Closure of a patent foramen ovale is associated with a decrease in prevalence of migraine

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A patent foramen ovale (PFO) is one of the major causes of right-to-left shunt, and a causal relationship between migraine and a PFO has been suggested.1 We evaluated whether percutaneous closure of a PFO was associated with changes in the prevalence of migraine.

Methods. Patient selection. Patients with a PFO who had a paradoxical embolic event or systemic desaturation and who underwent a percutaneous closure in our center between February 1999 and September 2002 were included. The medical files were reviewed. The ethical committee approved the study.

Evaluation of migraine. A questionnaire was composed in such a way that a neurologist could diagnose migraine with or without aura (MA+ and MA−) according to the criteria of the International Headache Society. The questionnaire was sent to all patients and focused on three periods: 1 year before and 2 months and at least 6 months after percutaneous closure. Two neurologists blinded to the patients’ files diagnosed MA+ and MA−.

Statistical analysis. Within-patient comparisons of the absence or presence of migraine were performed with McNemar’s paired χ2 test. Interobserver reliability was evaluated by measuring the kappa coefficient. p Value < 0.05 was considered significant. All statistical analyses were performed with GB Stat software (version 8.0; Dynamic Microsystems, Inc., Silver Spring, MD).

Results. Patient characteristics. Seventy-six patients (mean age, 50.7 ± 12.9 years) were selected, and 66 completed the questionnaire. In 57 patients, the period between PFO closure and completing the questionnaire was >6 months. The characteristics of patients who completed the questionnaire are summarized in the table.

Prevalence of migraine. The median time interval between the occurrence of a paradoxical embolic event and the closing procedure was 162 days (range, 0 to 3,613 days). The time between PFO closure and administration of the questionnaire was 579 days (range, 110 to 1,419 days).

Migraine was present in 26 of 66 patients (9 men and 17 women; 39.4%). Twelve (18.2%) had MA+, and 14 (21.2%) had MA−. Two months after closure, the prevalence of MA+ and MA− decreased to 6.1% (4/66) and 6.1% (4/66; p < 0.05 vs before closure). At 6 months or more, the overall prevalence of migraine was 15.8% (9/57; p < 0.05 vs before closure). The prevalences of MA+ and MA− were 5.3% (3/57; p < 0.05 vs before closure) and 10.5% (6/57; p = 0.11 vs before closure). The frequency of migraine attacks also decreased (p < 0.05). Seven patients were taking potential prophylactic migraine drugs 6 months after closure (six, β-blockers; one, calcium antagonists). The kappa coefficient for interobserver reliability for migraine was 0.8 (p < 0.05).

Discussion. Patients with migraine have a high prevalence of PFO.2 An increased rate of MA+ among stroke patients with PFO was found compared with patients with PFO.3 A causal relationship between PFO and migraine has been proposed. In individuals with a right-to-left shunt, a lower dose of venous trigger substances may be needed to induce migraine because the shunt permits the pulmonary filter to be bypassed. Moreover, the prevalence of migraine seems to decrease subsequent to PFO closure in patients with decompression illness.4

We evaluated whether PFO closure in patients who mainly had cryptogenic stroke would be associated with changes in the prevalence of migraine. We found a high rate of migraine in patients with PFO (39.4%) and documented a significant and persistent decrease in prevalence of MA+ ≥6 months after PFO closure. The frequency of migraine attacks also decreased significantly. Our data might fit with the recently reported experience that MA+ decreased after PFO closure.4

The prevalence of migraine also decreases with age; however, we believe that the changes in our study are too pronounced to be explained by the natural history of migraine.5 Most of our patients were treated with low-dose aspirin, which could also influence migraine prophylaxis. The effect of low-dose aspirin, if any, seems to be modest.6 The placebo effect in migraine therapy is potent, but the decrease in prevalence of migraine in our study seems to be larger than reported placebo effect rates of 20 to 40%.7

Nevertheless, this study has important limitations. It is a retrospective, nonrandomized trial of patients selected from a hospital-based database. The questionnaire may be influenced by recall bias.

