

# OPTIC NEURITIS

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- An acute inflammatory disorder of the optic nerve
- Typically presents with sudden monocular visual loss and eye pain
- In young adults, more commonly in women
- A common initial manifestation of multiple sclerosis (MS)

# OPTIC NEURITIS

## CLASSIFICATION:

## ETIOLOGICAL:

- a. Idiopathic
- b. Demyelination
- c. Infections (viral) etc.
- d. Para infections/Post viral syndrome

- e. Toxic-Drugs
- f. Intraocular inflammations
- g. Contiguous inflammations (Sinus, Orbit)
- h. Systemic disease-sarcoid, T.B, Syphilis

❖ Para infectious-after viral infections, immunization

❖ Infections-viral infections, cat scratch fever, syphilis, Lyme disease

# **AIDS usually causes-Neuroretinitis**

- ❖ Drugs-Ethambutol, Isoniazid, Interferon, Chloramphenicol
- ❖ I.O. Inflammations: Uveitis, APMPE
- ❖ Sinus related-Headache + ENT opinion

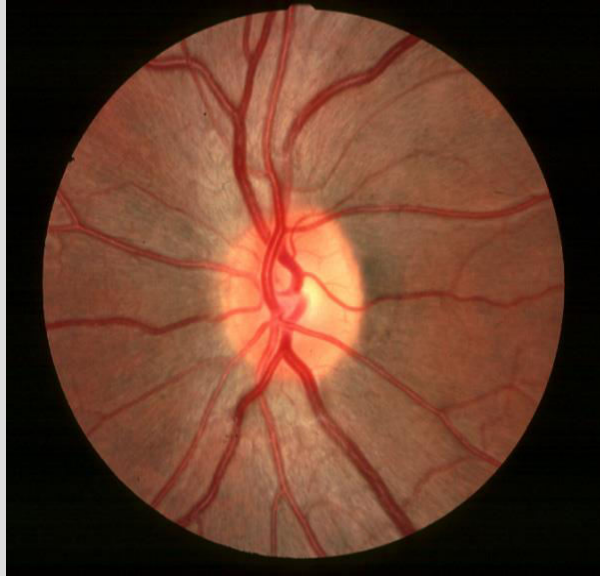
## **OPTIC NEURTIS**

### **CLINICAL TYPES**

- ❖ Retrobulbar (1/3rd cases)
- ❖ Papillitis (1/3rd cases)
- ❖ Neuroretinitis

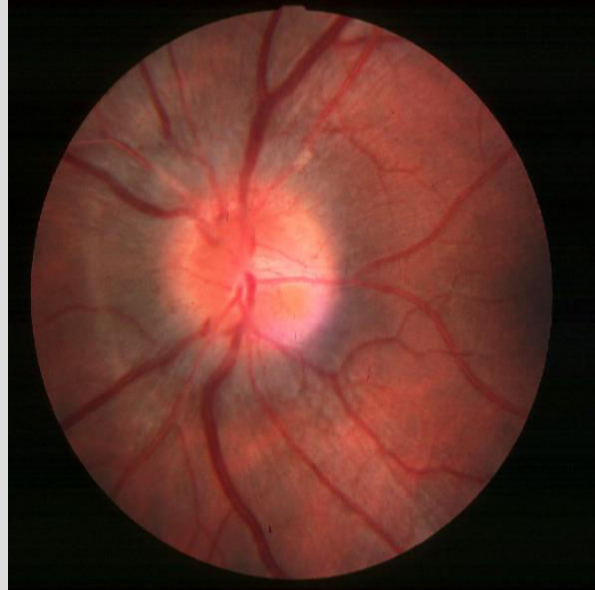
# Classification of optic neuritis

**Retrobulbar neuritis  
(normal disc)**



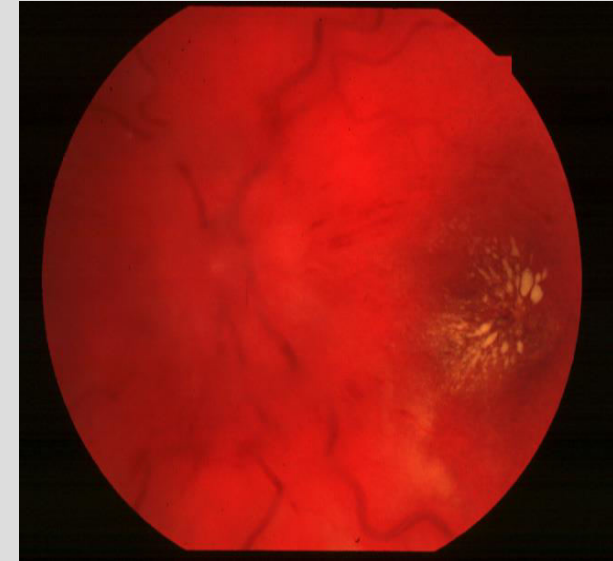
- **Demyelination - most common**
- **Sinus-related (ethmoiditis)**
- **Lyme disease**

