Visually induced analgesia during face or limb stimulation in healthy and migraine subjects

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Background: Visually induced analgesia (VIA) defines a phenomenon in which viewing one’s own body part during its painful stimulation decreases the perception of pain. VIA occurs during direct vision of the stimulated body part and also when seeing it reflected in a mirror. To the best of our knowledge, VIA has not been studied in the trigeminal area, where it could be relevant for the control of headache.

Subjects and methods: We used heat stimuli (53°C) to induce pain in the right forehead or wrist in 11 healthy subjects (HSs) and 14 female migraine without aura (MO) patients between attacks. The subjects rated pain on a visual analog scale (VAS) and underwent contact heat-evoked potential (CHEP) recordings (five sequential blocks of four responses) with or without observation of their face/wrist in a mirror.

Results: During wrist stimulation, amplitude of the first block of P1–P2 components of CHEPs decreased compared to that in the control recording when HSs were seeing their wrist reflected in the mirror (p = 0.036; Z = 2.08); however, this was not found in MO patients. In the latter, the VAS pain score increased viewing the reflected wrist (p = 0.049; Z = 1.96). Seeing their forehead reflected in the mirror induced a significant increase in N2 latency of CHEPs in HSs, as well as an amplitude reduction in the first block of P1–P2 components of CHEPs both in HSs (p = 0.007; Z = 2.69) and MO patients (p = 0.035; Z = 2.10). Visualizing the body part did not modify habituation of CHEP amplitudes over the five blocks of averaged responses, neither during wrist nor during forehead stimulation.

Conclusion: This study adds to the available knowledge on VIA and demonstrates this phenomenon for painful stimuli in the trigeminal area, as long as CHEPs are used as indices of central pain processing. In migraine patients during interictal periods, VIA assessed with CHEPs is within normal limits in the face but absent at the wrist, possibly reflecting dysfunctioning of extracephalic pain control.

Keywords: visually induced analgesia, CHEPs, migraine, vision, pain, mirror therapy

Introduction

Cortical networks engaged in the processing of sensory stimuli largely overlap for the various sensory modalities including nociceptive stimuli.1

The latter study using functional magnetic resonance imaging (fMRI) concluded that the largest part of networks activated by phasic nociceptive stimuli reflected non-nociceptive-specific cognitive processes mainly driven by salience of the applied stimulus and showed that the extent of cortical activation was largest for the visual modality.
Interactions between various sensory modalities (such as touch or vision) and pain processing have been suspected and partly studied for several decades, beginning with the gate control theory of Melzack and Wall.1,2 Hence, viewing one’s own body accelerates tactile reactions, increases tactile spatial acuity, and modulates somatosensory-evoked potentials.3–5 Touch can reduce pain levels and cortical pain processing.6–9 Finally, tactile discrimination training reduces chronic central pain,10,11 and this effect is enhanced by seeing the body.10–12

Therapeutic studies have suggested that vision of the body is able to reduce chronic phantom limb pain.13,14 In this disorder, the reflection of the damaged body part given by the mirror is thought to reorganize and integrate the mismatch between the subject’s proprioception and the actual visual feedback and thus to relieve phantom limb pain, probably via slow neuroplastic changes.15–17 Recently, the mirror therapy and partly studied for several decades, beginning with the gate control theory of Melzack and Wall.2 Hence, viewing one’s own body accelerates tactile reactions, increases tactile spatial acuity, and modulates somatosensory-evoked potentials.3–5 Touch can reduce pain levels and cortical pain processing.6–9 Finally, tactile discrimination training reduces chronic central pain,10,11 and this effect is enhanced by seeing the body.10–12

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Longo et al21,22 have recently demonstrated the existence of a visually induced analgesia (VIA) mechanism in healthy subjects (HSs). This physiological phenomenon is characterized by the reduction in pain perception and pain-related evoked potentials during a painful infrared laser stimulation when the subject sees the area of his or her body where the painful stimulus is applied.21,22 VIA is obtained during direct vision and also when indirectly seeing the body part reflected in a mirror; however, it is absent when viewing someone else’s body part. VIA was demonstrated in peripheral limbs but has not yet been studied in the face, ie, in the trigeminal area.

In migraine patients, interactions between vision and the trigeminal pain system are likely to occur. Photophobia is a hallmark of migraine attacks, but it is also prevalent between attacks. Its mechanisms have been addressed in several recent studies.23 We have recently shown in healthy volunteers that the nociception-specific blink reflex (nsBR), a trigeminal nociceptive brain stem reflex, is increased by inhibiting the visual cortex with low-frequency repetitive transcranial magnetic stimulation but decreased by visual activation with 8 Hz flash light stimulation, suggesting an inhibitory top–down relation between the visual cortex and second-order trigeminal nociceptors.24

Based on the assumption that pathways linking the visual and trigeminal systems might be impaired in migraine, we decided to study VIA in the trigeminal territory using thermoneceptive stimuli and comparing HSs with migraine patients between attacks.

