Correlation between Electromyographic Reflex and MR Imaging Examinations of the Trigeminal Nerve

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BACKGROUND AND PURPOSE: Previous studies have shown that clinical localization of trigeminal nerve lesions is inaccurate as compared with MR imaging findings. The purpose of our study was to ascertain the added value of electromyographic (EMG) investigation of the trigeminal nerve reflexes for the improvement of lesion localization and for the preselection of patients for MR imaging.

METHODS: We reviewed the EMG studies of the trigeminal reflexes and the MR imaging studies of 20 patients with unilateral symptoms and signs related to the trigeminal nerve (40 trigeminal nerves examined). The results of the two studies were compared to assess the value of EMG in predicting MR imaging outcome. Lesion localization as demonstrated by EMG was compared with localization at MR imaging. MR imaging was used as the standard of reference.

RESULTS: Eight (40%) of 20 patients had MR imaging findings related to presenting trigeminal symptoms, including five brain stem lesions and three peripheral lesions. Fourteen (70%) of 20 patients had EMG abnormalities related to presenting symptoms and signs. For brain stem lesions, lesion localization as shown by EMG corresponded well with MR imaging findings. EMG yielded a sensitivity of 100%, a specificity of 81%, a positive predictive value of 57%, and a negative predictive value of 100% in predicting MR imaging results. Interobserver agreement was good for both the EMG reflex and MR imaging examinations.

CONCLUSION: Our data suggest that EMG recordings of the trigeminal reflexes can be used to exclude structural lesions in patients with symptoms related to the trigeminal nerve. When a lesion is localized in the brain stem with EMG, a tailored MR imaging examination of this region may be sufficient.

Patients with symptoms related to the trigeminal nerve may present with a broad spectrum of clinical findings, including facial pain, either typical trigeminal neuralgia or atypical pain, numbness, paresthesias, and weakness or trismus of the masticator muscles. At physical examination, impaired pain, touch and temperature sensations, or a decreased or absent corneal reflex may be found (1, 2). MR imaging is considered the imaging method of choice for evaluation of patients with symptoms related to the trigeminal nerve (3, 4). These patients may have lesions anywhere from the brain stem nuclei to the distal extracranial branches (3, 4). The brain stem trigeminal nuclei extend from the upper midbrain to the lower medulla oblongata (1–4). In a previous radiologic study of patients with trigeminal neuropathy, it was suggested that clinical localization of trigeminal nerve lesions was extremely inaccurate as compared with MR imaging findings and that the entire course of the trigeminal nerve should always be visualized (4).

Electromyographic (EMG) investigation of the trigeminal nerve reflexes, including the blink reflex, the masseter inhibitory reflex, and the jaw-jerk reflex, may provide valuable additional information about the site of a lesion that cannot be obtained with physical information (5). When accurate localization of a lesion is possible with EMG, more tailored MR examinations might be possible, limiting MR imaging time. In a previous study of 112 consecutive patients with trigeminal nerve symptoms, 61% of the patients had abnormalities on MR images (6). Preselection of patients on the basis of EMG findings may increase the yield of MR imaging in the evaluation of patients with symptoms and signs related to the trigeminal nerve.
The purpose of our study was to compare the results of EMG examinations of the trigeminal nerve reflexes with MR imaging findings.

Patients and Methods

Patients

Between May 1992 and August 1997, 20 patients (11 men and nine women), aged 25 to 63 years (mean age, 42 years), with unilateral trigeminal nerve symptoms (nine with left-sided and 11 with right-sided symptoms) underwent EMG of the trigeminal reflexes and MR imaging. MR imaging diagnoses were confirmed at surgery and by histopathologic examination, by typical clinical course, or by further radiologic follow-up. Medical records were reviewed to assess type and laterality of presenting symptoms. For patients with negative MR imaging studies, follow-up findings were also evaluated.