**Papillitis (hyperemia and  
oedema)**



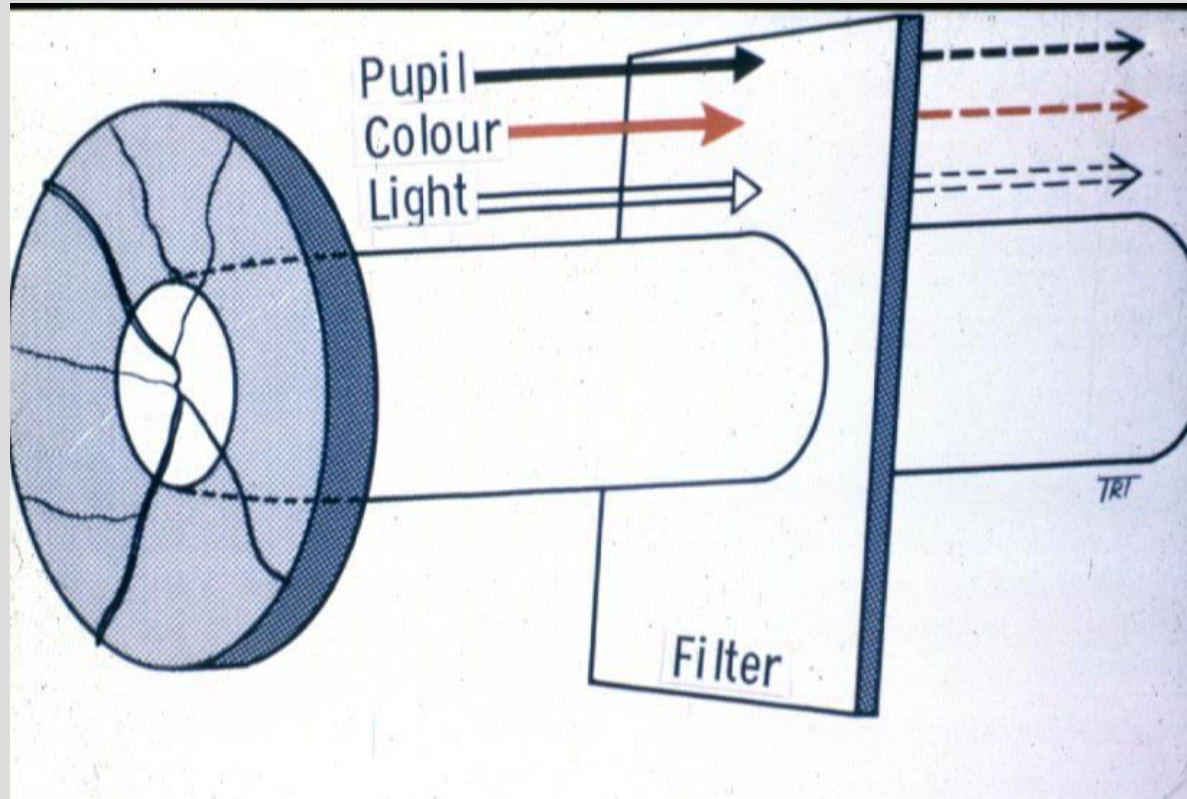
- **Viral infections and immunization in children (bilateral)**
- **Demyelination (uncommon)**
- **Syphilis**

**Neuroretinitis (papillitis  
and macular star)**



- **Cat-scratch fever**
- **Lyme disease**
- **Syphilis**

# Signs of optic nerve dysfunction



- **Reduced visual acuity**
- **Afferent pupillary conduction defect**
- **Dyschromatopsia**
- **Diminished light brightness sensitivity**

# SYMPTOMS: Triad

- ❖ Loss of vision
  - ❖ Ipsilateral eye pain
  - ❖ Dyschromatopsia
    - 70% - initially unilateral
    - 30% - 2nd eye may get involved
- (1) **Loss vision:** 58% - isolated symptom  
Range – 6/6 – PL-ve  
Vision decreases for 10-14 days
- Stabilize – Improvement (2-3 months)



**(II) PAIN:** Ipsilateral eye pain – 88%

with eye movements – 21% “Dull ache” in/behind the eye – 62% headache (involved region) 22% generalized headache- 13%

Pain – cause unknown with in 24 hrs. then recovers in 48-72 hrs.

