CASE REPORT

New-onset headaches secondary to spontaneous intracranial hypotension

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SUMMARY

We describe the case of a 54-year-old man who presented with new-onset positional headaches and seizures, which were determined to be secondary to spontaneous intracranial hypotension due to a cervicothoracic spinal cerebrospinal fluid leak, and its associated complications.

BACKGROUND

Spontaneous intracranial hypotension (SIH) is an under-recognised entity that can present with a myriad of neurological symptoms, the most common of which is headache. Certain complications of untreated SIH can be potentially life threatening. We would like to share our experience with the management of a patient with SIH.

CASE PRESENTATION First encounter

A 54-year-old man with no significant medical history was referred for the management of pressure-like holocranial, positional headaches with poor response to analgesics, and focal-onset generalised tonic clonic seizures, ongoing for a week prior to presentation. The headaches were orthostatic right from the onset. He already had



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Figure 1 MRI brain T2 FLAIR showing a right frontal hyperintensity—area of venous infarction. FLAIR, fluid attenuated inversion recovery.

a CT head and a lumbar puncture (LP), both of which were unremarkable. Opening pressure was not checked as the procedure was done with the patient in sitting position. Neurological examination showed no focal deficits. Magnetic resonance imaging/venogram of brain (MRI/V) done after the LP (figure 1) showed thrombosis of a right-hemispheric cortical vein, extending into the superior sagittal sinus, which was the likely explanation for the seizures. He was treated with levetiracetam, and heparin later transitioned to enoxaparin, and discharged home on symptom relief. MRI showed diffuse pachymeningeal enhancement typical of intracranial hypotension (figure 2).

Second encounter

He presented 4 days later for the recurrence of positional headaches and was found to have bilateral subdural hygromas (SDHs) (figure 3). His anticoagulation was reversed for CT myelogram, a blood patch was administered in the L3–L4 interspace and MRI of cervical/thoracic / lumbo-sacral (C/T/L-S) spine and CT myelogram (figure 4) located the site of the leak to be in the region of the lower cervical/upper thoracic spine. His anticoagulation was resumed, and he was discharged home.

Third encounter

He presented again 3 days later with progressive positional headaches, and this time he was found to have right-sided subdural haemorrhage at the site of the hygroma (figure 5). Once again, his anticoagulation was reversed, a second blood patch was administered and his SDH was managed conservatively.

INVESTIGATIONS

First encounter: MRI/V Second encounter: CT head, MRI C/T spine, CT myelogram Third encounter: CT head

DIFFERENTIAL DIAGNOSIS

Second encounter: the lumbar puncture could have contributed to the development of the SDHs in addition to the cervicothoracic spinal site of cerebrospinal fluid (CSF) collection identified on MRI spine/CT myelogram.

TREATMENT▶ Caffeine



Figure 2 MRI showing diffuse pachymeningeal enhancement.

- Strict bed rest
- Multiple blood patches

OUTCOME AND FOLLOW-UP

After discharge following the third hospitalisation, the patient has been headache-free and seizure-free, and does not have any focal neurological deficits at 3 months of follow-up.



Figure 3 CT head showing bilateral subdural hygroma.



Figure 4 CT myelogram showing contrast collection posterior to the thoracic spinal cord.

DISCUSSION

Dural enhancement with contrast/meningeal enhancement can be seen after an LP, but in our patient, MRI was done after the LP.

Our patient developed SIH and associated complications of cortical venous and venous sinus thrombosis, and SDHs. There are a few prior reported cases of venous sinus thrombosis in association with SIH.^{1 2} Cerebral venous sinus thrombosis has been described as an uncommon cause of iatrogenic postpartum headache after receiving epidural anaesthesia.³

The anticoagulation could have played a role in the development of SDH, but there have been reports⁴ of SDH in patients with SIH in absence of anticoagulation.



Figure 5 CT head showing right-sided subdural hygroma.

Headache Classification Committee of the International Headache Society in the third edition of the International Classification of Headache Disorders (ICHD-3) defined SIH as

a headache with low CSF pressure and absence of a procedure or trauma known to be able to cause CSF leakage, headache has developed in temporal relation to occurrence of low CSF pressure or CSF leakage, or has led to its discovery and is not better accounted for by another ICHD-3 diagnosis.⁵ The aetiology often remains elusive, but pre-existing dural weakness is thought to play a role. Clinical features of SIH and associated complications are a result of compression/traction/distortion of brain structures and/or venous engorgement, and include SDHs, cerebral venous and venous sinus thrombosis, and rebound intracranial hypertension, among others.⁶

Patient's perspective

I am still apprehensive about my condition, having sudden headaches are always at the back of my mind.

Learning points

- Spontaneous intracranial hypotension (SIH) can be a risk factor for the development of cortical venous and venous sinus thrombosis, and subdural hygromas.
- Serial blood patches are often required for the treatment of SIH.
- In most cases, there is no identifiable offsetting event leading to the development of SIH.

use of caffeine/theophylline and so on. Blood patching/surgical intervention are indicated in intractable cases. Epidural blood patch has been described to eliminate the determinants of cortical venous thrombosis (CVT) (vein dilatation and slow vein drainage) and can be exclusively used treat CVT by intracranial hypotension.⁷

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