Cerebral haemorrhagic infarction associated with acute otitis media in a 4-year-old boy

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DESCRIPTION

A 4-year-old boy had cough and purulent nasal discharge 2 days before admission; he also had fever, headache, vomiting and left ear pain. He received amoxicillin (40 mg/kg/day) and was diagnosed with left acute suppurative otitis media (AOM) by an otolaryngologist, the same day. The next day, he developed right clonus of the upper and lower limbs and was administered midazolam. He was immediately referred to our hospital; he was sedated at the time of admission due to midazolam. Physical examination revealed redness on the left eardrum and pus in the left ear canal, without a heart murmur, stiff neck or Kernig’s sign. Blood examination showed a white cell count of 12.2 × 10⁹/L, C-reactive protein level of 0.32 mg/mL, prothrombin time international normalised ratio of 1.11, activated partial thromboplastin time of 32.3 s, fibrinogen level of 524 mg/dL and D-dimer level of 0.6 µg/mL. Cerebrospinal fluid (CSF) examination showed a CSF pressure of 50 mm Hg, a cell count of 21 cells/µL, glucose level of 99 mg/dL (CSF/serum glucose ratio of 0.77) and protein level of 29.7 mg/dL. Brain diffusion-weighted and T2-star-weighted MRI showed two lesions of haemorrhagic infarctions in the left hemisphere (figure 1A–C). Brain T2-weighted MRI showed a high signal in the mastoid part and tympanic cavity (figure 1D). Brain magnetic resonance angiography (MRA) showed no abnormal findings in the cerebral artery. Head CT showed no anatomical abnormalities indicating direct central nervous system spread, including bone destruction. Echocardiography showed no signs of underlying cardiogenic-embolism-causing diseases, including infectious endocarditis. He was administered cefotaxime, ampicillin and corticosteroid. Blood and cerebrospinal cultures at admission were negative. Haemophilus influenzae was detected via CSF PCR. Transient aphasia, right facial nerve paralysis and right hemiplegia were observed. Although he had difficulty moving his right hand, he was discharged independently.

AOM can occasionally cause cerebral complications, including bacterial meningitis, brain abscesses and lateral sinus thrombosis.⁵ Haemorrhagic infarction caused by AOM is extremely rare. Massive ischaemic stroke after cerebral artery infarction caused by AOM has been reported.⁵ In the case, haemorrhagic infarctions were not considered arterial infarctions because MRA showed no abnormal findings and haemorrhagic infarctions were observed in different areas of the artery. Therefore, haemorrhagic infarctions were believed to be caused by venous infarctions. Cerebral venous infarction due to AOM is extremely rare, and a massive haemorrhagic infarction, as in our case, has not been reported.

In our case, H. influenzae was detected via CSF by PCR; however, we believe that the haemorrhagic infarction in our case was not caused by bacterial meningitis because it was observed on the side.

Patient’s perspective

We are glad that our son’s life was saved, although he had some neurological sequelae.

Learning points

► Causative organism can be identified by PCR of cerebrospinal fluid even with prior administration of antibiotics.
► Acute otitis media rarely causes cerebrospinal complications such as haemorrhagic infarction.
► It is important to empirically administer antibiotics in cases of haemorrhagic infarction.
Images in...

with otitis media, and the cell count in the CSF was inadequate for bacterial meningitis. It was speculated that the infarction was caused by the embolism of haematogenously spreading bacteria; the secondary bleeding after the infarction might have caused the leakage of bacteria into the CSF.

Paediatricians should remember that AOM rarely causes cerebrospinal complications.

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Disclaimer Case reports provide a valuable learning resource for the scientific community and can indicate areas of interest for future research. They should not be used in isolation to guide treatment choices or public health policy.

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