The prevalence of migraine in patients with a PFO is high. After ≥6 months, percutaneous PFO closure is associated with a decrease in the prevalence of MA+. Whether percutaneous PFO closure has the potential to manage migraine needs to be determined in a prospective randomized trial.

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References


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Prolactinoma presenting as painful postganglionic Horner syndrome

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Painful postganglionic Horner syndrome may be the initial presentation of an internal carotid artery (ICA) dissection and less frequently is a complication of cluster headache.1 However, it is not a frequent presentation of pituitary adenoma. We describe a patient with prolactinoma that initially manifested with painful Horner syndrome and improved on dopaminergic therapy. Eighteen months after treatment was discontinued, the patient sought treatment for a cavernous sinus syndrome that responded well to high-dose dopaminergic therapy.

Case report. A 52-year-old man was examined by one of our staff neurologists for an intermittent right hemicrania, retro-orbital pain and a narrowed palpebral fissure in March 1999. He had no history of alcoholism, head trauma, hypertension, diabetes mellitus, chronic headaches, impotence, or decreased libido. He smokes one pack of cigarettes daily. Examination revealed a 2-mm ptosis of the right upper eyelid and a 2-mm pupil that failed to dilate in the dark and after hydroxyamphetamine. The left pupil, in contrast, measured 3 mm and dilated to 5 mm in the dark and to 8 mm after hydroxyamphetamine. Both pupils reacted to light and accommodation. Visual acuity (V_-), color vision, visual fields, funduscopic, and remaining physical examination were normal. Brain MRI showed a discrete mass surrounding the carotid artery (figure, left). Serum prolactin level was elevated at 566.4 ng/mL (normal range 0.5 to 18.1 ng/mL). Bromocriptine 5 mg/d was initiated, and the patient experienced symptomatic improvement during the ensuing 6 months.

Thirty-two months later, he again sought treatment at our institution for worsening right-sided headache and diplopia. Eighteen months earlier, the patient decided to discontinue his bromocriptine. During the interval, he had no change in his history. Examination revealed a 4-prism diopter right exophoria and a partial right oculomotor nerve paresis with partial ptosis of the upper eyelid; symmetric limitation of infra- and supraduction, and adduction; and a 4-mm pupil with preservation of pupillary responses. Additionally, there was complete right sixth nerve palsy with hyposthesia in the right V1 and V2 dermatomes. Color vision, funduscopic, visual fields, and V_- were normal bilaterally. During the next 2 days, he developed a complete ophthalmoplegia of the right eye.

Brain MRI showed a 2.2 × 1.8 × 3-cm lesion filling the right cavernous sinus adjacent to, but not compressing, the optic chiasm (figure, middle). The tumor now encircled the cavernous portion of the right ICA. Serum prolactin at that time was 281.3 ng/mL. The patient was started on bromocriptine 15 mg daily and cabergoline 1 mg twice weekly. Ten days after cabergoline was added, serum prolactin level decreased to 22.1 ng/mL.

Four months later, follow-up MRI showed a 50% reduction in tumor bulk with the tumor still encircling the cavernous right ICA (figure, right). Clinically, he had resolution of the Horner syndrome and third and sixth nerve palsies, and return of facial sensation. In addition, there were no signs of aberrant regeneration. The patient was able to remain compliant with dual dopamine agonist therapy, except for nausea managed with ondansetron.

Discussion. A pituitary adenoma, presenting as a focal mass in the cavernous sinus, without diffuse sellar involvement, is distinctly uncommon. We are unaware of previous reports of pituitary tumors manifesting initially as postganglionic Horner syndrome. Although our patient did not have symptoms of hyperprolactinemia, a markedly increased prolactin level directed us to this diagnostic possibility. The successful effect of dopaminergic therapy, with resolution of the presenting symptoms and signs and virtual normalization of the serum prolactin level, left no doubt as to the diagnosis.