EMG

Blink reflex response latencies to supraorbital nerve stimulation on either side were recorded with a fine concentric needle electrode in the orbicularis oculi muscle, according to a technique described previously (7–11). The supraorbital nerve and the sensory ophthalmic root form the common afferent limb; the facial nerve, the common efferent limb. In this reflex arc, there is an early unilateral, R1, response at about 10 milliseconds, relayed through an oligosynaptic pathway through the principal trigeminal nucleus in the middle third of the pons. The later bilateral, R2, responses at about 30 milliseconds are relayed through a more complex pathway in the medullary region and terminate in the most caudal part of the spinal trigeminal nucleus. From this area, R2 is conducted through ipsilateral and contralateral polysynaptic pathways through the lateral tegmental field before making connections with the facial nuclei (5, 10, 13–15).

Masseter inhibitory reflex latencies were recorded simultaneously from both masseter muscles by needle or surface electrodes after stimulation of the mental nerve on either side during maximal clenching of the teeth. The masseter inhibitory reflex consists of early symmetrical and late phases of EMG silent periods, with the first silent period (SP1) occurring at 10 to 15 milliseconds and the second silent period (SP2) at 40 to 50 milliseconds, interrupting the voluntary EMG activity in the ipsilateral and contralateral masseter muscles (16, 17). After stimulation of the mental nerve, impulses reach the pons via the sensory mandibular root of the trigeminal nerve. The S1 response is mediated by one inhibitory interneuron projecting onto jaw-closing motoneurons bilaterally. The whole circuit lies in the midpons (5, 16, 17) (Fig 2). The afferents for S2 descend in the spinal trigeminal tract and connect with a polysynaptic chain of excitatory interneurons, located in the lateral tegmental field, at the level of the pontomedullary junction. The last interneuron of the circuit is inhibitory and gives rise to ipsilateral and contralateral collaterals that ascend medially to the right and left spinal trigeminal complexes, to reach the trigeminal motoneurons (5, 16, 17) (Fig 2).

To elicit the jaw-jerk reflex, the examiner holds one finger on the subject’s chin and taps it with a reflex hammer. EMG responses are recorded simultaneously from the two sides by surface electrodes placed on the masseter muscle belly (5). The jaw-jerk reflex circuit involves the ipsilateral midbrain and midpons. The afferent nerve fibers have their cell bodies in the trigeminal mesencephalic nucleus, which has collateral links with the trigeminal motor nucleus in the pons (5, 18, 19) (Fig 3).

Topodiagnostic implications of trigeminal reflex abnormalities have been described previously (5, 12, 15, 20).

EMG studies were reviewed by a neurologist and a neurology resident with expertise in the brain stem reflexes who had knowledge of the clinical or MR imaging findings described herein. The EMG study was considered positive when a delay
in the peak or an absent latency was found. Interobserver variability was assessed with \( \kappa \) statistic. Discrepancies were resolved by consensus.

**MR Imaging**

MR imaging was performed on a 1.5-T unit. The patients underwent a standard trigeminal nerve MR imaging protocol, which included axial proton density– and T2-weighted spin-echo images (3800/22–90/1 [TR/TE/excitations]) or fast spin-echo images (3500/22–90/1) with a 3-mm section thickness, a 23-cm field of view, and a 192 \( \times \) 256 matrix through the pons (including the orbit and maxillary sinus), extending to the inferior mandible if the third division of the trigeminal nerve (V3) was involved; and coronal T1-weighted spin-echo images (570–610/14–15/2) with a 3-mm section thickness, a 23-cm field of view, and a 192 \( \times \) 256 matrix through the pons (including the orbit and maxillary sinus), extending to the inferior mandible if the third division of the trigeminal nerve (V3) was involved; and cor- 

**EMG/MR Imaging Correlation**

Lesion localization as demonstrated with EMG was compared with localization at MR imaging. MR imaging was the standard of reference. EMG findings were correlated with MR imaging results in a 2 \( \times \) 2 table to determine the sensitivity, specificity, negative and positive predictive value, and accuracy of the EMG relative to MR imaging.

**Results**

The clinical, EMG, and MR imaging findings are summarized in Table 1.