O.N.T.T. (Pain) – 92% cases.

Persistent Pain 5 days – Atypical

**(III) DYCHROMATOPSIA:** Impaired color vision

Color desaturation – No macular lesions highly suggestive of O.N disease.

Color defect – worse than expected.

## **OTHER FEATURES:**

- 1) **RAPD:-** Ipsilateral – Neutral density filter
- 2) **CONTRAST SENSITIVITY:** Most sensitive (Even if V.A 6/6)
- 3) **VISUAL FIELD DEFCTS:** Central scotoma – may be altitudinal, arcuate, diffuse etc.
- 4) **PULFRCH EFFECT:** Pendulum movements appears elliptical when observed in front of eyes. [Delayed transmission + weak stimulation]

- 5) **UHTHOFF'S SYMPTOM:** episodic transit worsening of vision with exertion [ Exertion, hot food/drink, Tired end day ]  
Recovery – within 5-6 min/sometimes 2hrs. Pts with Uthhoff's –  
Higher incidence of MS, May be present in lebers optic N,  
Toxic optic N (Chloramphenicol).
- 6) **INVERSE UHTHOFF's SYMPTOM** – Improved vision with exercise, beer etc.
- .

7) Visual obscuration in bright light: O.N Pts vision in bright light – see better in dim light.

**8) MOVEMENT PHOSPHENES:**

**Phosphenes:** Seeing brief flash of light (1-2 sec). Almost exclusively and horizontal eye movements better seen in dark/dim room with closed eyes ipsilateral and unilateral suggests demyelination

**9) SOUND PHOSPHENES:** Produced by sudden noise when pt. is resting in the dark. May occur in optic neuritis or compressive neuropathy

**PROGNOSIS:** 75 – 6/9 OR BETTER V.A

- 85% 6/12 OR BETTER V.A
- Color vision, contrast sensitivity, light brightness appreciation often
- remain abnormal. Mild RAPD may stay and optic atrophy may start.

**OPTIC NEURTIS TREATMENT TRIAL:**

- Multicentered, Randomized, Prospective clinical trial – 457 cases of
- O.N (81-45 years- Acute O.N for 8 days + visual)
- Field defects + RAPD – Included (Typical)

# OPTIC NEURITIS TREATMENT TRIAL

- I): Oral prednisone – 1 mg/kg/day – 14 days
- II): I/V Methyl prednisolone (1000 mg/day) for 3 days f/b oral prednisone (1mg/kg/day) for 11 days.
- III): Oral placebo for 14 days.

# OPTIC NEURITIS TREATMENT TRIAL

## Results

- ❖ Oral steroids – recurrence of O.N in affected or contralateral eye.
- ❖ I/V steroids → Fast recovery of vision for the first year but then after no difference.

## **TREATMENT:**

- ❖ Mild cases – No treatment
- ❖ Cases with V.A > 6/12 I/V steroids f/b oral steroids – speed up recovery and lower the incidence of MS in first two years.

No long-term benefit on the final V.A.

## **WORKUP**

- All patients with O.N – MRI – if normal no further work.
- MRI show 2 or > 2 typical lesion then I/V steroids f/b oral may incidence of MS in first two years.



## **ASSOCIATION OF MS & OPTIC NEURITIS:**

- 1) 15 – 20% pts with MS will present with O.N.
- 2) 35 – 40% pts with MS will develop optic neuritis during course of their disease.
- 3) 74% female and 34% male with optic neuritis will develop MS when followed up to 15 years

- 4) 50% -70% of clinically isolate optic neuritis have abnormal MRI similar to that seen in MS.
- 5) Risk of MS is increased in pts of optic neuritis when there is winter onset, HLA DR2 positively and Uhthoff's phenomenon.
- 6) 36 eyes of 18 pts – when alive only 8 of 18 pts had diagnosis of unilateral or bilateral optic neuritis.

On Autopsy 35 eyes out of 36 eyes showed evidence of demyelination.

**Thank You**