.5, SD = 16.34,  $z = -1.13$ ,  $p.adj = 1.000$ ).

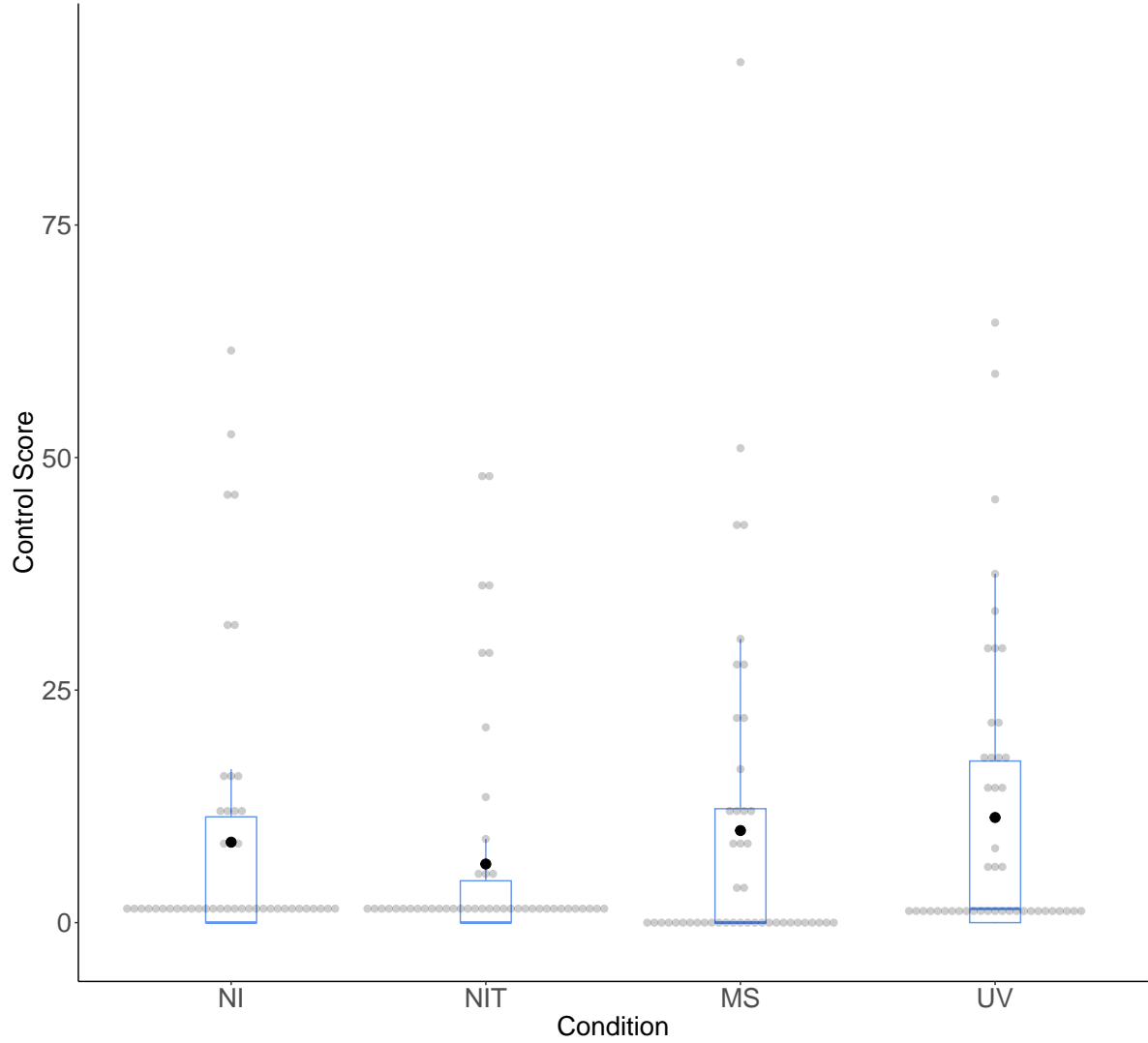


Figure 10: Combined Control Scores Across Conditions (NI: Non-Illusion; NIT: Non-Illusion Tactile; MS: Multisensory; UV: Unimodal Visual). Scores below 50 indicate disagreement with experience of control statements, whilst scores above 50 indicate agreement. A continuous visual analogue scale was used in data collection, with agreement and disagreement statements located at each end of the scale. Box plots show means, medians and inter-quartile ranges of data. Medians are indicated with a horizontal line whilst means are indicated by a black dot. Data points are shown in grey jitter binned along the y-axis, grouped by condition.

