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Case Report

Magnetic Resonance Imaging of Kernohan's Notch in Chronic Subdural Hematoma: Significance of Coronal Images for Preoperative Diagnosis

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A 67-year-old man presented with headache and gait disturbance. On admission, he was mildly confused but aroused by verbal stimulus, with normal motor function. A CT scan showed bilateral subdural hematomas (right >> left) and a midline shift to the left. One hour after admission, he suddenly became somnolent and developed right hemiparesis. While repeated CT examinations failed to reveal new findings, coronal MRI clearly depicted the left cerebral peduncle pressed against the free edge of the tentorium. Craniotomy was immediately performed to remove the right chronic subdural hematoma. Soon after the operation, neurological functions were markedly improved.

A 67-year-old man was admitted to our hospital on 31 May 2000, with headache and difficulty of walking. On admission, the patient was mildly confused, rating a Glasgow Coma Scale (GCS) of 14/15 (E3V5M6). Pupils were isocoric and no definite hemiparesis was evident. The initial CT scan revealed bilateral subdural hematoma with a 10-mm-shift of the septum pellucidum to the left. The right hematoma was 15-mm-thick and appeared as a two-layer pattern of subdural collection (figure 1 A). The patient was hospitalized the same day for surgery. One hour after admission, the patient suddenly developed drowsiness with a GCS of 8/15 (E2V1M5) and right (ipsilateral to the lesion) hemiparesis. Pupils became slightly anisocoric (right > left). Repeated CT studies failed to determine any new findings. Therefore, we postulated that Kernohan's notch might play a role in this paradoxical motor deficit. To confirm this hypothesis and to exclude the possibility of cerebral ischemia, MRI and MRA studies were immediately performed. Coronal MRI clearly demonstrated the left cerebral peduncle pressed against the free edge of the tentorium (figure 1 B). The signal patterns observed suggested the recurrence of hemorrhaging in the subdural hematomas. No significant ischemic lesions were observed in MR studies, including diffusion-weighted images and MRA. Directly after MR studies, a craniotomy was performed to remove the right subdural hematoma, which was a typical chronic hematoma with outer and inner membranes. Following surgery,

Key Words: Kernohan's notch, chronic subdural hematoma, magnetic resonance image

Introduction

Due to a supratentorial mass, the contralateral cerebral peduncle is occasionally pressed against the free edge of the tentorium, resulting in paradoxical (ipsilateral to the lesion) hemiparesis. While this phenomenon is well known as Kernohan's notch (KN), few radiological examinations have been reported. Preoperative conventional magnetic resonance imaging (MRI) studies are particularly difficult to perform, because KN is usually only seen in emergency cases with severe head injury, such as large acute epidural hematomas (AEDH) or acute subdural hematomas (ASDH). KN is relatively rare to observe in cases of chronic subdural hematomas (CSDH). In this study, we report on an unusual case of KN secondary to bilateral CSDH, distinctly identified by preoperative coronal MRI.

Case report

A 67-year-old man was admitted to our hospital on 31 May 2000, with headache and difficulty of walking. On admission, the patient was mildly confused, rating a Glasgow Coma Scale (GCS) of 14/15 (E3V5M6). Pupils were isocoric and no definite hemiparesis was evident. The initial CT scan revealed bilateral subdural hematoma with a 10-mm-shift of the septum pellucidum to the left. The right hematoma was 15-mm-thick and appeared as a two-layer pattern of subdural collection (figure 1 A). The patient was hospitalized the same day for surgery. One hour after admission, the patient suddenly developed drowsiness with a GCS of 8/15 (E2V1M5) and right (ipsilateral to the lesion) hemiparesis. Pupils became slightly anisocoric (right > left). Repeated CT studies failed to determine any new findings. Therefore, we postulated that Kernohan's notch might play a role in this paradoxical motor deficit. To confirm this hypothesis and to exclude the possibility of cerebral ischemia, MRI and MRA studies were immediately performed. Coronal MRI clearly demonstrated the left cerebral peduncle pressed against the free edge of the tentorium (figure 1 B). The signal patterns observed suggested the recurrence of hemorrhaging in the subdural hematomas. No significant ischemic lesions were observed in MR studies, including diffusion-weighted images and MRA. Directly after MR studies, a craniotomy was performed to remove the right subdural hematoma, which was a typical chronic hematoma with outer and inner membranes. Following surgery,
neurological functions improved considerably, and after 25 days the patient was discharged with no significant motor deficit. A postoperative MRI examination revealed marked improvement in the deformity of the left cerebral peduncle (figure 1 C).

