## Title: Optic Nerve Disease Across Ages

- 1. Objectives:
  - I. Provide an overview of optic neuropathies that present in different age groups.
  - II. Review how to assess a patient with suspected optic neuropathy
  - III. Describe the different characteristics of the optic neuropathies and key features to recognize
  - IV. Discuss the different treatment/management for these conditions of the optic nerve
- 2. Optic Nerve Assessment
  - I. Case History
  - II. Entrance Testing
    - i. Visual Acuity
    - ii. Color Vision
    - iii. Pupil testing
  - III. Ocular Health Assessment
  - IV. Special testing
    - i. Threshold visual field testing
    - ii. Fluorescein angiography
    - iii. Electrodiagnostics
    - iv. Imaging Studies

## 3. Under 20yo

- I. Papillitis
  - i. Prevalence / Etiology
    - 1. Most cases are due to viral infections or post-immunization
  - ii. Characteristics
    - 1. Commonly associated with headache
    - 2. Children may not notice vision changes
    - 3. Vision changes are worse by days 2-3 to 2-3 weeks. Visual recovery starts soon thereafter.
    - 4. Majority of posterior segment findings involve only the optic nerve
  - iii. Treatment / Management
    - 1. Visual prognosis is good without treatment
- II. Leber's Hereditary Optic Neuropathy
  - i. Prevalence / Etiology
    - 1. Inherited from maternal mitochondrial DNA
      - a. All of the children of the mother will receive the trait, but only the female children are able to transmit the trait to the next generation.
    - 2. 9:1 (men over women)
    - 3. Age of onset is in an individual's teens or twenties.
  - ii. Characteristics
    - 1. Painless, vision loss with central scotoma are typically the first symptoms.

- 2. Initially present unilaterally but over a period of weeks to months becomes bilateral
- 3. Fundus findings include: circumpapillary telangiectasia but 1/3 will have normal ONH appearance.
- 4. Visual field defect typically a central scotoma
- III. Treatment / Management
  - i. None at this time
- 4. 20yo thru 40yo
  - I. Optic Neuritis associated with Multiple Sclerosis (MS)
    - i. Prevalence
      - 1. Young health adults
      - 2. Female > Male
      - 3. Incidence of MS associated with Optic Neuritis is highest in people living at higher latitudes (Northern US, Northern & Western Europe, New Zealand, and Southern Austrailia) and reduced significantly closer to the equator.
    - ii. Etiology
      - Is an inflammation of the optic nerve secondary to demyelinating Disease
      - 2. Other systemic conditions that can cause optic neuritis in this age group are: systemic lupus erythematosus, sarcoidosis, and syphilis.
        - a. Typically observe perineuritis and/or neuroretinitis
    - iii. Characteristics
      - 1. Sudden unilateral vision loss
      - 2. Pain or discomfort with eye movement (90%)
      - 3. Optic nerve appears normal in 2/3 of patients
      - 4. Optic disc edema in 20-40% of patients (\*does not correlate with severity of the disease process)
      - 5. Variety of visual field defects
      - 6. Uhtoff's sign worsening of symptoms when the body gets overheated
    - iv. Treatment / Management
      - 1. Optic Neuritis Treatment Trial
      - 2. CHAMPS Study
- 5. 40yo thru 60yo
  - I. Non-arteritic Anterior Ischemic Optic Neuropathy (NAION)
    - i. Prevalence
      - 1. Males = Females
    - ii. Etiology
      - 1. Inefficient blood supply to the ONH
      - 2. Prognosis > AION
    - iii. Characteristics
      - 1. Unilateral sudden painless (pain in 10%) vision loss

- 2. Dyschromotopsia
- 3. Altitudinal visual field defect
- 4. Crowded optic nerve appears swollen with sectorial hemorrhaging
- 5. Medical hx demonstrates h/o HTN, hypercholesterolemia, DM, obstructive sleep apnea, or other vascular risk factors.
- iv. Treatment / Management
  - 1. Manage the systemic condition
- 6. Over 60yo
  - I. Arteritic Anteror Ischemic Optic Neuropathy (AION)
    - i. Prevalence
      - 1. Females > Males
    - ii. Etiology
      - 1. Due to Giant Cell Arteritis (GCA)
      - 2. Strong association with Polymyalgia Rheumatica
    - iii. Characteristics
      - 1. Severe Vision Loss
      - 2. Jaw Claudication
      - 3. Scalp Tenderness
      - 4. Arthralgia
      - 5. Headache
      - 6. Amaurosis fugax
      - 7. Fever
      - 8. Weight loss
    - iv. Additional information
      - Diagnosis is made with laboratory testing: C-reactive protein / Westergren ESR.
      - 2. Temporal artery biopsy
    - v. Treatment / Management
      - 1. Systemic corticosteroids