Idiopathic Intracranial Hypertension Update

Christopher J. Borgman, OD, FAAO
Southern College of Optometry
Memphis, TN
Disclosures...

- I have no disclosures to report.
Case:

“Doc these headaches are killing me!... will new glasses help?”
26 YO AAF------ “fits the profile”

CC: “progressive loss of side vision and now central vision”
  OS>>OD, (+)HA’s behind eyes, onset 2-3 mo, referred by PCP after no improvement with oral antibiotics for “sinus infection”

BCVA = 20/25- OD, 20/1000 OS with Snellen

Color Vision = 1/7 OD, UTT OS with HRR Plates

CT = ortho at D & N

IOP = 16 mmHg OD, 18 mmHg OS with Goldmann

BP= 114/73 in-office

PMH = (+)HTN—controlled, (-)DM, (-)pregnant

Meds = Atenolol

Allergies = NKDA
MRI results for patient

- MRI = WNL
- MRA = WNL
- MRV = bilateral stenosis of transverse sinuses
- LP opening pressure = 295 mm/H20
- LP Cytology = normal
Idiopathic Intracranial Hypertension (IIH)

- Aka: Pseudotumor Cerebri (PTC)
- Defn: increased ICP without a mass effect and with normal CSF composition
- MOA: intracranial venous drainage obstruction; decreased CSF drainage
- *Headaches = 90% of cases*
  - Most common Sx
- Blurred vision, *loss of VF (up to 96%), visual obscurations, permanent visual loss (25%)*
- *Papilledema = most common Sn; 89-95% of cases*
- F>>M (90% vs. 10%); females of child-bearing age
- Risk factors = obesity (70% of IIH), delayed CSF absorption, venous outflow abnormalities/increased cerebral venous sinus pressure
Modified Dandy’s Criteria (Revised 2008)

1. Absence of mass lesion or hydrocephalus with CT or MRI
2. Elevated CSF opening pressure upon lumbar puncture with normal CSF profile
   - Non-obese patient >200 mmH2O = Abnormal
   - Obese patient >250 mmH2O = Abnormal
3. Intact neurological exam with the exceptions of visual disturbances, and/or 6th nerve palsy, and/or papilledema
PTC/IIH Symptoms

1. *Headache* (worse upon awakening) (90%)
2. Transient Vision Loss (62%)
3. Pulsitile Tinnitus (48%)
4. Blurred Vision
5. Vomiting
PTC/IIH signs...

- **Papilledema!** In up 95% of cases! (Puffer et al. 2014)

- “With rare exception, all PTC/IIH patients have papilledema, a hallmark of subacute intracranial hypertension.” ---Galgano et al. (2013)

- Although papilledema is present in the vast majority of PTC/IIH patients, its absence it not an exclusionary criteria.” ---Galgano et al. (2013)
Ocular work-up in IIH...

- Visual Acuity
- **Visual Fields**
- EOM’s
- Fundus Exam
- Retinal Imaging (FP, OCT, etc.)
- Color Vision
- Contrast Sensitivity

Most important to assess in IIH
Visual Acuity and Color Vision in IIH

- **Visual Acuity:**
  - Acuity tests foveal function
  - Not typically affected unless edema extends into central 10° of fixation

- **Color Vision:**
  - Only been found to be abnormal in ~20% of cases
  - Ishihara defects only noted in the existence of moderate to marked visual loss and optic atrophy
  - Not the most reliable way to follow patients
EOM’s/VF’s in IIH...

- **EOM’s:**
- If present, uni/bilateral 6th nerve palsies are present 2° stretching nerve between apex of clivus bone/Dorello’s canal and exit zone of 6th nerve on brainstem
- **Dilation required** in all 6th nerve palsies to rule out/in papilledema per Will’s Eye

- **Visual Fields:**
- *Most important test to follow for changes*
- Enlarged blindspot first to show, followed by generalized constriction, and nasal defects.
- Any kind of defect is possible though...
VF’s in neuro-optometry....
Is testing the central 30° enough?

“Humphrey SAP has replaced Goldmann perimetry in clinical practice despite fears that peripheral visual field defects may be missed. This fear seems unwarranted as only 1-2% of patients with nonglaucomatous VF defects have abnormalities in the peripheral field beyond 30° degrees in the absence of central field defect.”

Alternatively said....98-99% of neurological VF defects will show up in the central 30° when tested....pretty good odds!

Neuroimaging/workup in IIH...Order is important...why???

1. Order MRI/MRV first

2. Followed by lumbar puncture if MRI/MRV is normal
   - >200 mmH2O in nonobese patients = abnormal
   - >250 mmH2O in obese patients = abnormal

- Herniation through foramen magnum can compress upper medulla which is where the respiratory and cardiovascular centers are located → Death

What does herniation look like?
What are we looking for in work up?

- **MRI**
  - Rules out space occupying mass, hemorrhage, etc.
  - Empty sella, pituitary deformities, distention of ON, posterior globe flattening

- **MRV**
  - Rules out transverse sinus stenosis and/or venous sinus thrombosis

- **Lumbar Puncture (LP)**
  - Document elevated opening/intracranial pressure

- **LP cytology**
  - Rule out infectious meningitis, blood, and other possible issues/causes
CSF Hydrodynamics

- Oh crap......you’re kidding me right?
Where is CSF made again???