Prolactinoma is not initially thought of as a cause of an isolated involvement of postganglionic sympathetic fibers. Raeder2 first reported simultaneous impairment of sympathetic and trigeminal nerve fibers in a patient with an extrinsic mass lateral to the cavernous sinus. Our patient mimicked this presentation with a mass intrinsic to the cavernous sinus. Unlike his case, there was no involvement of motor trigeminal fibers.

Our patient had already been treated successfully with a single dopamine agonist but now presented with cavernous sinus syndrome after discontinuing therapy for 18 months.3 The main impetus to try simultaneous, high-dose dopamine agonists was to elicit a rapid response given the severity of the pain and ophthalmoplegia and to rule out the possibility of a prolactin-secreting carcinoma. The remarkable clinical and imaging response to dual dopamine agonists essentially served to elucidate the benign neuroendocrine character of the tumor, allowing a diagnosis without need for tissue biopsy or malignant tissue markers.4

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Figure. T1-weighted, contrast-enhanced MRI of brain. (Left) April 1999, performed after initial visit. (Middle) August 2002, ~1 month after starting dual agonist therapy. (Right) November 2002, only minimal symptoms present with reported 50% reduction radiographically in tumor bulk.
IV insulin during acute cerebral infarction in diabetic patients

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Although hyperglycemia can be a nonspecific response to stress, tight glycemic control during acute myocardial infarction and in postoperative ventilated patients has been shown to significantly reduce mortality. Animal and human studies suggest that hyperglycemia also augments acute cerebral ischemia, and clinical efficacy of tight glycemic control during acute cerebral infarction is being investigated. Good glycemic control during acute cerebral infarction in nondiabetic patients appears feasible and safe, but this has not been reported in patients with diabetes mellitus.

Methods. Our Institutional Review Board approved this study, and all subjects signed a valid informed consent. We prospectively administered an IV insulin protocol initiated within 12 hours after onset of cerebral infarction to 24 consecutive patients with admission hyperglycemia. Three revisions may further limit individualized doses of rapidly acting subcutaneous insulin, given right after meals, should reduce postprandial hyperglycemia within our protocol.

Discussion. Our insulin protocol, although still suboptimal, represents a reasonable combination of simplicity, safety, and glycemic control. Given the dynamic nature of cell death during acute cerebral infarction, it is important to note that the glucose level was lowered into the target range within 5 hours after starting insulin.

Postprandial hyperglycemia appears to be a substantial contributor to recurrent hyperglycemia. Individualized doses of rapidly acting subcutaneous insulin, given right after meals, should reduce postprandial hyperglycemia within our protocol. Protocol deviations can result in recurrent hyperglycemia. The deviations will likely be minimized if the protocol is kept simple and used frequently. Three revisions may further limit hyperglycemia: 1) higher frequency of glucose monitoring or continuous infusion, and 2) including glucose in the insulin solution, as was done in other studies, and 3) using a higher target range.

Good glycemic control during acute cerebral infarction in patients with diabetes mellitus type 2 is feasible and appears relatively safe. We plan to make additional modifications to our insulin protocol during a pilot clinical trial that is in progress. In this pilot trial, we will evaluate the feasibility of randomized, blinded treatment with IV (aggressive) vs subcutaneous (usual care) insulin in patients with acute cerebral infarction and admission hyperglycemia.

Additional material related to this article can be found on the Neurology Web site. Go to www.neurology.org and scroll down the Table of Contents for the April 27 issue to find the title link for this article.

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Meniere disease, and acoustic neurinoma.1-3 Toxic drugs, autoimmune inner ear disease, perilymph fistulae, viral disease, syphilis, Lyme disease, vascular disease, HIV, otosquelae.

evaluation, the unilateral deafness persisted without any other symptoms, revealed total deafness on the right side. There were no other abnormalities, including no nystagmus (also not present after use of Frenzel glasses) and no ataxia.

