INCIDENTAL DISCOVERY OF CAROTID ARTERY DISSECTION AS A CAUSE OF HORNER’S SYNDROME AND HEADACHE IN A PATIENT PRESENTING FOR FOLLOW-UP OF LUMBAR EPIDURAL STEROID INJECTION: A CASE REPORT WITH A THREE YEAR FOLLOW-UP

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Objective: To describe management challenges and follow-up data for three years in a case with spontaneous carotid artery dissection (SCAD).

Case Report: We present a case of SCAD that presented with Horner’s syndrome and headache. The patient developed the SCAD, Horner’s syndrome and headache one month after a lumbar epidural steroid injection for radicular pain. Three years follow-up experience with therapeutic challenges is described.

Conclusion: A life threatening cause of headache is described which occurred one month after receiving an uneventful lumbar epidural steroid injection. Despite no carotid arterial luminal flow restriction and anticoagulation, the headache and Horner’s syndrome persisted for three years. Stellate ganglion blocks were able to relieve both head pain and headache temporarily. In patients with new onset headache and Horner’s syndrome SCAD must be ruled out. If SCAD is detected, appropriate therapy including anticoagulation must be instituted immediately, since the potential for continued dissection or embolization remain significant.

Keywords: Headache, spontaneous carotid artery dissection (SCAD), lumbar epidural steroid injection

Spontaneous carotid artery dissection (SCAD) has been associated with a number of clinical syndromes including transient ischemic attack (TIA), cerebral vascular accident (CVA), cranial neuropathies and migraine headache. This case report presents an unusual presentation of SCAD that included cluster headache and Horner’s syndrome in a patient who presented to our pain clinic for follow-up of lumbar epidural steroid injection (LESI) for radicular low back pain.

Case Report

A 44-year-old Caucasian woman presented to the pain clinic for evaluation and treatment of radicular low back pain. Workup revealed a bulging disk at the L3-4 level for which she received a lumbar epidural steroid injection (LESI) using 80 mg methylprednisolone acetate in 2 ml of 1% lidocaine hydrochloride with resolution of the radiculopathy pain (VAS diminished from 8/10 to 1/10 at 48 hours). Two months later, she came to the pain clinic for follow-up at which time we noted that she had developed a right-sided Horner’s syndrome. We elicited a history of her having “bumped” her head in a supermarket one week following the LESI. The following day, she noted an intense headache confined to the right side of her face and cranium. The patient denied concomitant photophobia, nausea, vomiting, neurological symptoms or visual scintillation. The headache worsened over the next two days. She then developed a drooping of her eyelid. Her primary care physician prescribed Acetaminophen and Tramadol for her headache, which afforded no significant relief. As the headache got progressively worse over the next few weeks, she visited a neurologist who diagnosed cluster headaches and began an empirical regimen of prednisone and verapamil, also without reduction of her symptomatology. The patient was an office manager, without reduction of her symptomatology. The patient was an office manager, without reduction of her symptomatology.

The remainder of the physical examination was essentially unremarkable, including cranial nerve assessment. Of note, the carotid arterial pulses were normal and no bruits were auscultated.

Our workup included computed tomography (CT) scans of the head and neck and chest X-ray, which all failed to identify any pathology. A magnetic resonance angiogram (MRA) (Fig 1) revealed a dissection of the right internal carotid artery (ICA) with an intimal flap. Ultrasound of neck confirmed the MRA findings, which showed dissection in the right ICA and right common carotid artery (CCA) without significant narrowing of the ICA or the CCA.

Differential diagnosis for the spontaneous dissection of internal carotid artery included vasculitis and connective tissue disorders. Blood was sent for serum complement levels and rheumatoid factor, which were within normal limits. The patient was anticoagulated using heparin. Following anticoagulation,
the patient reported improvement in her ptosis, conjunctival injection, and a modest reduction in her headache pain score (30% to 40%).

At 3 months, a follow-up MRA revealed no interval change. The intimal flap at the origin of the ICA was again noted protruding into the lumen narrowing it approximately 65% involving a length of 8 mm. There was normal flow distally within the ICA. A follow-up MRI brain also noted no interval changes but revealed punctuate high T2 signal areas within subcortical white matter and faint increased T2 signal within the peritrigonal white matter, which could be related to ischemic or demyelinating disease. A MRA of the brain at the same time was normal and did not reveal extension of the intimal tear into the brain. She maintained regular follow-up in our Pain Clinic at monthly intervals for symptomatic headache and low back pain. She was being managed conservatively on medications which included Tylenol #3, clonazepam and Elavil. Her anticoagulation was still therapeutic and no further LESI were planned until her coagulation status normalized.

She continued to receive a multidisciplinary approach to management of her pain and SCAD issues which included a neurologist, pain psychologist and vascular surgeon. At 6 months follow-up, the vascular surgeon was considering placement of a vascular stent in her ICA, but decided to wait for some more time considering a risk-benefit assessment.