**EMG Findings**

Overall, EMG findings were abnormal in 14 trigeminal nerves (in 14 patients, unilaterally) of the 40 nerves (in 20 patients, bilaterally) examined. All abnormalities were found along the course of symptomatic nerves. Blink reflex recordings were obtained in all 40 nerves (20 patients, bilaterally) and were abnormal in 11 symptomatic nerves (11 patients, unilaterally) and normal in nine symptomatic nerves (nine patients, unilaterally); they were also normal in all 20 asymptomatic nerves (20 patients, unilaterally). Masseter inhibitory reflex recordings were obtained in 24 nerves (12 patients, bilaterally) and were abnormal in seven nerves (seven patients, unilaterally) and normal in five symptomatic nerves (five patients, unilaterally); they were also normal in all 12 asymptomatic nerves (12 patients, unilaterally). Jaw-jerk reflex recordings were obtained in 22 nerves (11 patients, bilaterally) and were abnormal in four nerves (four patients, unilaterally) and normal in seven symptomatic nerves (seven patients, unilaterally); they were also normal in all 11 asymptomatic nerves (11 patients, unilaterally).

**MR Imaging Findings**

Eight of the 20 patients with unilateral symptoms related to the trigeminal nerve had an abnormality along the course of the symptomatic trigeminal nerve. Five brain stem lesions were found, including infarcts in the dorsolateral medulla oblongata in three patients (in the midpons in or near the principal trigeminal nucleus in one patient) and hemorrhage in the midpons, which included the principal nucleus in another patient. Three extraaxial lesions were found, including inflammation of the infraorbital nerve in the first patient, inflammation of the trigeminal ganglion extending into the proximal mandibular division in the second patient, and a cerebellopontine angle cistern epidermoid in the third patient. No abnormalities were seen along 12 symptomatic nerves (12 patients, unilaterally) or in any of the 20 asymptomatic nerves (20 patients, unilaterally). Six of the 12 patients with negative MR imaging studies had similar clinical findings after 3 to 36 months (mean, 14 months), and one of these six had a repeat MR imaging examination after 36 months, which was also negative. Four were lost to follow-up, and two had no symptoms, one after 4 months and one after microvascular decompression for trigeminal neuralgia.
Table 1: Clinical, EMG reflex, and MR imaging findings in 20 patients with symptoms related to the trigeminal nerve