Answer: Choroid Plexus in Lateral, 3rd, & 4th Ventricle
Where does CSF drain to?

- CSF is absorbed from the subarachnoid space across the arachnoid villi into the venous circulation.

- The arachnoid villi act as one-way valves between the subarachnoid space and the dural venous sinuses. The rate of absorption correlates with the CSF pressure.

  - “Pressure gradient valves”

- Dural Venous Sinus Thrombosis important to rule out with MRV!
Pathophysiology of IIH...

- Any blockage along this CSF drainage pathway or in the venous sinus drainage can result in increased ICP!

- Bottom Line MOA = obstruction of intracranial venous drainage
PTC /IIH Treatment Options...

1. Weight loss (5-10% is sometimes curative!)
2. Carbonic anhydrase inhibitors
   - Acetazolamide and/or Topiramate
   - No oral steroids → weight gain
3. Ventriculoperitoneal Shunt / Lumboperitoneal Shunt
   - Headaches only; vision stable
4. Optic Nerve Fenestration
   - Vision/Visual Field worsening; no headaches
5. Venous Sinus Stenting
   - In venous sinus stenosis
Topamax vs. Diamox?

- **Acetazolamide** = CAI inhibitor; works on ciliary body and choroid plexuses

- **Topiramate** = novel anticonvulsant with many MOA; epilepsy/migraines

- Sulfonamide drug....be careful of sulfa allergies

- Also has carbonic anhydrase inhibition component; and decreases appetite
  - Weight loss of 5-10% alone may be curative in some cases of IIH
  - Average weight loss of 7.3% was obtained in one year on medication

- Monitor for angle closure glaucoma and myopic shift!!!
  - 85% of this happens within first 2 weeks of therapy

- Ciliochoroidal infusion occurs
  - **MOA** = lenticular/uveal effusion and ciliary edema causing forward displacement of the lens-iris diaphragm with resultant narrowing of the anterior chamber.

- D/c med; cycloplege and IOP lowering meds may be needed. LPI not helpful.

How long should CAI’s be maintained?

- Can/Should Tx ever be discontinued once Sn/Sx are under control?

- A long-term follow up study was done in PTC patients using a CAI (acetazolamide) over 6.2 years.

- 54 total patients followed for over 6 years

- 60% of patients experienced multiple recurrent episodes over this time span

- None of the recurrences occurred while maintained on acetazolamide!

- Good evidence to maintain longterm Tx???

Pregnancy Considerations...

- **Topamax** = FDA Category D; evidence shows up to 10-20% of dose can be found in infants who are nursing; avoid in pregnancy

- **Acetazolamide** = FDA Category C; case reports of placenta crossing; has been avoided in pregnancy in the past.....new evidence to suggest otherwise?

- If both are avoided in pregnancy, then sometimes repeat LP’s may be necessary in short term to keep ICP down; inherent risks...

“The use of carbonic anhydrase inhibitors (CAIs) has a large pool of human data on which to base clinical decisions. The source is the National Collaborative Perinatal Project (NCPP) conducted by the NIH from 1959 through 1974. This study monitored more than 50,000 mother-child pairs and 1,024 instances of systemic usage of acetazolamide during pregnancy. In the resulting offspring, there were 18 instances of malformations. The predicted number due to chance was 18.06. This suggests that the incidence of malformations from acetazolamide exposure during pregnancy is no greater than the natural incidence. In the same study, there were 12 documented first trimester exposures to acetazolamide. No anomalies were observed in the resulting offspring.”

---Steven Odrich, MD (Bronx, NY)

- 12 patients on Diamox 500 mg BID PO during pregnancy
- No adverse side effects nor congenital malformations noted
- Cited the results of the Collaborative Perinatal Project as well

- “In summary, there is no convincing evidence from the literature for the recommendation to limit the use of acetazolamide for IIH in pregnancy. Although the use of acetazolamide might be restricted in the first trimester, this recommendation may have a more medicolegal than medical rationale. It is our recommendation that acetazolamide be considered if the risk of nontreatment (e.g., progressive visual loss) is sufficiently high to warrant its use.”
Question to y’all:

Is it appropriate for O.D.’s to prescribe Diamox for these patients long term???
Surgical Considerations...

1. Headaches only, vision stable (can be used for both HA’s and vision too)
   ----- LP shunt, VP shunt
2. Vision loss/VF worsening despite maximal medical Tx
   ----- Optic Nerve Fenestration
3. Venous sinus thrombosis
   ----- Anticoagulants
4. Venous sinus stenosis
   ----- Venous sinus stenting

- Majority can be managed via weight loss and oral meds (Diamox)
1A) Ventriculo-Peritoneal Shunts (VPS) and Ventriculo-Atrial Shunts (VAS) in IIH
1B) Lumboperitoneal shunts in IIH

- L3/L4 or L4/L5 spaces most commonly used
- Drain into peritoneal space like VPS
VPS vs. LPS...

