# **HINTS Exam: Determining the Cause of Acute Vestibular Syndrome**

# **Acute Vestibular Syndrome (AVS)**<sup>1,2</sup>

- Rapid onset of dizziness, nausea/vomiting, head motion intolerance, nystagmus, and unsteady gait
- Continuous dizziness for ≥ 24 hours
- Symptoms can last days to weeks

#### Causes of AVS

- Peripheral
  - o Acute peripheral neuropathy (APN) affecting CN VIII or the labyrinth
    - Vestibular neuritis or labyrinthitis
    - Trauma/surgical injury
    - Tumor, structural lesion (e.g. vestibular schwannoma, compression from tortuous vertebral artery)
- Central
  - o Posterior circulation stroke vertebrobasilar arterial system supplies the brainstem and cerebellum
    - PICA, AICA, Pontine arteries, SCA
    - 25% of patients in ED with AVS have posterior circulation stroke<sup>2-4</sup>
  - o Progressive neurologic conditions (e.g. MS, Friedreich's ataxia), tumor/structural lesion, paraneoplastic syndrome, vestibular migraine, medication reaction

### Posterior Circulation Associated with AVS

- Strokes in any of these territories can present as isolated AVS; patients can also have other neurological signs/symptoms
- Posterior Inferior Cerebellar Artery (PICA)
  - o From vertebral arteries supplies medulla and posterior inferior portion of cerebellum
  - o Most common artery for cerebellar infarcts
  - o Possible signs/symptoms: isolated AVS
  - Other possible neuro signs/symptoms: truncal ataxia<sup>5</sup>, dysarthria, dysphagia, contralateral sensory loss, decreased coordination<sup>6,7</sup>
- Anterior Inferior Cerebellar Artery (AICA)
  - o From basilar artery supplies pons, anterior inferior portion of cerebellum, and inner ear<sup>8</sup>
  - o Possible signs/symptoms: isolated AVS<sup>9</sup>
  - o Other possible neuro signs/symptoms: dysarthria, sudden hearing loss, ipsilateral CN V and VII involvement, dysmetria of UE more than LE, contralateral sensory loss<sup>5,9,10</sup>

- Labyrinthine Artery
  - o From AICA most commonly, or direct branch of basilar artery supplies inner ear
  - o Possible signs/symptoms: isolated AVS (if vestibular loss, peripheral pattern of nystagmus)
  - o Other possible neuro signs/symptoms: sudden hearing loss
- Pontine Arteries
  - o From basilar artery supply the pons
  - o Possible signs/symptoms: isolated AVS<sup>11</sup>
  - Other possible neuro signs/symptoms: motor involvement (UE>LE; distal >proximal), sensory involvement,<sup>31</sup> dysarthria, ataxia, CN V VII palsy<sup>10,11</sup>
- Superior Cerebellar Artery (SCA)
  - o From the basilar artery supplies the superior cerebellum, cerebellar white matter, midbrain, and pons
  - o Possible signs/symptoms: isolated AVS
  - o Other possible neuro signs/symptoms: dysarthria, diplopia, ataxia, contralateral loss of CN IV or pain/temp<sup>10</sup>

#### **Other Posterior Circulation Not Associated with AVS**

- Posterior Cerebral Artery (PCA)
  - o From the basilar artery supplies the occipital lobe, midbrain, thalamus, posterior inferior parietal lobe<sup>10</sup>
  - A stroke isolated to the PCA will not cause AVS
  - o It is possible to have multiple lesions sites in the posterior circulation. If there is a lesion in the PICA, AICA, Labyrinthine artery, Pontine arteries or SCA, in addition to the PCA, the patient could have S/S of AVS as well as the neurological S/S associated with the occipital lobe and midbrain.
  - Possible signs/symptoms associated with PCA: visual field loss, contralateral sensorimotor loss, CN III-IV palsy, ataxia, contralateral tremor, choreoathetosis, thalamic pain <sup>10</sup>