Subjects and methods

Subjects

We recruited 11 female HSs (age, 29.45 ± 10.25 years, mean ± SD) and 14 patients suffering from migraine without aura (MO; code 1.1) according to ICHD-3 beta criteria (MO, 14 females; age, 26.4 ± 4.55 years, mean ± SD).25 HSs had no personal or familial history of primary headaches and were devoid of any significant somatic or psychiatric diseases and of any drug treatment except the contraceptive pill. They were recruited among medical students and hospital personnel. MO patients were recruited among those attending our outpatient headache clinic and were not allowed to take preventive therapy or any other drug between attacks. They suffered from migraine since 11.07 ± 4.9 years and had on average 2.02 ± 1.52 attacks per month with a mean attack duration of 21.57 ± 17.85 hours. They were recorded in the interictal phase, ie, they were attack free for at least 3 days before and after the recording sessions, which was verified by phone or e-mail.

To avoid changes of cortical excitability due to hormonal variations, females were recorded during mid-cycle.26

This study was approved by the local ethics committee of the CHR Citadelle Hospital, Liège, Belgium, and conducted following the principles of the Declaration of Helsinki. All participants gave written informed consent prior to testing.

Methods

We assessed heat pain using a 0–10 visual analog scale (VAS) for the subjective evaluation of pain intensity and contact heat-evoked potentials (CHEPs) as an objective index of pain processing.27–30

CHEPs

The CHEPs stimulation unit (Medoc Ltd., Ramat Yishai, Israel) is composed of a thermode applied to the skin and covers a cutaneous area of 572.5 mm² (diameter 27 mm). The thermode comprises an external layer consisting of a heating foil and a subjacent Peltier element with two thermistors (electronic thermal sensors). The heating thermofoil (Minco Products, Inc., Minneapolis, MN, USA) is covered with a 25 µm layer of thermoconductive plastic (Kapton®; DuPont, Wilmington, DE, USA; thermal conductivity at 23°C of 0.1–0.35 W/m/K) that separates the external foil from the skin. Two thermocouples are embedded at 10 µm within this conductive coating, which is in direct contact with the skin, allowing an estimation of skin temperature at the thermode surface.30,31
Baseline skin temperature was kept constant at 35°C. Twenty brief heat stimuli were delivered. The peak was set at 53°C, the heating rate was 70°C/s, and the cooling decrement that started as soon as 53°C was reached was 40°C/s, for a total stimulus duration of 707 ms. The interstimulus interval varied randomly between 10 and 22 seconds. The thermode was placed on the right volar wrist or right forehead.

The cortical responses evoked by the heat stimuli (CHEPs) were recorded using pin electrodes inserted at Cz (active) and Fz (reference) according to the international 10–20 system, with a band pass of 0.15–100 Hz (CED™ 1902 preamplifier and CED™ Micro1401 converter; Cambridge Electronic Design Ltd, Cambridge, UK). A ground electrode was fixed to the right hand. The impedance for all electrodes was kept <5 kΩ. Twenty responses were averaged off-line and partitioned into five blocks of four responses using Signal™ software version 4.11 (Cambridge Electronic Design Ltd). We focused on the following Aδ fiber-related CHEP components identified according to latency: P1 as the first most positive point around a latency of 200 ms for the wrist and 150 ms for the face, N2 as the following negative peak around a latency of 280 ms for the wrist and 250 ms for the face, and P2 as the second-most positive point around a latency of 400 ms for the wrist and 350 ms for the face (Figure 1).

Subjects were asked to rate the global heat-evoked pain on a VAS from 0 to 10 just after each series of 20 stimuli.

**Study protocol**

Subjects sat with open eyes in a comfortable armchair in the neurophysiological laboratory room with dimmed light. CHEPs and heat pain intensity were recorded at baseline without sight of the corresponding body part, first during stimulation of the wrist followed after ±10 minutes by forehead stimulation. Thereafter the recordings were repeated in the same sequence and with similar intervals while subjects were seeing the wrist or forehead in a mirror (Figure 2).

