Hepatocellular carcinoma metastasizing to the skull base involving multiple cranial nerves

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Abstract

We describe a rare case of HCV-related recurrent multiple hepatocellular carcinoma (HCC) metastasizing to the skull base involving multiple cranial nerves in a 50-year-old woman. The patient presented with symptoms of ptosis, fixation of the right eyeball, and left abducens palsy, indicating disturbances of the right oculomotor and trochlear nerves and bilateral abducens nerves. Brain contrast-enhanced computed tomography (CT) revealed an ill-defined mass with abnormal enhancement around the sella turcica. Brain magnetic resonance imaging (MRI) disclosed that the mass involved the clivus, cavernous sinus, and petrous apex. On contrast-enhanced MRI with gadolinium-chelated contrast medium, the mass showed inhomogeneous intermediate enhancement. The diagnosis of metastatic HCC to the skull base was made on the basis of neurological findings and imaging studies including CT and MRI, without histological examinations. Further studies may provide insights into various methods for diagnosing HCC metastasizing to the craniospinal area.

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Key words: Hepatocellular carcinoma; Distant metastasis; Skull base; Cranial nerve
0.04 kAU/L) for protein induced by vitamin K absence (PIVKA II). Incremental dynamic computed tomography (CT) revealed multiple HCC nodules in both lobes (Figure 1). Chest X-ray showed extrahepatic metastases to both lungs; however, imaging studies did not show metastases to the bone or adrenal gland. Brain contrast-enhanced CT (Figure 2) revealed an ill-defined mass with abnormal enhancement around the sella turcica. On unenhanced brain magnetic resonance imaging (MRI) the mass was shown to involve the clivus, cavernous sinus, and petrous apex. Inhomogeneous intermediate enhancement of the mass was confirmed by contrast-enhanced MRI with gadolinium-chelated contrast medium. Meningeal thickening from the posterior of the sella turcica to the clivus and dilation of the superior carotid vein were observed (Figure 3).

**DISCUSSION**

The third, fourth, and sixth cranial nerves may be involved in lesions affecting their nuclei or fibers of efferent in the pons or the mesencephalon, or in the course of nerves through the subarachnoid space, the cavernous sinus, and the superior orbital fissure. Different syndromes occur with lesions at each of these sites. From neurological findings such as ptosis, fixation of the right eyeball, and left abducens palsy, we suspected the involvement of the multiple cranial nerves. Brain CT and MRI with contrast medium revealed abnormal enhancement around the sella turcica, and brain MRI revealed that the mass involved the clivus, cavernous sinus, and petrous apex, therefore we made the definite diagnosis of HCC metastasizing to the
Metastasis appears to be a late event in the natural history of HCC, and most patients die of liver failure attributed to liver replacement by the tumor. Extrahepatic metastases are commonly found at autopsy in over half of the cases, the lung, being the most frequent (approximately 50%), and the regional lymph nodes often being the sites of metastasis. In addition, bone metastases are common, and patients may show a bony metastasis as the initial presentation of occult HCC. Adrenal metastases are found in up to 15% of autopsies, and some cases of pedunculated HCC have been reported as actual metastasis to the right adrenal gland from a nearby tumor. Conversely, the brain is less common metastatic sites. Incidental extrahepatic lesion at such more uncommon site is less confidently viewed as potential metastases when metastatic disease is not detected at the more common sites (the lungs, lymph nodes, and bone). According to neurological findings and imaging studies of brain CT and MRI, our case was confirmed as metastatic HCC to the skull base involving multiple cranial nerves including the third, fourth, and sixth.

The central nervous system is an unusual site of metastatic HCC. Seven cases of HCC presenting as cranial metastasis without obvious hepatic involvement have been described: metastatic spread of HCC to the cerebrum in one case, and to the cranium in six cases, all manifesting mildly abnormal liver function when first evaluated, leading to the conclusion that in cranial metastasis of unknown origin in a geographical area where HCC is a common disease, HCC should be considered in the differential diagnosis. In Japan as in Taiwan, where HCC is a common disease, however, metastatic HCC to the skull base involving multiple cranial nerves has not been reported so far, except for a case of skull metastasis of HCC associated with acute epidural hematoma.

Before 1990, the diagnosis of metastatic HCC to the craniospinal area was confirmed by histological examinations of biopsy, surgical and autopsy specimens. Recently diagnosis is reached by neurological findings and imaging studies, such as CT and MRI because of improvements in these modalities. With the recent advances in the nonsurgical treatment of HCC, such as percutaneous ablation including percutaneous ethanol injection therapy (PEITT), radiofrequency ablation (RFA) and transcatheter embolization, longer-term survival has been attained, even in cases of advanced HCC compared with older autopsy populations. This may be due to current treatment regimens of chemotherapy or chemoembolization, or both, that result in the longer survival of patients diagnosed with HCC, as was in our case. Although HCC metastasizing to the skull base involving multiple cranial nerves is very rare, clinicians need to be vigilant and would do well to conduct imaging studies such as CT and MRI, when neurological findings reveal suspected cranial nerve involvement.

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