An MRI scan showed a small cerebellar infarct in the territory of the right posterior inferior cerebellar artery (PICA) on the T2-weighted image (figure, A). On the sagittal T1-weighted images, the distal part of the right vertebral artery was hyperintense, and no flow void was seen (not shown). A contrast-enhanced MR angiography showed a normal proximal right vertebral artery that showed irregularities and tapering in the high cervical portion. The distal portion of the right vertebral artery was occluded (figure, B and C).

The combination of image findings suggest vertebral dissection.

In this patient, sudden deafness, vertigo, and neck pain were caused by a spontaneous right vertebral artery dissection. The patient was treated with low molecular weight heparin, especially to prevent new neurologic signs and symptoms. During follow-up evaluation, the unilateral deafness persisted without any other sequelae.

The differential diagnosis of sudden deafness includes trauma, viral disease, syphilis, Lyme disease, vascular disease, HIV, ototoxic drugs, autoimmune inner ear disease, perilymph fistulae, Meniere disease, and acoustic neurinoma.1,3

The symptoms of sudden deafness combined with vertigo and neck pain suggested the diagnosis of a vertebral artery dissection.

Discussion. Sudden deafness has been described in anterior inferior cerebellar artery (AICA) infarction. Lee et al. described the signs and symptoms of 12 patients with unilateral AICA infarction.1 Vertigo was the initial symptom in all patients, accompanied by a horizontal-rotatory nystagmus beating toward the healthy side. Sensorineural hearing loss was present in 11 patients. Unlike our patient, all 12 patients had gait and limb ataxia caused by involvement of the middle cerebellar peduncle or anterior inferior cerebellum.1

Another recent article described bilateral deafness in vertebral insufficiency caused by stenosis of the basilar artery. Vertigo was present, but no nystagmus was observed in this patient as well.4

Our patient had unilateral sudden deafness caused by a right vertebral dissection. In most patients, the internal auditory artery originates from the AICA, but in a few, it branches off the PICA. In our patient, most likely, small emboli originating from the dissection caused a small selective cerebellar and cochlear infarction, sparing the vestibular apparatus. This led to unilateral sensorineural hearing loss without nystagmus. In conclusion, for patients presenting with neck pain and sudden unilateral sensorineural deafness, with or without vertigo, a vertebral artery dissection should be considered and MR angiography performed.

References

Sudden unilateral deafness due to a right vertebral artery dissection

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Case report. We present a 42-year-old woman without a known connective tissue disorder, who sought treatment at the emergency department because of sudden right-sided deafness, vertigo, and tendency to fall to the right. She also reported right-sided neck pain for 3 days.

Neurologic examination, performed 1 day after the onset of symptoms, showed irregularities and tapering in the high cervical portion. After use of Frenzel glasses, her vision was normal, and no flow void was seen (not shown). A contrast-enhanced MR angiography showed a normal proximal right vertebral artery that showed irregularities and tapering in the high cervical portion. The distal portion of the right vertebral artery was occluded (figure, B and C).

The combination of image findings suggest vertebral dissection.

In this patient, sudden deafness, vertigo, and neck pain were caused by a spontaneous right vertebral artery dissection. The patient was treated with low molecular weight heparin, especially to prevent new neurologic signs and symptoms. During follow-up evaluation, the unilateral deafness persisted without any other sequelae.

The differential diagnosis of sudden deafness includes trauma, viral disease, syphilis, Lyme disease, vascular disease, HIV, ototoxic drugs, autoimmune inner ear disease, perilymph fistulae, Meniere disease, and acoustic neurinoma.1,3

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Figure. (A) MRI (T2-weighted) showing a small right-sided posterior inferior cerebellar artery (PICA) infarction. (B and C) Contrast-enhanced MR angiography showing a normal proximal right vertebral artery with irregularities and tapering in the high cervical portion.
Plasmapheresis improves outcome in postinfectious cerebellitis induced by Epstein–Barr virus

Jeremy D. Schmahmann, MD

Postinfectious cerebellitis (PIC) is a rare complication of Epstein–Barr virus (EBV) infection. Whereas its course is usually regarded as self-limited, it can be prolonged and produce considerable residual disability. We report two adults with persistent PIC after EBV infection who demonstrated complete and rapid recovery with a course of plasmapheresis (total plasma exchange [TPE]).