At 7 months follow-up, the patient presented to our Pain Clinic for the first time without Horner’s syndrome. She still complained of intermittent headache. Mood was upbeat and she was planning to return to work in the next week. At 8 months the anticoagulation was discontinued. She continued to have headache and occasional Horner’s syndrome. Her LBP was continuing to bother her. She had lost 20 pounds due to exercise and her mood was upbeat.

At 9 months follow-up, MRA neck again re-identified the flap in the ICA associated with 65% intimal stenosis. The flap now spanned approximately 9 mm in craniocaudal length. MRI and MRA of the brain were unremarkable. She had now increasing low back pain with radiculopathy. Considering that she was off anticoagulation and the dissection was stable, we performed LESI #2 using 80 mg methylprednisolone acetate in 2 ml of 1% lidocaine hydrochloride with resolution of the radiculopathy pain. She had a third LESI again using the same combination of medications at 11 months. At 13 months, her radiculopathy pain had decreased but she complained of increasing neck pain. She had a superficial cervical plexus block performed using 10 mL of 0.25% bupivacaine with 80 mg of methylprednisolone acetate with excellent neck pain relief. At 14 months, she presented with “mild” Horner’s syndrome with a new finding of “facial burning”. A stellate ganglion block using 8 mL of 0.25% bupivacaine and 80 mg of methylprednisolone acetate was done with excellent pain relief. The stellate ganglion block was repeated again at 16 and 18 months, with resolution of her neck and facial pain.

At the last follow-up at 36 months, her Horner’s syndrome had returned full-blown and she continued to complain of moderate headaches, which were intermittent.

**DISCUSSION**

Spontaneous carotid artery dissection (SCAD) typically occurs without any antecedent history of trauma or tumor. However, in a prospective study by the Canadian Stroke Consortium (1), careful history taking in patients with so-called spontaneous dissection has invariably revealed minor trauma. It is likely that the etiology and pathogenesis of spontaneous carotid artery and vertebral artery dissections are multifactorial and that mechanical factors, such as sudden neck movements, and underlying arteriopathy have a role. The underlying arteriopathy may be transient and may be related to a recent infection, or the arteriopathy may be related to a genetic abnormality. Our patient had a history of incidental head “bumping” one week after receiving a LESI. If the trauma really contributed to the dissection is conjecture. The temporal relationship of the Horner’s syndrome following the headache may imply the initial trauma as being related to the dissection. Typically, following trauma, a hematoma forms and causes a dissection of the arterial wall that result in compression and collapse of the arterial lumen. Nerve damage occurs secondary to an interruption of blood flow and the compression that results from local inflammatory re-

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Fig 1. **MRA of the neck revealing dissection of the internal carotid artery with a luminal flap**

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Dissection of the carotid artery is a dynamic process. The radiographic findings may change dramatically within a period of days or even hours. Although the radiographic appearance may worsen during the acute phase of dissection, about 90 percent of stenoses eventually resolve, two thirds of occlusions are recanalized, and one third of aneurysms decrease in size. This improvement takes place largely within the first two or three months after the dissection and is rare after six months. However, in our case the dissection has persisted in follow-up. Surgical or endovascular treatment should be reserved for patients who have persistent symptoms despite adequate anticoagulation. Surgical treatment consists of ligation of the carotid artery with an in situ or extracranial-to-intracranial bypass. These procedures can be associated with a high morbidity rate. Endovascular treatment, consisting of percutaneous balloon angioplasty and placement of one or more metallic stents, has gained favor over surgery as the initial therapy of choice once medical therapy fails (8, 9). However, the long-term results of carotid stenting are unknown, and the treatment of stent-related complications can be complex (10).

In summary, this case is fascinating in that a spontaneously occurring, and potentially life-threatening syndrome developed post-procedurally in an otherwise young healthy person seen in the pain clinic for lumbar radiculopathy. One week after receiving a completely uneventful LESI, she developed a sudden onset of intense headache accompanied by Horner’s syndrome. The diagnosis of SCAD was expeditiously made, and the patient underwent treatment for it. Despite no carotid arterial luminal flow restriction and acceptable anticoagulation, the patient’s headache and Horner’s syndrome persisted for three years. Additionally, she became emotionally labile and her symptomatology worsened during periods of melancholy or tearfulness. She is slowly improving, and fortuitously for her, never developed irreversible neurological sequelae from SCAD. In patients with new onset headache and Horner’s syndrome, detailed evaluation, including MRA, must be immediately undertaken to rule-out the possibility of SCAD. If there is a spontaneous dissection, appropriate therapy must be instituted immediately, since the potential for continued dissection or embolization remains significant until such therapy is instituted.

References