<table>
<thead>
<tr>
<th>Patient</th>
<th>Age (y)</th>
<th>Sex</th>
<th>Trigeminal Nerve Symptoms</th>
<th>Other Symptoms</th>
<th>Reflex</th>
<th>Masseter Inhibition</th>
<th>Jaw Jerk</th>
<th>EMG Findings</th>
<th>Location of EMG Findings</th>
<th>MR Findings</th>
<th>Location of MR Findings</th>
<th>Diagnosis/ follow-up*</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>54</td>
<td>F</td>
<td>L pain, sens def (V2–3)</td>
<td>L deafness</td>
<td>Blink</td>
<td>+</td>
<td>+</td>
<td>–</td>
<td>+ L sens trig root</td>
<td>+ L cpa cistern</td>
<td>–</td>
<td>Epidermoid</td>
</tr>
<tr>
<td>2</td>
<td>52</td>
<td>M</td>
<td>L sens def (V1–3)</td>
<td>R body side sens def</td>
<td>+</td>
<td>. . . . . . . .</td>
<td>+</td>
<td>+ L med obl, dorsolat</td>
<td>+ L med obl, dorsolat</td>
<td>Infarction</td>
<td></td>
<td></td>
</tr>
<tr>
<td>3</td>
<td>56</td>
<td>F</td>
<td>L sens def (V2)</td>
<td>None</td>
<td>+</td>
<td>. . . . . . . .</td>
<td>+</td>
<td>L ncl princ, midpons</td>
<td>+ L ncl princ, midpons</td>
<td>Infarction</td>
<td></td>
<td></td>
</tr>
<tr>
<td>4</td>
<td>42</td>
<td>F</td>
<td>R trig neuralgia (V1–3)</td>
<td>R hemifacial spasm</td>
<td>–</td>
<td>. . . . . . . .</td>
<td>–</td>
<td>None</td>
<td>– None</td>
<td>– None</td>
<td>No symptoms at 4 mo</td>
<td></td>
</tr>
<tr>
<td>5</td>
<td>34</td>
<td>M</td>
<td>L sens def (V1–2)</td>
<td>L deafness</td>
<td>–</td>
<td>–</td>
<td>–</td>
<td>–</td>
<td>– None</td>
<td>– None</td>
<td>Lost to follow-up</td>
<td></td>
</tr>
<tr>
<td>6</td>
<td>27</td>
<td>M</td>
<td>R pain, sens def (V2–3)</td>
<td>None</td>
<td>+</td>
<td>+</td>
<td>+</td>
<td>R lat tegm field</td>
<td>– None</td>
<td>– None</td>
<td>No symptoms at 4 mo</td>
<td></td>
</tr>
<tr>
<td>7</td>
<td>50</td>
<td>M</td>
<td>R paresthesias (V2–3)</td>
<td>None</td>
<td>–</td>
<td>–</td>
<td>. . . . .</td>
<td>None</td>
<td>– None</td>
<td>– None</td>
<td>Lost to follow-up</td>
<td></td>
</tr>
<tr>
<td>8</td>
<td>36</td>
<td>F</td>
<td>L paresthesias (V1–2)</td>
<td>None</td>
<td>+</td>
<td>–</td>
<td>+</td>
<td>L pons, rostral</td>
<td>– None</td>
<td>– None</td>
<td>Lost to follow-up</td>
<td></td>
</tr>
<tr>
<td>9</td>
<td>40</td>
<td>M</td>
<td>L paresthesias (V1–3)</td>
<td>None</td>
<td>+</td>
<td>. . . . . . . .</td>
<td>+</td>
<td>R sens trig root</td>
<td>– None</td>
<td>– None</td>
<td>Similar symptoms at 3 y, repeat MR negative</td>
<td></td>
</tr>
<tr>
<td>10</td>
<td>48</td>
<td>M</td>
<td>L sens def (V1–3)</td>
<td>None</td>
<td>–</td>
<td>+</td>
<td>. . . . .</td>
<td>L V3</td>
<td>– None</td>
<td>– None</td>
<td>Similar symptoms at 5 mo</td>
<td></td>
</tr>
<tr>
<td>11</td>
<td>41</td>
<td>M</td>
<td>L sens def (V2–3)</td>
<td>None</td>
<td>–</td>
<td>+</td>
<td>+</td>
<td>L V3</td>
<td>+ L V2–3</td>
<td>Inflammation</td>
<td></td>
<td></td>
</tr>
<tr>
<td>12</td>
<td>42</td>
<td>F</td>
<td>L sens def (V1–3)</td>
<td>L III palsy</td>
<td>–</td>
<td>–</td>
<td>–</td>
<td>None</td>
<td>– None</td>
<td>– None</td>
<td>Similar symptoms at 12 mo</td>
<td></td>
</tr>
<tr>
<td>13</td>
<td>25</td>
<td>F</td>
<td>L pain (V1), sens def (V1–3)</td>
<td>L III, VI, VII palsies</td>
<td>+</td>
<td>+</td>
<td>+</td>
<td>L trig root, prox V3</td>
<td>+ L trig ggl, prox V3</td>
<td>Inflammation</td>
<td></td>
<td></td>
</tr>
<tr>
<td>14</td>
<td>26</td>
<td>M</td>
<td>R pain, sens def (V1)</td>
<td>None</td>
<td>+</td>
<td>. . . . . . . .</td>
<td>+</td>
<td>L midpons, dorsolat</td>
<td>– None</td>
<td>– None</td>
<td>Lost to follow-up</td>
<td></td>
</tr>
<tr>
<td>15</td>
<td>41</td>
<td>F</td>
<td>R paresthesias (V1–3)</td>
<td>None</td>
<td>–</td>
<td>–</td>
<td>–</td>
<td>None</td>
<td>– None</td>
<td>– None</td>
<td>Similar symptoms at 3 mo</td>
<td></td>
</tr>
<tr>
<td>16</td>
<td>29</td>
<td>M</td>
<td>R sens def (V2)</td>
<td>R sens def, paresthesias hand</td>
<td>–</td>
<td>+</td>
<td>–</td>
<td>L lat tegm field, pontomed junction</td>
<td>– None</td>
<td>– None</td>
<td>Similar symptoms at 5 mo</td>
<td></td>
</tr>
<tr>
<td>17</td>
<td>51</td>
<td>F</td>
<td>R sens def (V1), absent CR</td>
<td>L body side sens def</td>
<td>. . . . .</td>
<td>+</td>
<td>. . . . .</td>
<td>R lat tegm field, med obl</td>
<td>+ R med obl</td>
<td>Infarction</td>
<td></td>
<td></td>
</tr>
<tr>
<td>18</td>
<td>63</td>
<td>F</td>
<td>R sens def (V1–3)</td>
<td>L body side sens def</td>
<td>. . . . .</td>
<td>+</td>
<td>. . . . .</td>
<td>R med obl, dorsolat</td>
<td>+ R med obl, dorsolat</td>
<td>Infarction</td>
<td></td>
<td></td>
</tr>
<tr>
<td>19</td>
<td>49</td>
<td>M</td>
<td>R sens def (V1–2)</td>
<td>R VI palsy, sens def L hand</td>
<td>. . . . .</td>
<td>+</td>
<td>. . . . .</td>
<td>R ncl princ, midpons</td>
<td>+ R pons</td>
<td>Hemorrhage</td>
<td></td>
<td></td>
</tr>
<tr>
<td>20</td>
<td>58</td>
<td>M</td>
<td>R sens def (V2–3)</td>
<td>None</td>
<td>–</td>
<td>. . . . . . . .</td>
<td>–</td>
<td>None</td>
<td>– None</td>
<td>– None</td>
<td>Similar symptoms at 20 mo</td>
<td></td>
</tr>
</tbody>
</table>

