Traumatic trigeminal neuropathy after whiplash injury
A case report
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Abstract

\textbf{Rationale:} Many studies using diffusion tensor tractography (DTT) have reported trigeminal neuropathy in various neurological diseases. However, no study on traumatic trigeminal neuropathy following whiplash has been reported.

\textbf{Patient concerns:} A 51-year old female suffered an indirect head trauma resulting from a flexion-hyperextension injury. At approximately 30 minutes after onset, she began to sense a headache in the left frontal area and sensory changes in the left facial area, signs that intensified with the passage of time. At 7 days after onset, she visited the rehabilitation department of our university hospital and described the characteristics and severity of pain as follows: headache on the left frontal area including the forehead with intermittent squeezing and numbness sensations. Her visual analog scale pain score was 6 with her left cheek having a continuous, dull, swelling sensation (visual analog scale score: 1). On neurological examination, she revealed mild allodynia without hyperalgesia or somatosensory change on the head, cheek, tongue, and oral cavity.

\textbf{Diagnosis:} Diffusion tensor imaging data were acquired 7 days after onset. On DTT, the left trigeminal nerve showed discontinuation in the middle portion compared to that of the right trigeminal nerve. Traumatic trigeminal neuropathy was diagnosed based on her clinical features and DTT findings.

\textbf{Intervention:} She was prescribed carbamazepine (200 mg/day) and pregabalin (150 mg/day), and her facial pain was well-controlled to a tolerable level.

\textbf{Outcomes:} These drugs were stopped after approximately 7 month’s administration, however, she did not complain of facial pain.

\textbf{Lessons:} By using DTT, we demonstrated traumatic trigeminal neuropathy in a patient with whiplash. We suggest that DTT would be a useful tool for detection of traumatic trigeminal neuropathy in patients who show clinical features of trigeminal neuropathy following whiplash.

\textbf{Abbreviations:} DTI = diffusion tensor imaging, DTT = diffusion tensor tractography, MRI = magnetic resonance imaging.

\textbf{Keywords:} diffusion tensor tractography, head trauma, traumatic trigeminal neuropathy, trigeminal nerve, whiplash

1. Introduction

Whiplash is a bony and/or soft tissue injury resulting from acceleration–deceleration energy transfers in the neck.\cite{1} Recently, several studies have demonstrated brain injury following whiplash.\cite{2} Many studies have reported clinical evidence of facial symptoms which suggests the presence of trigeminal neuropathy after whiplash.\cite{3,4,5} However, because conventional brain magnetic resonance imaging (MRI) is limited in its capability to demonstrate trigeminal neuropathy, trigeminal neuropathy after whiplash could not be clearly diagnosed. By contrast, diffusion tensor tractography (DTT) which is a derivation of diffusion tensor imaging (DTI), allows 3-dimensional reconstruction and estimation of the trigeminal nerve.\cite{10} Many previous studies using DTT have demonstrated trigeminal neuropathy in various neurological diseases, particularly trigeminal neuralgia.\cite{11,12,13,14,15,16,17,18,19,20} However, no study on traumatic trigeminal neuropathy following whiplash has been reported.

In this study, we report on a whiplash patient who showed traumatic trigeminal neuropathy, which was demonstrated on DTT.

2. Case report

A 51-year old female with no history of neurological, physical, or psychiatric illness suffered an indirect head trauma resulting from
a flexion-hyperextension injury after being hit from behind by a moving vehicle while stopping at an intersection. At the time of the head trauma, she did not experience loss of consciousness or posttraumatic amnesia, and her Glasgow Coma Scale was 15. Approximately 30 minutes after onset, she began to sense a headache in the left frontal area and experience sensory changes in the left facial area, signs that increased with the passage of time. At 7 days after onset, she visited the rehabilitation department of our university hospital, the characteristics and severity of her pain were as follows: headache in the left frontal area including the forehead; a spontaneous intermittent squeezing and numbness sensation (visual analog scale score: 6) with a spontaneous, continuous, dull, swelling sensation in the left cheek (visual analog scale score: 1). On neurological examination, she revealed mild allodynia without hyperalgesia or somatosensory change on the head, cheek, tongue, and oral cavity. In addition, she did not show weakness of the left masseter and temporalis muscles. Conventional brain MRI and DTI were recommended for the patient. Conventional brain MRI including T1-weighted, T2-weighted, and fluid-attenuated inversion recovery images, obtained at 7 days after onset, showed no abnormality (Fig. 1A). She was prescribed carbamazepine (200 mg/day) and pregabalin (150 mg/day), and her facial pain was well-controlled to a tolerable level. Her facial pain was gradually relieved with the passage of time. These drugs were stopped after approximately 7 month’s administration, however, she did not complain of facial pain. The patient provided written and informed consent, and the study protocol was approved by the institutional review board of our university hospital.