**Data processing and statistical analysis**

The latencies (in milliseconds) of the three CHEP components and peak-to-peak P1–N2 and N2–P2 and amplitudes (in microvolt) were measured. To assess more reliably the global brain activity generated by the thermonociceptive stimulus, we also determined the P1–P2 area under the curve (AUC; µV·ms) of each block of four averaged responses. As we found no difference between the results obtained for amplitude and for AUC measures, we chose to report only the latter.

Habituation was defined as the change in P1–P2 AUC over the five successive blocks of averaging expressed as the slope of the linear regression line of AUC changes. In addition, we measured habituation as the percentage change in P1–P2 AUC between the fifth and the first block. The two methods yielded similar results; thus, we report only on slope values.

Statistica for Windows version 8.0 (StatSoft, Inc. Tulsa, OK, USA) was used for all statistical analyses. Wilcoxon’s
test was applied to search for differences in latencies, P1–P2 AUC, and slope between baseline and mirror vision. The Mann–Whitney U-test was used to make comparisons between groups. The significance level was \( p \leq 0.05 \).

## Results

Table 1 displays all experimental data as mean ± SD values and \( p \) values for changes between baseline recordings and those with vision of body part with the mirror.

All CHEP recordings obtained after right forehead stimulation were analyzable, while the recordings after wrist stimulation of two subjects (one HS and one MO) were excluded because of insufficient signal quality.

During wrist stimulation, we found a significant decrease in the first block P1–P2 AUC in HSs when they were seeing their wrist reflected in the mirror compared to the baseline recording (\( p = 0.036; Z = 2.08; \) Figure 3A) where there was no change in pain ratings (Figure 3B). By contrast, in MO patients, using the mirror had no effect on CHEP amplitudes, but the VAS pain score increased significantly at the wrist (\( p = 0.049; Z = 1.96; \) Figure 3B). There were no significant changes in P1, N2, and P2 latencies between baseline recordings and those with the mirror (Table 2).

When the heat stimulus was applied to the forehead, there was a significant decrease in N2 latency in HSs when they watched their face in the mirror (\( p = 0.04; Z = 1.99; \) Table 2). In both HSs and migraine patients, using the mirror significantly decreased first block CHEP P1–P2 AUC (\( p = 0.007; Z = 2.69 \) and \( p = 0.035; Z = 2.10 \), respectively; Figure 3C). Global average over all 20 P1–P2 AUC responses was significant in HSs using the mirror with a decreased effect (\( p = 0.001; Z = 3.17; \) Table 2). Conversely, there was no difference in VAS pain scores between baseline and vision with the mirror (Figure 3D).

The habituation slope of CHEP P1–P2 AUC over the five blocks of four averaged responses was not modified by seeing the wrist or forehead in the mirror in either the HSs or migraine patients (Table 2 and Figure 4).

Finally, there were no significant differences between HSs and migraine patients in any of the baseline CHEPs or pain score values.

## Discussion

Our study shows that VIA can be demonstrated in healthy and migraine subjects in the trigeminal area, as far as CHEPs are used as indices of central pain processing. In healthy volunteers, but not in migraine patients, we also found that VIA for CHEPs can be obtained by thermonociceptive stimulation of the wrist, which is in line with the reports by Longo et al.\(^ {21,22} \) who used laser heat-evoked potentials.

Contrary to the latter, however, pain ratings were not significantly attenuated in our study by vision of the stimulated body part, despite a numerical decrease in pain ratings in healthy volunteers. Another difference is that all our subjects were females, while in the cohort of Longo et al.,\(^ {22} \) only three out of 14 subjects were females. Gender differences in pain perception were well documented. Women reported indeed a higher pain sensitivity with various types of noxious stimuli (eg, ischemic, pressure, electrical, and thermal), with the magnitude of gender differences depending on various factors such as sample size, nature of the noxious stimulus, or use of behavioral or verbal indices of pain sensitivity.\(^ {34–38} \)

The reason for enrolling only women in our study was that we aimed at comparing HSs with migraine patients who are predominantly females. A weakness of our study is that we

## Table 1: Mean demographic data ± SD.

<table>
<thead>
<tr>
<th></th>
<th>HS ( n = 11 )</th>
<th>MO ( n = 14 )</th>
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<tbody>
<tr>
<td>Women ( n )</td>
<td>11</td>
<td>14</td>
</tr>
<tr>
<td>Age (years)</td>
<td>29.45 ± 10.25</td>
<td>26.5 ± 4.55</td>
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<tr>
<td>Duration of history of migraine (years)</td>
<td>11.07 ± 4.9</td>
<td></td>
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<tr>
<td>Attack frequency/month (n)</td>
<td>2.02 ± 1.52</td>
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<tr>
<td>Attack duration (hours)</td>
<td>21.57 ± 17.85</td>
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</tbody>
</table>

Abbreviations: HS, healthy subject; MO, migraine without aura; SD, standard deviation.
Figure 3 CHEP amplitudes (P1–P2 AUC of first block) (A and C) and VAS pain ratings (B and D) after heat stimulation at the wrist (A and B) or forehead (C and D) in HSs and migraine patients before (baseline: dark bars) and during vision of the hand or face reflected in a mirror (light bars; mean ± SD).