Case 1. A 20-year-old woman (Case 15 in reference 1) developed fever and myalgia, followed 2 weeks later by onset over 3 days of slurred speech, gait instability, and difficulty judging distances. Examination revealed hypermetric saccadic eye movements, cerebellar dysarthria, dysmetria, dysdiadochokinesia, and ataxic gait. She had flattened affect, decreased verbal output and phonemic fluency, perseveration, and poor visual–spatial planning. Results of strength, reflex, and sensory testing were normal. Gadolinium-enhanced brain MRI and EEG were normal. CSF revealed three lymphocytes, normal protein, and glucose. Heterophile antibody was positive; serum EBV titer was 1:586; EBV viral capsid was 1:1230; restricted immunoglobulin (Ig) G was 1:320; and EBV nuclear antigen (EBNA) was 1:20, diagnostic of chronic or recurrent EBV infection. CSF viral capsid, diffuse and restricted IgG, and EBNA were normal. The persistence of motor and cognitive symptoms at 1 month prompted treatment with plasmapheresis. There was marked improvement in a stepwise fashion after each of six sessions of TPE. One week after treatment, the examination was only minimally abnormal, and 1 month after treatment, all cerebellar motor tests and neuropsychological tests had returned to normal, except for tests of complex and unstructured list learning.

Case 2. A 28-year-old man developed fever, myalgia, and pharyngitis. Two to 3 weeks later he had vertigo, ataxia, dysarthria, untidy handwriting, and incoordination of his limbs. He had difficulty concentrating and making quick decisions, and his family noted a flattened affect. Brain MRI and EEG were normal. EBV IgG titer was 1:640; IgM was elevated (1/10); and EBNA was negative, consistent with primary infection ~6 weeks previously. Symptoms persisted, and examination 3 months after the onset of the illness revealed hypermetric saccades, cerebellar dysarthria, upper extremity rebound, dysmetria and dysdiadochokinesia, widened stance, and ataxic gait. Strength, reflexes, and sensation were normal. Attention, declarative learning, and phonemic fluency were reduced; thinking was concrete; and he perseverated on the Luria diagram. This lack of recovery prompted treatment with plasmapheresis. During the course of five TPE sessions, the cerebellar motor syndrome abruptly improved, and thinking and personality reverted to the premorbid state. Three months after TPE, the examination was normal.

Discussion. Acute cerebellar meningoencephalitis from EBV infection is rare, occurring more commonly in children than in adults. It is characterized by an acute cerebellar syndrome sometimes with brainstem signs or diffuse encephalitis, meningismus, CSF pleocytosis, positive serologic tests for EBV in the CSF, and MRI abnormalities.

In contrast, PIC is thought to be an immune-mediated response to viral infection occurring 1 to 3 weeks later (range, 1 to 43 days), characterized by ataxia and dysmetria with minimal CSF pleocytosis, no systemic features, and negative CSF EBV titers despite positive serum titers. Neuropsychiatric features are also observed.1,2 MRI is usually normal, but SPECT scans may show decreased3 or increased perfusion.4

Reports generally stress the benign nature of postinfectious cerebellar ataxia,5 but Connolly et al.6 found that residual motor and cognitive deficits persisted for months or years, and in some cases deficits were permanent. Persistent cerebellar ataxia has also been noted in adults.6

Both our patients remained prominently compromised 1 month and 3 months after the onset of symptoms, and this time course, together with serologic tests in both patients and CSF analysis in Case 1, supported the diagnosis of EBV PIC. Continued clinical disability prompted the decision to treat. Recovery was rapid, complete, and occurred within days of the full course of conventional TPE (five to six exchanges), indicating a cause-and-effect relationship.