Figure 1. (A) T2-weighted brain magnetic resonance (MR) images obtained 7 days after whiplash onset show no abnormality (upper row). Results of diffusion tensor tractography: the left trigeminal nerve is discontinued (arrow) (lower row). (B) Results of diffusion tensor tractography for a control subject (53-year old female).
The DTI data were acquired 7 days after onset by using a sensitivity-encoding head coil on a 1.5 T Philips Gyroscan Intera (Hoffman-LaRoche, Best, Netherlands) with single-shot echo-planar imaging and a navigator echo. For each of the 32 non-collinear diffusion-sensitizing gradients, 67 contiguous slices (acquisition matrix = \(96 \times 96\); reconstructed to matrix = \(192 \times 192\) matrix; field of view = \(240 \times 240\) mm; repetition time = 10,398 ms; time echo = 72 ms; parallel imaging reduction factor (ENSE factor) = 2; echo planar imaging factor = 59; \(b = 1000 \times \) \(\text{mm}^2\)) were acquired parallel to the anterior commissure-posterior commissure line. Eddy current-induced image distortions were removed by using affine multi-scale 2-dimensional registration as included in the Oxford Centre for the Functional Magnetic Resonance Imaging of the Brain software library (www.fmrib.ox.ac.uk/fsl).[211] DTI-Studio software (CMRM, Johns Hopkins Medical Institute, Baltimore, MD) was used for evaluation of the trigeminal nerve. For nerve fiber delineation, the seed region of interest was placed on the preopticine cistern and the target region of interest was placed on isolated distal branches. Fiber tracking was performed by applying a fractional anisotropy threshold of >0.1 and a direction threshold <70°.[10] On DTI, the left trigeminal nerve showed discontinuation in the middle portion compared to that of the right trigeminal nerve (Fig. 1).

3. Discussion

In this study, we detected a discontinuation of the left trigeminal nerve, an observation that was in accord with the trigeminal neuropathy symptoms exhibited by our patient with whiplash. The patient showed clinical features that satisfied the definitive criteria for presence of peripheral painful traumatic trigeminal neuropathy (i.e., A, spontaneous pain; B, develops within 3 months of an identifiable traumatic event; C, allodynia present; D, imaging or neurophysiology results demonstrate a neurologic lesion and its location; E, not attributed to other disorders). As a result, the traumatic trigeminal neuropathy signs appear to coincide with the DTI results showing a neuropathy of the left trigeminal nerve.[22]

Several studies have reported patients who presented facial pain after whiplash which appeared to be related to traumatic trigeminal neuropathy.[3,5,6,8] In 1988, McGlone et al[3] reported on a patient who showed delayed onset of facial pain after a whiplash. In 2011, Haggman-Henrikson et al[5] found that 88% of whiplash patients complained of the face and jaw pain which were examined by questionnaire for pain. Subsequently, Genesu[6] reported on a patient who showed the right facial numbness and cheek pain after a whiplash. The patient's facial symptoms were alleviated by treating the right-sided strain of the trigeminal nerve.[3,5,6,8] In 2011, Samim and Epstein[9] reported on a patient who complained of shooting pain in the jaw, cheek, and forehead beginning 7 days after a whiplash. However, these previous studies could not demonstrate trigeminal neuropathy because the authors suspected trigeminal neuropathy based on questionnaire or neurological examination without radiologic evidences. Since introduction of DTI, many DTI-based studies have demonstrated trigeminal neuropathy in various neurologi- cal diseases including classical trigeminal neuralgia, brain tumor, neurovascular compression, and multiple sclerosis.[11–20] As a result, to the best of our knowledge, this is the first case study to report on the presence of traumatic trigeminal neuropathy following whiplash. However, the limitation of this study regarding DTI should be considered; DTI can produce both false positive and false negative results throughout the white matter of the brain due to complex fiber configurations such as crossing or kissing fibers and/or partial volume effects.[2,3]

In conclusion, by using DTI, we demonstrated the presence of traumatic trigeminal neuropathy in a patient with whiplash. We suggest that DTI can be a useful tool for detection of traumatic trigeminal neuropathy in patients who show clinical features of trigeminal neuropathy following a whiplash.

Author contributions

Sung Ho Jang, study concept and design, manuscript development, writing, data acquisition, and drafting/revising the image. Jeong Pyo Seo, study support, data acquisition, and drafting/revising the image. Younghyeon Kwon, critical revision of manuscript for intellectual content, drafting/revising the image, writing, and funding.

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