Abbreviations: CHEP, contact heat-evoked potential; AUC, area under the curve; VAS, visual analog scale; HS, healthy subject.

Table 2 Synopsis of experimental data (significant changes are in bold; mean ± SD).

<table>
<thead>
<tr>
<th></th>
<th>HSs (n = 10)</th>
<th>MO patients (n = 13)</th>
<th>p-value</th>
<th>HSs (n = 11)</th>
<th>MO patients (n = 14)</th>
<th>p-value</th>
</tr>
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<tbody>
<tr>
<td><strong>Wrist heat stimulation</strong></td>
<td></td>
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<tr>
<td>P1 latency (ms)</td>
<td>208.74 ± 31.36</td>
<td>212.01 ± 48.40</td>
<td>0.76</td>
<td>203.28 ± 39.26</td>
<td>192.29 ± 44.19</td>
<td>0.64</td>
</tr>
<tr>
<td>N2 latency (ms)</td>
<td>272.75 ± 39</td>
<td>272.97 ± 42.82</td>
<td>0.77</td>
<td>278.54 ± 54.25</td>
<td>273.35 ± 43.47</td>
<td>0.55</td>
</tr>
<tr>
<td>P2 latency (ms)</td>
<td>404.15 ± 27.79</td>
<td>388.48 ± 27.70</td>
<td>0.18</td>
<td>422.61 ± 75.14</td>
<td>403.11 ± 74.88</td>
<td>0.08</td>
</tr>
<tr>
<td>First block AUC P1–P2 (µV ms)</td>
<td>2.17 ± 1.38</td>
<td>1.63 ± 1.37</td>
<td>0.03</td>
<td>1.47 ± 0.66</td>
<td>1.47 ± 0.86</td>
<td>0.97</td>
</tr>
<tr>
<td>Average on 20 responses' AUC P1–P2 (µV ms)</td>
<td>1.44 ± 1.51</td>
<td>1.22 ± 1.07</td>
<td>0.16</td>
<td>0.98 ± 0.60</td>
<td>1.03 ± 0.65</td>
<td>0.64</td>
</tr>
<tr>
<td>Slope P1–P2 over 5 blocks</td>
<td>−0.27 ± 0.18</td>
<td>−0.20 ± 0.17</td>
<td>0.33</td>
<td>−0.16 ± 0.15</td>
<td>−0.12 ± 0.25</td>
<td>0.50</td>
</tr>
<tr>
<td>VAS (0–10)</td>
<td>5.90 ± 2.28</td>
<td>5.10 ± 2.54</td>
<td>0.12</td>
<td>5.58 ± 1.57</td>
<td>6.25 ± 1.51</td>
<td>0.04</td>
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<tr>
<td><strong>Forehead heat stimulation</strong></td>
<td></td>
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<td></td>
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<tr>
<td>P1 latency (ms)</td>
<td>165.09 ± 12.13</td>
<td>160.94 ± 31.36</td>
<td>0.87</td>
<td>164.62 ± 9.64</td>
<td>162.24 ± 11.09</td>
<td>0.53</td>
</tr>
<tr>
<td>N2 latency (ms)</td>
<td>247.31 ± 19.02</td>
<td>244.37 ± 31.45</td>
<td>0.049</td>
<td>253.52 ± 36.26</td>
<td>248.70 ± 29.95</td>
<td>0.85</td>
</tr>
<tr>
<td>P2 latency (ms)</td>
<td>376.14 ± 29.69</td>
<td>373.24 ± 29.98</td>
<td>0.24</td>
<td>395.68 ± 39.5</td>
<td>376.46 ± 45.83</td>
<td>0.12</td>
</tr>
<tr>
<td>First block AUC P1–P2 (µV ms)</td>
<td>2.60 ± 1.55</td>
<td>1.63 ± 0.7</td>
<td>0.007</td>
<td>2.75 ± 1.97</td>
<td>2.23 ± 1.42</td>
<td>0.03</td>
</tr>
<tr>
<td>Average on 20 responses' AUC P1–P2 (µV*ms)</td>
<td>1.87 ± 1.25</td>
<td>1.20 ± 0.60</td>
<td>0.001</td>
<td>1.94 ± 1.40</td>
<td>1.55 ± 0.87</td>
<td>0.18</td>
</tr>
<tr>
<td>Slope P1–P2 over 5 blocks</td>
<td>−0.18 ± 0.33</td>
<td>−0.17 ± 0.14</td>
<td>0.86</td>
<td>−0.20 ± 0.38</td>
<td>−0.21 ± 0.27</td>
<td>0.47</td>
</tr>
<tr>
<td>VAS (0–10)</td>
<td>5.91 ± 1.92</td>
<td>5.00 ± 2.88</td>
<td>0.26</td>
<td>5.92 ± 2.6</td>
<td>6.38 ± 1.74</td>
<td>0.47</td>
</tr>
</tbody>
</table>