Note.—Sens def indicates sensory deficit; III, oculomotor nerve; V1, V2, V3, first (V1), second (V2), and third (V3) division of trigeminal nerve; VI, abducens nerve; VII, facial nerve; . . . , not performed; +, abnormal; –, normal; med obl, medulla oblongata; ncl, nucleus; cpa, cerebellopontine angle. Only the EMG and MR imaging findings of the 20 symptomatic trigeminal nerves are shown, the EMG and MR imaging findings of all 20 asymptomatic nerves were normal.

*Follow-up only for patients with negative MR imaging findings.
Table 2: EMG versus MR imaging results in 40 trigeminal nerves (20 symptomatic)

<table>
<thead>
<tr>
<th>MR Imaging</th>
<th>+</th>
<th>-</th>
<th>Total</th>
</tr>
</thead>
<tbody>
<tr>
<td>EMG</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>+</td>
<td>8</td>
<td>6</td>
<td>14</td>
</tr>
<tr>
<td>-</td>
<td>0</td>
<td>26</td>
<td>26</td>
</tr>
<tr>
<td>Total</td>
<td>8</td>
<td>32</td>
<td>40</td>
</tr>
</tbody>
</table>

+ indicate abnormal; −, normal. Sensitivity, specificity, negative and positive predictive value, and accuracy of EMG in predicting MR imaging results were as follows: sensitivity, 100% (8/8) (95% CI: 63%–100%); specificity, 81% (26/32) (95% CI: 64%–94%); negative predictive value, 100% (26/26) (95% CI: 87%–100%); positive predictive value, 57% (8/14) (95% CI: 29%–82%); accuracy, 85% (34/40) (95% CI: 70%–94%).

EMG/MR Imaging Correlation

The results of the EMG/MR imaging correlation of the 40 trigeminal nerves examined are summarized in Table 2.

There were 14 positive EMG studies, corresponding to eight positive and six negative MR imaging studies. Of the eight patients with positive EMG and MR imaging studies, five had brain stem lesions. In these five patients, lesion localization, as demonstrated with EMG, corresponded well with lesion localization as shown by MR imaging (Figs 4 and 5). Three of these eight patients had extraaxial lesions; EMG correctly identified these lesions, but could not exactly localize the abnormalities as MR imaging did (Fig 6). In six patients, EMG was positive and MR imaging was negative. At clinical follow-up, three of these six patients had similar symptoms after 5, 12, and 36 months, respectively. One had a repeat MR imaging study, which was also negative; the other three were lost to follow-up. None of the trigeminal nerves that produced a negative EMG study showed a positive MR imaging result. As compared with MR imaging, EMG had a sensitivity of 100% (95% confidence interval [CI]: 63% to 100%) and specificity, 81% (95% CI: 64% to 93%); a positive predictive value of 57% (95% CI: 29% to 82%); and a negative predictive value of 100% (95% CI: 87% to 100%) (Table 2).

