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# **Ocular Toxoplasmosis**

- Caused by Toxoplasma gondii
- Obligate intracellular protozoan
- 500 million have antibodies worldwide
- 50% of adult population in US have asymptomatic infection
- 28-55% of all cases of posterior uveitis
- First isolated from the brain of a "gondii" (North African Rodent)
- Cat family is definitive host, can infect other mammals and birds
- Oocysts found in intestinal tracts of cats
- Cysts ingested (most likely)
- Poor hygiene, infected pork, chicken but probably not beef
- Can survive outside host for up to 1 year
- Two forms in humans: cysts or tachyzoites.
- Propensity for cardiac and skeletal muscle and neural tissue (brain & eye)

# **Clinical Manifestations**

### Acquired vs. Congenital

- Most cases are presumed reactivation of congenital infection
- 2 to 6/1000 women acquire infection while pregnant, 40% risk of transmission to fetus.
- Of infected infants: 70% chorioretinal scars, 5% will die or severe disability, 1-2% severe visual impairment
- Northern Brazil has high rate of acquired disease

## **Systemic**

- 90% lympadenopathy
- · fever, malaise, occasional sore throat
- immunocompromised- fulminant CNS disease

### **Ocular**

- Keratic Precipitates, anterior chamber cell and flare, posterior synechia, cataract
- Retinochoroiditis
- Vitritis- concentrated over lesion, scaffolding of vitreous strands
- Macular edema
- · Retinal vasculitis
- VF defect in area of scar
- FFA of active lesion- early blockage and subsequent leakage

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# **Atypical Presentations**

- In early infection: gray-white fine punctate lesions in deep retina and RPE, progress to more classic lesions
- Papillitis, vitreal inflammation, nerve fiber bundle defects
- Bullous like inflammatory lesions in mid-periphery
- Wide ring-like lesion near extreme periphery resembling pars planitis
- Scleritis

### **Reasons for Vision Loss**

- Vitreal inflammation causing clouding
- Lesion in posterior pole with edema affecting fovea
- · Lesion in fovea
- Subsequent CNVM

# **Diagnosis**

- Typical lesions
- Toxoplasmosis titers are supportive
  - IgM titers- can be missed
  - IgG titers- high rate of false positives
- Immunoflourescence, ELISA
- Western blot for Toxo antigens
- PCR and Southern Blot for Toxo DNA
- Angiography: flourescein tagged Ab (successful in rabbit studies)

# **Therapy**

- Should you treat it at all?
  - Lesion within temporal arcade
  - Lesion next to optic nerve or large vessel
  - Lesion has induced large degree of hemorrhage
  - Vision drop of > two lines
  - Multiple recurrences with vitreal contraction
- No truly randomized, controlled clinical trials to compare efficacy
- Generally 4-6 weeks, multi-drug regimens

## **Medications**

- Sulfadiazine 1g PO QID
- Pyrimethamine 75-100mg load and 25-50mg PO BID (bone marrow suppression, nausea)

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- Use concurrently: Folinic Acid 3-5mg PO 3 times/week (Baseline CBC, follow q week)
- Clindamycin 150-300mg PO TID-QID (?reduces recurrence, risk of pseudomembranous colitis)
- Trimethoprim-sulfamethoxazole (DS) I PO BID
- Atovaquone (Mepron) 750mg PO BID (kills cysts in vitro)
- Tetracycline 2g load and 250mg PO QID (to replace clindamycin)
- Prednisone 20-60 mg/day not used alone

# Rothova et.al. (The Netherlands) Am J Ophth, 115:515-523. April, 1993

### Treatment Regimen #1

- Pyrimethamine, Sulfadiazine, Folinic Acid, Prednisone
- Best at reducing size of lesions (49%),
- more medication side effects (26%),
- recurrence rate at 3 years (42%)

### **Treatment Regimen #2**

- Clindamycin, Sulfadiazine, Prednisone
- Reducing size of lesions >1/2 DD (28%)
- med side effects (17%)
- recurrence rate at 3 years (67%)

### **Treatment Regimen #3**

- Co-trimoxazole, Prednisone
- Reducing size of lesions >1/2 DD (11%)
- med side effec!ts (4%)
- recurrence rate at 3 years (40%)

### **Treatment Regimen #4**

- No treatment (peripheral lesions)
- Reducing size of lesions >1/2 DD (20%)
- no med side effects
- recurrence rate at 3 years (53%)

#### **Comments**

- Recurrence rates not statistically significant
- Size of retinal lesion correlated with duration of inflammation
- All side-effects were reversible
- Delay in starting medication (even up to 1 week) did not alter duration of inflammation
- Sub-tenon's steroid- risk of increasing activity of organism
- Vitrectomy- for vitreal haze reducing vision, perioperative antibiotics advocated
- Cryotherapy and Laser photocoagulation generally not successful

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