**Abbreviations**: HS, healthy subject; MO, migraine without aura; AUC, area under the curve; VAS, visual analog scale.
did not correct VAS pain ratings for certain features such as
social context, BMI, and height, which can have an effect
on pain perception.39

More importantly, a lack of correlation between amplitude
of nociceptive-evoked cortical potentials and subjective pain
perception was demonstrated in several studies undermin-
ing previous suggestions of a linear correlation between
the two.40–42 Pain perception engages a complex multifocal
interconnected network in the brain, while CHEPs more
simply reflects activity in the cingulate gyrus. This led to
the now prevailing concept that amplitude of pain-evoked
potentials is driven by saliency of the stimulus rather than
by pain perception.40,43

VIA is a complex phenomenon for which there are
several physiological explanations. It involves chiefly the
primary somatosensory cortex (SI) and the operculo-insular
cortex.22 Visuotactile stimulation increases activation of the
somatosensory cortex more than does touch alone.44 An
fMRI study with infrared laser stimulation showed that VIA
was not associated with an overall reduction in the cortical
response elicited by the painful stimulus but that it increased
connectivity between the brain’s pain network (formerly the
“pain matrix”) and posterior brain areas activated by the
visual perception of the body (or “visual body network”),
resulting in modulation of the experience of pain.22 Recent
studies have focused on the role of the extrastriate cortex
in VIA.45 Interestingly, in the latter study, excitatory anodal
tDCS over extrastriate areas increased the VIA phenomenon,
while cathodal tDCS had no effect. These results are in line
with our previous study where excitatory flash light stimula-
tion increased the pain threshold.24

The connection between visual and somatosensory corti-
ces is probably modulated by the thalamus. This is supported
by a study showing that visual stimulation activates the
orbitofrontal but not the somatosensory cortex in thalamecto-
mized cats, suggesting that projections from the occipital
cortex have to pass through the thalamus to be conveyed to
the somatosensory cortex.46 Interestingly, in this study, the
lateral geniculate nucleus was intact, showing that it is not
an obligatory relay for visuo-somatosensory connections.
However, in our work we included only MO patients. To
better understand the role of the normal or dysfunctioning
of visual cortex in the phenomenon of VIA, it would be of
interest in future studies to explore patients suffering from
migraine with visual aura and complex neurological auras.

There were several differences between HSs and migraine
patients during interictal periods in our study. While in both
groups, VIA was demonstrable with CHEPs derived from
forehead stimulations, there was a concordant numerical
decrease in pain ratings in HSs but rather an increase in
migraineurs. Moreover, contrary to the former, there was no
VIA for CHEPs at the wrist in the latter, and this was accom-
panied by a significant increase in pain scores. These results
in migraine patients came as a surprise, as we expected the
opposite given that sensitization of the trigeminal nociceptive
system is considered a hallmark of migraine.47 Admittedly,
while cranial cutaneous allodynia may persist in migraine
patients between attacks, it is more prevalent during the
attack and in patients with frequent or chronic migraine.48,49
In our study, the mean attack frequency was rather low (2.02/
month) and patients were recorded at a delay of at least 3 days
from an attack. Hence, it is unlikely that they had persistent

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**Figure 4** Habituation of CHEPs P1–P2 (AUC) over five blocks of four averaged responses in healthy women (blue) and migraine women (orange) after stimulation of the
wrist (A) or the forehead (B) without (light color) and with (dark color) mirror. There was no statistical difference.

**Abbreviations:** CHEP, contact heat-evoked potential; AUC, area under the curve.
This phenomenon might be useful for treating an attack, maybe as an add-on to increase efficacy of acute drug therapies. Further studies are needed to verify this hypothesis.

**Acknowledgment**

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**Disclosure**

The authors report no conflicts of interest in this work.

**References**