Figure 1 A. CT scans on admission, shows a large subdural hematoma in the right frontotemporoparietal area with a midline shift to the left. A small subdural hematoma is also observed on the left.

Figure 1 B. A T2-weighted coronal MR image 1 hour after admission, clearly shows the compressed lateral part of the left brain stem by tentorial edge (arrow).

Figure 1 C. Three months after surgery coronal T2WI reveals improvement in the deformity of the left cerebral peduncle and no T2-elongated lesion in the left brainstem.

Discussion

Kernohan’s notch (KN) has been well known as a possible cause of paradoxical (ipsilateral to the lesion) motor deficits usually seen in patients with severe head injury. Few MRI studies of these lesions have been reported, probably due to the serious condition of the patients and rapid deterioration of the clinical course. We were able to find ten reports of KN demonstrated by MR imaging (Table 1), including two cases of CSDH. In most cases, postoperative axial images revealed T2 elongated lesions in the contralateral cerebral peduncle. Itoyama et al. described a deformity of the cerebral peduncle in preoperative T1-weighted axial MR images. In the present case, however, axial images would not have provided sufficient diagnostic information. While several reports emphasized the significance of coronal MR images for detecting KN, their imaging data were postoperative.

Table 1. Summary of MRI findings in reported cases of Kernohan’s notch

<table>
<thead>
<tr>
<th>Author</th>
<th>Sex/Age</th>
<th>Diagnosis</th>
<th>MR Images</th>
</tr>
</thead>
<tbody>
<tr>
<td>1990 Cohen</td>
<td>M/19</td>
<td>ASDH</td>
<td>T2 hyperintensity</td>
</tr>
<tr>
<td>1991 Jones</td>
<td>M/25</td>
<td>ASDH</td>
<td>T1 hypointensity, T2 hyperintensity</td>
</tr>
<tr>
<td>1992 Iwama</td>
<td>M/39</td>
<td>ASDH</td>
<td>T1 hypointensity, T2 hyperintensity</td>
</tr>
<tr>
<td>1993 Itoyama</td>
<td>M/49</td>
<td>CSDH</td>
<td>deformity of the crus cerebri</td>
</tr>
<tr>
<td>1997 Yamazaki</td>
<td>M/43</td>
<td>ASDH</td>
<td>T2 hyperintensity</td>
</tr>
<tr>
<td>1997 Ninomiya</td>
<td>F/16</td>
<td>ASDH</td>
<td>T2 hyperintensity</td>
</tr>
<tr>
<td>1997 Zafonte</td>
<td>M/42</td>
<td>ASDH</td>
<td>T1 hypointensity</td>
</tr>
<tr>
<td>1998 Murotsu</td>
<td>M/16</td>
<td>ASDH</td>
<td>T1 hypointensity</td>
</tr>
<tr>
<td>1998 Takahara</td>
<td>F/27</td>
<td>ASDH</td>
<td>T1 hypointensity, T2 hyperintensity</td>
</tr>
<tr>
<td>1999 Mastronardi</td>
<td>F/73</td>
<td>ASDH</td>
<td>T1 hypointensity, T2 hyperintensity</td>
</tr>
</tbody>
</table>

M: male, F: female, ASDH: acute subdural hematoma, CSDH: chronic subdural hematoma
to explain the mechanism of paradoxical motor deficit. To our knowledge, this report is the first demonstration of preoperative coronal MR images for diagnosing KN. In the left cerebral peduncle, no pathological signals were observed even in the diffusion-weighted images, suggesting reversibility of the lesion as previously reported. In present case study, sudden deterioration in neurological status first led us to suggest ischemic complications in the contralateral brain, because there were no significant changes in the hematoma and a midline shift appeared on repeated CT scans. Counter pressure from the contralateral CSDH might produce a minimal shift and higher ICP than typically expected. As a result, relatively little renewed hemorrhaging may cause KN in the absence of definite changes in CT images.

In such cases, serial examinations by MRI and MRA are necessary, and should be immediately performed in order to determine the next option of the treatment, such as thrombolysis or surgical removal of the hematoma.

In general, neurological symptoms in CSDH gradually progress over several weeks following head injuries. However, Kotwica and Brzezinski reported cases of sudden onset and rapid deterioration suggesting cerebral stroke, observed in 14 (11%) of 131 adult patients with CSDH. Further, in 9 (7%) patients, CSDH led to transtentorial herniation within a very short period. Of particular interest, 10% of their patients presented focal signs ipsilateral to CSDH. These observations suggest that atypical clinical pictures of CSDH as encountered in this case are not as uncommon as expected, and should always be taken into consideration in the management of CSDH.

References

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