**Ventriculo-Peritoneal Shunt (VPS):**
- Infection rate of 7-15%
- 20% revision rate q2 yrs

**Lumbar Peritoneal Shunt (LPS):**
- Infection rate of 1%
- 50% revision rate q2 years

- “In short, most shunted PTC patients require multiple revision surgeries during their lifetime.”
  ---Galgano MA et al. (2013)
2) **ON Sheath Fenestration...**

- **Defn:** make slits in ON sheath to reduce the local pressure around the optic nerves.
- ~50% of unilateral ON sheath fenestration procedures results in resolution of visual symptoms in both eyes.
  - Both optic nerves are connected via the subrachnoid tissue around the optic chiasm
- Typically only done for visual Sn/Sx *without* headaches...
  - If headaches → shunt procedure is better option
- Safe and effective *up to 10 years* per several studies
- Revision rate is usually very low; 1 procedure per lifetime generally
DVS Thrombosis Considerations...

- Blood clots in young people are NOT normal...
- If DVST occurs, hematological workup and anticoagulant therapy is required.
  ---Subramanian PS et al. (2014)
- Consider: CBC with diff, CMP, lipid panel, PT/PTT, Protein S, Protein C, Homocysteine levels, Lupus anticoagulant, anticardiolipin, Factor V Leiden, Prothrombin mutation, Antithrombin III mutation, Sickledex screen, hemoglobin electrophoresis
DVS Thrombosis Treatment...

- Rule out clotting disorder, infection, etc.
- Aggressive anti-coagulation (heparin, warfarin, clopidogrel)
- Not a candidate for DVS Stenting in vast majority of cases...
- If anticoagulation and oral CAI’s do not work, then may need shunt surgery
Dural Venous Sinus Stenosis
Dural Venous Sinus Stenosis (DVSS)

- **Defn:** focal, narrowed section of dural venous sinuses causing back up/turbulent venous blood flow

- *Most common at junction of Sigmoid and Transverse sinuses*

- Not a true blood clot like DVST is...

- Treatment = weight loss, oral CAI, and/or DVS Stenting procedure
Focal stenosis has been demonstrated in 90+% of IIH patients using advanced imaging techniques.

Furthermore, focal stenosis in the same sinus territory was only demonstrated in 6.8% of asymptomatic control subjects.

Might be on to something here....
4) Dural Venous Sinus Stenting

- Right transverse sinus is dominant in 73% of cases

- **MOA:** Increases drainage of venous blood from venous sinus system which helps with the pressure dependent valves, arachnoid villi granulations, allowing them to clear CSF into the venous system more efficiently/quickly → decreasing intracranial pressure

- High frequency of resolved or improved HA’s and papilledema with this method
DVS Stenting...

- Not every patient is a candidate for DVS Stenting

- **Criteria Needed:**
  1. Presence of venous sinus stenosis (MRI/MRV) ; not thrombosis...
  2. Transvenous manometry across the stenosis >10 mmHg differential

- Catheter with stent and manometer placed in femoral vein and “fished” upwards to location of sinus stenosis

- **Post-Op Medications:**
  - Plavix 75 mg x 6-12 weeks then d/c
  - ASA 325 mg for life
Transvenous Manometry in IIH

- Significant = >10 mmHg pressure differential of proximal vs. distal locations in respect to stenosis

<table>
<thead>
<tr>
<th>Location</th>
<th>Pre Stent</th>
<th>Post Stent</th>
</tr>
</thead>
<tbody>
<tr>
<td>Superior Sagittal Sinus</td>
<td>45 mmHg</td>
<td>21 mmHg</td>
</tr>
<tr>
<td>Right Transverse</td>
<td>46 mmHg</td>
<td>20 mmHg</td>
</tr>
<tr>
<td>Right Sigmoid</td>
<td>15 mmHg</td>
<td>19 mmHg</td>
</tr>
<tr>
<td>Difference</td>
<td>31 mmHg</td>
<td>1 mmHg</td>
</tr>
</tbody>
</table>

Dural Venus Sinus Stenting for IIH
Suboccipital/Subtemporal Cranial Decompression

- Very invasive; but historically pretty successful...
- Remove part of skull to allow for more room inside...
- Not gold standard anymore
- Can be used in severely refractory cases unresponsive to traditional surgical procedures for ICP and IIH
Optometry IIH/PTC Summary...

- Make the diagnosis
- Get MRI/MRV
- Refer for LP
- Neurology should start Diamox/Topamax

- Monitor x 1 month post med Tx, then q3-4 months until resolution/stability (Varies)
- Serial OCT, FP, and HVF’s are necessary to gauge Tx/stability
- Relay findings to managing neurologist/PCP regularly.
- Encourage weight loss
What happened to patient?

- “No-showed” to neurologist twice!
- Has not returned phone calls or letters...
  - Phone has been disconnected...
- Possible candidate for DVSS? Diamox only? Both?
- “You cannot care more for a patient than what they care for themselves.” ----Joseph Sowka, OD and Alan Kabat, OD
References

Questions???

- Thanks!!!

- Email: cborgman@sco.edu