# **Diagnostic Tools** – HINTS exam is the best tool for ruling out central cause of AVS

- CT:
  - o Only 16% sensitivity for ischemic stroke<sup>8</sup>
  - o Good for tumors and hemorrhagic stroke, but only 4% of AVS causing strokes are hemorrhagic<sup>3</sup>
- Early MRI in patients with AVS and at least 1 stroke risk factor<sup>1</sup>
  - o 88% sensitive, 96% specific when done within 48 hours of symptom onset
  - o 80% sensitive, 97% specific when done within 24 hours of symptom onset
- HINTS exam<sup>1</sup>
  - o 3-part oculomotor exam: **H**ead **I**mpulse Test, **N**ystagmus, **T**est of **S**kew
  - o >96% sensitive and specific in patients with AVS and at least 1 stroke risk factor<sup>1</sup>
  - o Only for patients with AVS: dizziness present for ≥24 hours and nystagmus in at least 1 gaze evoked position
  - o Valuable exam during first few days of symptoms: ED, acute hospital, subacute setting if new symptoms arise

	Test Procedure	Stroke/Central Pathology	Unilateral Peripheral Pathology	Clarifying Information
<u>H</u> ead <u>I</u> mpulse Test	Patient keeps eyes on examiner's nose; passively rotate head back/forth at varying speed From ~ 20° rotation quickly rotate head back to midline	Normal HIT (no corrective saccade) or     Abnormal HIT (corrective saccade)	Abnormal HIT in only 1 direction	<ul> <li>Assesses the VOR</li> <li>Tests the side the head is rotated towards</li> <li>Abnormal HIT: eyes move in the direction the head was rotated requiring a corrective saccade to return to target</li> <li>Normal HIT in pts with AVS single best indicator of central pathology</li> </ul>
<b>N</b> ystagmus	With head stationary in midline the patient looks straight ahead, to the R and to the L for several seconds each	<ul> <li>Direction changing nystagmus (e.g., R-beating with R gaze and L-beating with L gaze)         or</li> <li>Direction fixed nystagmus</li> </ul>	Direction fixed nystagmus     Alexander's Law:     strongest when looking     in direction of the fast     beat & most     subtle/absent when     looking in direction of     slow beat	<ul> <li>Named for the direction of the fast phase</li> <li>Fast phase is away from the side of the lesion and must be in opposite direction of the abnormal HIT with a peripheral pathology</li> </ul>
<u>T</u> est of <u>S</u> kew	Alternate cover test; pt looks straight ahead, examiner covers 1 eye then covers the other eye	<ul> <li>Abnormal: Vertical refixation of at least 1 eye or</li> <li>Normal: No refixation</li> </ul>	No vertical realignment	<ul> <li>Imbalance of vestibular input to oculomotor system</li> <li>Least common central finding</li> </ul>
Interpretation of Results		Only 1 of the results in <b>bold</b> must be present to indicate a central pathology	All 3 must be present to indicate an unilateral peripheral vestibular pathology	

Dangerous HINTS = INFARCT (Impulse Normal, Fast-phase Alternating, Refixation on Cover Test)<sup>1</sup>

	Stroke/Central Pathology	Unilateral Peripheral Pathology
Actions Based on Results	<ul> <li>Document findings</li> <li>Discuss with MD</li> <li>Acute: MD may prescribe meds (vestibular suppressant or antiemetic)</li> <li>Patient education: stroke and treatment options</li> <li>Provide interventions as necessary for deficits</li> </ul>	<ul> <li>Document findings</li> <li>Discuss with MD</li> <li>Acute: MD may prescribe meds (vestibular suppressant or antiemetic)<sup>12</sup></li> <li>Acute/Chronic: Vestibular rehab as long as symptomatic<sup>12</sup></li> </ul>
	<ul> <li>Prognosis typically worse for those with central cause as compared to peripheral cause; ability for CNS to compensate may be reduced</li> <li>Possible referral to neuro-ophthalmologist or neuro-optometrist for persistent diplopia</li> <li>Possible referral for vehicle driving assessment if persistent ataxia, dynamic visual acuity deficit or diplopia</li> <li>Vestibular rehab may include same interventions as those for a peripheral cause<sup>13</sup></li> <li>Postural control, balance and gait exercises are important to improve visual and somatosensory input<sup>13,14</sup></li> </ul>	Components Peripheral Vestibular Rehab¹²  • Gaze stability exercises (e.g. VOR)  • Habituation exercises  • Balance/gait training  • Walking for endurance

#### Resources

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