Cervicomedullary Hematoma: Diagnosis by MR

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CT is often useful for diagnosing primary or secondary brainstem hematoma. However, subacute hemorrhage may be isodense with the normal brain substance, causing a false negative CT study. MR tends to be positive during this phase of hemorrhage. We present an unusual case of an intramedullary hematoma that was not detected by CT but was detected by MR. MR has shown great promise as a neuroradiologic tool in difficult situations by virtue of its excellent tissue-contrast resolution, a lack of bone artifacts, and direct multiplanar imaging capability [1]. The lack of signal from bone has the advantage that neighboring brain structures, particularly those in the posterior fossa, are not obscured by the linear bone artifacts common to CT scans [2].

Primary hemorraghes of the brainstem commonly involve the pons, less frequently the cerebral peduncle, and rarely, the medulla oblongata [3]. Primary bulbar hemorrhage is usually first identified at autopsy. To our knowledge, there is no known reported case of cervicomedullary hematoma in which antemortem diagnosis was established by any imaging technique. In our case, CT failed to show the lesion but MR clearly identified the hematoma. Timely surgical intervention with evacuation of the hematoma resulted in marked postoperative functional recovery.

Case Report

A 21-year-old right-handed woman was admitted with persistent hiccups, nausea, headache, and generalized weakness of 5 days duration and difficulty in swallowing for the past 2 days. She denied any fever and vomiting.

Shortly after admission, a lumbar puncture revealed 20 WBC/high-power field with 60% polyps and 40% lymphs, and 7500 RBC/high-power field. The patient was empirically treated for tuberculosis. A second lumbar puncture was normal. Electroencephalography revealed background slowing (6-7 Hz) with intermittent frontotemporal delta wave-burst activity. CT brain scans on the day of admission (not shown) were normal; however, the lowest cut was at the level of the lower pons. Bilateral vertebral angiography was normal (Fig. 1). Within 36 hr, the patient became unresponsive and quadriplegic. Generalized tonic clonic seizures developed. Respiratory embarrassment led to tracheostomy. An MR study was performed (Fig. 2), and T1-weighted short spin-echo images (TR=500 msec, TE=30 msec) showed a very high signal mass within the central medulla consistent with subacute hemorrhage.

A repeat CT examination of the posterior fossa and the upper cervical spine (Fig. 3) revealed slight dilatation of the cervicomedullary junction. The night before surgery, the patient developed a fever of 104°F, so the surgery was postponed. Blood cultures showed staphylococcus epidermidis, and she was started on oxacillin. She was still drowsy and quadriparetic. A portable chest X-ray showed pneumonitis in the left lung base. After slight improvement in her general condition, the patient was rescheduled for surgery.

Under general anesthesia, a suboccipital craniotomy and C1–C2 laminectomy were done. The cervicomedullary junction was discolored and bulging. With the aid of an operating microscope, a small myelotomy was made in the right posterior column. Brownish smoky fluid (subacute hemorrhage) was released. The dura was closed with a graft. The patient quickly awakened and began moving all extremities spontaneously. A repeat bilateral vertebral angiogram was normal. She was transferred to a rehabilitation unit, where she showed continued improvement. She is currently ambulatory without assistance.

Discussion

Hematomyelia is defined as a focal extravasation of blood within the substance of the spinal cord, resulting in a nonextending, expansile, limited saucular clot [4]. Hematomyelia can be primary or secondary. The cause of primary hematomyelia is either unknown or due to capillary telangiectasia. The most common cause of secondary hematomyelia is vascular malformation. Other predisposing conditions include polomyelitis, toxic states, blood dyscrasias (especially when associated with purpura), asphyxia as a sequelae of severe convulsions, and spinal cord injury [5].

A similar classification can be applied to brainstem hematoma, but no such grouping is clearly stated in the literature. Brainstem hematomas can be defined as subependymal or intraaxial. Intraaxial hematomas are not clinically distinguishable from intrinsic brainstem tumors, which are generally not considered suitable for surgery. Primary spontaneous hematomas constitute 13% of the cases in one series [6]. The
rostral parts of the brainstem are mostly affected because vascular malformations are more frequent in this region [7]. Medullary hematomas are extremely rare and the pathogenic mechanisms of primary hematomas are not clear [8].

We do not have complete pathologic proof of a bleeding source, but it seems likely that the hematoma was due to the rupture of a microaneurysm of one of the penetrating branches of the basilar or vertebral arteries [7] as suggested by Cole and Yates [9] (though they described a pontomedullary hematoma).

The pathologic examination did not reveal any capillary telangiectasia in our patient. A similar case is described by Donald and Lysia [10].

MR is clearly superior to CT for the identification of subacute hemorrhage. At the time that blood clot is isodense to brain on CT, T1-weighted MR images show very high signal. The only other tissue exhibiting such characteristics on MR is fat. Lipoma is not of serious consideration in the presence of acute intramedullary brainstem expansion, as in this case. MR has also been shown to be superior to CT for the detection of subacute subdural hematoma [11], although acute hemorrhages, which are dense on CT studies, may be less obvious on MR [11].

While the exact figures are unknown, there are isolated case reports with good postoperative results after the evacuation of intramedullary hematoma [7, 12]. An early diagnosis of hematomyelia with successful postoperative results are obtained in six of nine cases in one series [13].
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REFERENCES

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