Successful improvement of PIC with IV Ig has been noted, but to our knowledge there are no published accounts of the use of TPE in this disorder.4,7 The success of TPE in our patients suggests that immune-modulating therapy may limit the duration of illness and potential long-term motor and cognitive deficits in patients with PIC from EBV. Prospective clinical trials could address this possibility more rigorously. The rapid recovery of PIC with TPE in 3 months implies that there is a protracted stage of antibody-mediated, reversible neuronal toxicity before Purkinje cell death.

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References

Insulinoma misdiagnosed as intractable epilepsy

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Many conditions may mimic epileptic seizures, including movement disorders, cardiac arrhythmias, psychiatric disorders, or metabolic abnormalities. Among these, insulinoma is a rare but curable cause of hypoglycemia often misdiagnosed as epilepsy or a psychiatric disorder. We report three patients with hypoglycemic spells caused by insulinoma. All patients were diagnosed during a video-EEG monitoring session initially performed for the evaluation of intractable epilepsy.

Case reports. Case 1. A 37-year-old woman had a 6-year history of stereotyped paroxysmal episodes of paresesthesias of the left arm, spreading to the face and lower limbs, and lasting ~20 minutes. Episodes could last up to 2 hours and included gait incoordination, mild agitation, emotional lability, and incoherent speech, which was followed by fatigue. Frequency of the attacks gradually increased up to twice a week. The patient was initially diagnosed with partial epilepsy, refractory to carbamazepine. No interictal activity was recorded during a prolonged video-EEG session. Brain MRI was normal, and interictal SPECT revealed bifrontal and right posterior temporal hypoperfusion. A fasting blood glucose level was obtained and showed mild hypoglycemia at 3.2 mmol/L (normal, 3.8 to 6.1 mmol/L). A prolonged fast reproduced a typical attack after 8 hours, along with concomitant severe hypoglycemia (0.9 mmol/L) and high insulin levels, confirming the diagnosis of insulinoma. An endoscopic ultrasound detected a small 1.5-cm nodular lesion over the pancreatic tail. Surgical removal of the tumor led to full remission of the symptoms.
Case 2. A 44-year-old man had a 2-week history of spells characterized by awkward behavior (crawling, swimming, circling, slamming, and vocalizations), unresponsiveness, and paranoia. Most of these episodes occurred late at night or early morning, lasting from 30 minutes to 6 hours, followed by sleepiness. The attacks increased in intensity and frequency up to a daily basis. These spells were initially misdiagnosed as anxiety disorder and then as complex partial seizures refractory to valproic acid. The EEG, head CT, and glucose levels were normal. A prolonged fast was performed during concomitant video-EEG monitoring. A typical attack was induced after 5 hours associated with ictal diffuse slow waves and significant low blood glucose level (1.7 mmol/L). Concomitant insulin and C-peptide measures confirmed the inappropriate secretion of insulin. An abdominal CT revealed a 2-cm mass. The patient was cured by a distal pancreatectomy.

Case 3. A 55-year-old woman had a 7-month history of episodic stereotyped confusional spells characterized by psychomotor slowing, unresponsiveness, abnormal posturing, and perseverative unpurposeful behavior. The attacks lasted from a few minutes to 6 hours and occurred up to three times a day despite treatment with carbamazepine and clobazam. Routine serum glucose levels and brain MRI were normal. The patient was admitted to our epilepsy unit with a working diagnosis of psychogenic nonepileptic seizures. No epileptic interictal activity was observed during prolonged video-EEG monitoring. A typical attack was recorded, associated with diffuse slow waves (figure, A), during spontaneous hypoglycemia (1.4 mmol/L). A prolonged fast was performed and induced a spell after 9 hours with simultaneous low blood glucose (1.3 mmol/L) and high insulin and C-peptide levels. Abdominal MRI demonstrated a 1-cm lesion in the pancreatic tail (figure, B). The patient was cured by resection of the lesion.