There was a high degree of agreement between the observers with regard to the presence of an appropriate lesion at EMG (κ = .94) and at MR imaging (κ = .92).

Discussion

Previous authors have shown that clinical localization of a trigeminal nerve lesion is poor as compared with MR imaging results and that the entire course of the trigeminal nerve should always be visualized (4). The results of our study indicate that electrodiagnostic testing of the trigeminal reflexes can be used for improved lesion localization. EMG of multiple trigeminal nerve reflexes can accurately localize lesions in the brain stem because the trigeminal reflex circuits are located at different brain stem levels—mesencephalon (jaw-jerk reflex); pons (blink reflex-R1, masseter inhibitory reflex-SP1); pontomedullary junction (masseter inhibitory reflex-SP2); and lower medulla (blink reflex-R2)—enabling accurate assessment of these regions (5, 11, 12, 15, 16, 18, 20).

In six of our patients, EMG reflex studies were positive and MR imaging studies were negative. Very small lesions or impaired physiologic functions may be found with trigeminal reflex testing,
even though structural abnormalities cannot be depicted with MR imaging (20, 21). This may explain the relatively high number of positive (6/14) EMG findings relative to MR imaging results in our study. We postulate that in some of these patients the lesions were too small to be seen with MR imaging. Probable causes for the trigeminal nerve deficits in these patients are microinfarctions in the brain stem, ischemia of the nerve itself, viral inflammation, and autoimmune disorders (10, 22, 23).

Previous investigators have noted that EMG can distinguish between idiopathic trigeminal neuralgia (ie, neurovascular) and symptomatic trigeminal neuralgia (ie, due to a structural cause, such as neoplasms or multiple sclerosis) (8, 24). Cruccu et al (24) recorded the trigeminal reflexes, including blink, masseter inhibitory, and jaw-jerk, in 30 patients with idiopathic trigeminal neuralgia and in 20 patients with symptomatic trigeminal pain. Of the 30 patients with idiopathic trigeminal neuralgia, only two showed slight delays of short-latency reflexes. In the other 28 cases, the trigeminal reflexes were completely normal. These results correspond well with the negative EMG studies in the patient with trigeminal neuralgia in our study. In the study by Cruccu and coworkers, all the patients with symptomatic trigeminal pain had trigeminal reflex abnormalities (24).

The results of our study indicate that EMG reflex studies can play a role in the preselection of patients with trigeminal nerve dysfunction who are referred for MR imaging. The high negative predictive value of EMG suggests that if a normal EMG result is obtained, it is unlikely that MR imaging will show a structural cause for the symptoms and is therefore not required for this purpose. Nonetheless, the high sensitivity and negative predictive value of EMG in predicting MR imaging results in our study may be due to selection bias of the patients in this retrospective study. The prevalence of positive MR imaging studies was relatively low in our series (abnormalities found in 40% of all patients and in 20% of all nerves examined) as compared with a previous study of 112 consecutive patients who underwent MR imaging for symptoms and signs related to the trigeminal nerve (6). In that study, abnormalities along the trigeminal nerve were found in 68 patients (61%) (6). Furthermore, to compare EMG reflex and MR imaging studies of the trigeminal nerve in a blinded manner, the trigeminal nerves of the clinically normal sides were also included in our study, which contributed to the low prevalence.

To our knowledge, assessment of interobserver variability of EMG reflex studies of the trigeminal nerve has not been performed previously. The good interobserver agreement for MR imaging we obtained corresponds to the results of a previous study (6). The high $\kappa$ values for both the EMG reflex studies and the MR imaging examinations indicate that these are both robust tests for evaluating trigeminal nerve lesions.

Conclusion

Our data suggest that EMG of the trigeminal reflexes can be used to exclude structural lesions and to localize accurately lesions of the trigeminal system in the brain stem. When a lesion is localized in the brain stem with EMG reflex studies, a more tailored MR examination of this region might be sufficient. We recognize, however, that our study represents preliminary work and that a large prospective study is warranted to validate these conclusions.
References