Spearman's correlations were run to identify any correlations between participant's illusion score and their SSEP amplitude across electrodes of interest (F1 & FC1) for each condition found no significant correlations across conditions.

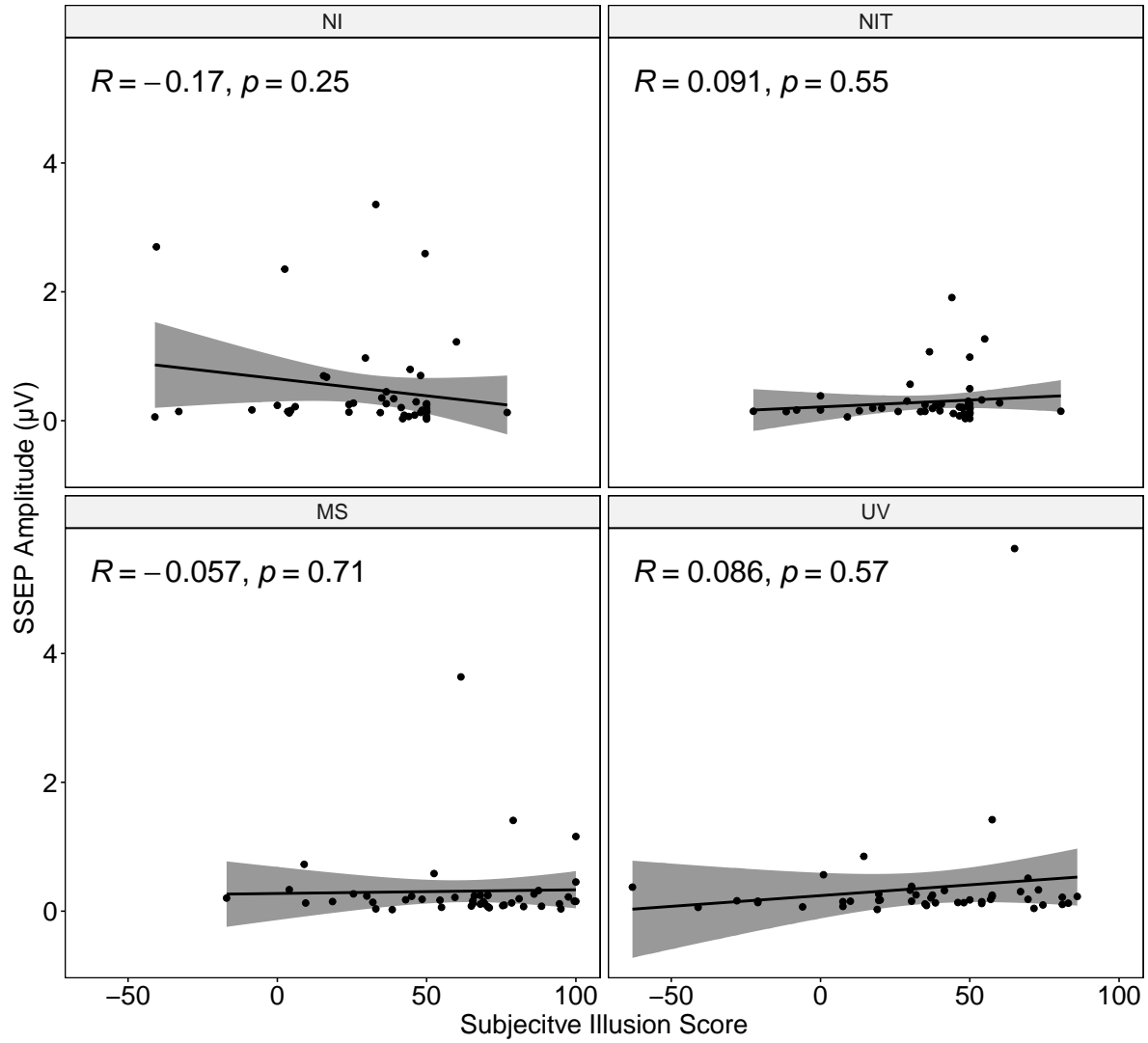


Figure 11: Correlation Between Amplitude and Subjective Illusory Score for Each Condition (NI: Non-Illusion; NIT: Non-Illusion Tactile; MS: Multisensory; UV: Unimodal Visual).

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