Discussion. Insulinomas may lead to fasting hypoglycemic spells caused by excessive (and time-inappropriate) secretion of insulin. Insulinomas are rare, with 4 cases per 1 million patients each year. The majority (>90%) are benign solitary adenomas, whereas up to 10% are associated with multiple endocrine neoplasia type I syndromes. The main challenge of these rare tumors is their recognition. Patients are frequently misdiagnosed with epilepsy or a psychiatric disorder, which was the case in our three patients. It is possible that some of the spells were complex partial seizures secondary to hypoglycemia, although this is doubtful because spells were atypical in semiology, lengthy in duration, unresponsive to anticonvulsant agents, and not associated with ictal epileptic activity during monitoring. In a recent series, median interval from onset of symptoms to diagnosis is 2 years, with a range of 1 month to 30 years. The diagnostic delay in our patients ranged from 6 months to 8 years. Fortunately, none experienced serious consequences. Late recognition may be caused by many factors, including the rarity of the condition, the lack of specificity of most symptoms, and the blunt of autonomic symptoms from...
Changes in the appearance of venous sinuses after treatment of disordered intracranial pressure

Debra B. Baryshnik, MD; and Richard I. Farb, MD

Cerebral angiographic findings in idiopathic intracranial hypertension (IIH) and spontaneous intracranial hypotension (SIH) have been described. Conventional cerebral venography has demonstrated tapered stenoses and filling defects in the transverse sinuses of patients with IIH and SIH. The stenoses have also been found in >90% of IIH patients imaged with auto-triggered elliptic-centric-ordered three-dimensional gadolinium-enhanced MR venography (ATECO MRV). Conventional cerebral angiography demonstrated prominence of intracranial venous structures in patients with SIH. To date, MRI correlates of these dilated venous channels have not been described. This report discusses the use of ATECO MRV to demonstrate the normalization of venous sinus caliber associated with the resolution of these two disorders of intracranial pressure (ICP).

Case reports. Case 1. A 21-year-old woman was diagnosed with IIH. CSF opening pressure was 32.5 cm H₂O. Months later, she was admitted to our institution with persistent headache and blurred vision despite maximal doses of acetazolamide. Examination revealed an obese woman with bilateral papilledema. CSF pressure was normal (19 cm H₂O). A CSF nuclear medicine flow study was normal. MRI of the brain demonstrated dural enhancement secondary to the shunt. ATECO MRV demonstrated normal-appearing venous sinuses with resolution of dural sinus narrowing (figure, A). A ventriculoperitoneal shunt was inserted, resulting in symptom resolution.

Eighteen months later, the patient had refractory headaches exacerbated by activity, photophobia, nausea, and vomiting. Her medications included acetazolamide 250 mg twice daily and acetaminophen 300 mg/codeine 30 mg two to nine tablets daily for 3 months. CSF pressure was normal (19 cm H₂O). A CSF nuclear medicine flow study was normal. MRI of the brain demonstrated dural enhancement secondary to the shunt. ATECO MRV revealed normal-appearing venous sinuses with resolution of dural sinus narrowing (figure, B). The patient was diagnosed with transfemoral vertebral angiogram and treated with percutaneous transluminal angioplasty.

Case 2. A previously healthy 19-year-old woman with headache and diplopia was transferred to our institution. Her headache was sudden in onset, exacerbated by movement, and severe in the upright position. Examination revealed bilateral abducens palsies. MRI revealed findings typical of intracranial hypotension, with marked venous sinus dilatation and increased intracranial sinus caliber (figure, C). The patient was discharged on acetazolamide 250 mg twice daily and a percutaneous dural shunt was inserted, resulting in symptom resolution.

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including inferior displacement of the cerebellar tonsils, flattening of the anterior aspect of thepons, small bilateral subdural hygromas, thickened appearance of the dura, and diffuse pachymeningeal enhancement after gadolinium injection. An ATECO MRV demonstrated dilatation of the dural venous sinuses (figure, C). A diagnosis of SIH was made. She was treated with a blood patch, and the headache improved. One year later, repeat imaging revealed resolution of the aforementioned stigmata of intracranial hypotension. Specifically, the dural venous sinuses returned to normal caliber (figure, D).

Discussion. These cases demonstrate normalization of venous sinus caliber associated with the resolution of IIH and SIH. The improved imaging and noninvasive nature of ATECO MRV over other venographic techniques have made this possible.\(^3\) The dynamic changes in venous sinus caliber can be explained using the Monro–Kellie hypothesis.\(^4\) In IIH, the increase in CSF pressure is compensated for by a decrease in the intracranial blood volume. We propose that the dural venous sinuses act as a volume-pressure reservoir that can accommodate these changes. This is seen on ATECO MRV by apparent venous sinus stenosis in patients with IIH. Using the same reasoning, in SIH, a decrease in CSF volume (and thus decrease in pressure) would be accompanied by an increase in the intracranial blood volume. This is seen on ATECO MRV as enlargement of the dural venous sinuses.

Although CSF leakage into extradural space has been accepted as the pathophysiology of IIH, the pathophysiology of IIH has slowly been delineated. Cerebral venous manometry in patients with IIH demonstrated a pressure gradient along the transverse sinuses, suggesting the presence of a functional obstruction to venous outflow.\(^5\) Subsequent investigations demonstrated that removal of 20 to 25 mL of CSF corresponded with decrease in central venous sinus pressure, suggesting that the obstruction to venous outflow in IIH is a secondary phenomenon.\(^6\) These researchers were unable to use venography to demonstrate the normalization of the appearance of the dural venous sinuses with improvement of CSF pressure.

Case 1 provides evidence that the narrowing in the transverse sinuses seen with elevated ICP is reversible when ICP normalizes. This supports the hypothesis that the deformity of the venous sinuses in IIH is a secondary event. Case 2 demonstrates the complementary phenomenon of normalization of venous sinus caliber after clinical and imaging resolution of SIH. These findings suggest that there is a direct and dynamic relationship between CSF pressure and the appearance of the venous sinuses.

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Auditory dysfunction in chronic inflammatory demyelinating polyradiculoneuropathy

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Cranial nerve involvement, most commonly affecting the facial and oculomotor nerves, has been reported in chronic inflammatory demyelinating polyradiculoneuropathy (CIDP).\(^1\) Symptomatic involvement of the vestibulocochlear nerve is well documented in chronic inflammatory demyelinating polyradiculoneuropathy. Brainstem auditory evoked potentials (BAEPs) confirmed bilateral retrocochlear dysfunction. Treatment with high-dose prednisone was associated with nearly complete recovery clinically and neurophysiologically.

Case report. A 66-year-old man sought treatment for sudden bilateral hearing loss and paresthesias of the lower extremities. Ten months earlier, CIDP had been diagnosed based on clinical signs and symptoms (symmetric muscle weakness, sensory ataxia, distal hypesthesia, and absent tendon reflexes), results of nerve conduction studies (i.e., motor distal latency and F waves of the ulnar and median nerves >130% of the upper limits of normal, slowing of nerve conduction velocities in ulnar and peroneal nerves <70% of the lower limit of normal, and presence of a conduction block in the peroneal nerve), and CSF analysis (elevated protein without pleocytosis).\(^2\) He was treated with IV gam-maglobulin, which had resulted in a nearly full recovery. At present, he does not use any medications besides prednisone for his biventricular use of ciprofloxacin for recurrent cystitis and vitamin B12 supplementation once a month.

At presentation, the patient had sudden bilateral hearing loss and hypesthesia of the feet. He had not noted a change in muscle strength or vertigo. Physical examination showed no abnormalities of the cranial nerves, with the exception of bilateral sensorineural hearing loss. There was areflexia and hypesthesia in the
Abnormal BAEPs in patients with CIDP have been described, but none of these patients had auditory symptoms. Moreover, in these studies a second BAEP investigation was not performed after management of the CIDP, thus leaving unanswered the question whether BAEP abnormalities are reversible, as might be expected in an immune-mediated demyelinating disorder. The results of the second BAEP study in our patient clearly showed that the BAEP abnormalities were reversible.

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References


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