بسم الله الرحمن الرحيم

Treatment of infections of peripheral nervous system

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Treatment of ophthalmic Infections

1- Herpes zoster ophthalmicus

- It is a reactivated latent <u>varicella-zoster virus</u> (VZV) infection involving the eye (<u>eyelid</u>, <u>cornea</u>, <u>conjunctiva</u>) & the ophthalmic division of the <u>trigeminal nerve</u> (V).→wtreated→death
- Itreatment should be <u>started as early as possible</u> (within 72 hours but up to 7 days after onset of rash). >early is better
- ☐ Oral aciclovir (800mg 5 times per day) or oral valaciclovir

 1g three times per day for 7 days duration.

N.B: Topical anti-viral drugs are NOT routinely required.

□If recommended by ophthalmologist: Ganciclovir eye gel 5 times daily (risk of teratogenicity).

2- Blepharitis (eyelid bacterial infection)

Treatment:

- 1- Good hygiene and treating associated seborrheic dermatitis or rosecia.
- 2- Topical Chloramphenicol or fusidic acid > kills gram +ve

3- For severe cases oral azithromycin.

3- Corneal Ulcers (Microbial Keratitis)

Empirical Treatment:

Ulcer <1mm: Ofloxacin eye drops

Ulcer ≥1mm: Gentamicin + Cefuroxime eye drops or Ofloxacin eye drops

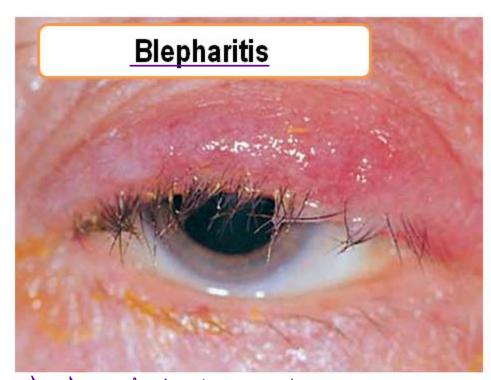
Specific Treatment:

- □<u>Bacterial</u> (according to <u>culture</u> and <u>sensitivity</u>)
- Herpes simplex keratitis (dendretic ulcer) Ganciclovir eye gel 5 times daily until healing then 3 times daily for 7 days (Maximum duration 2 weeks).

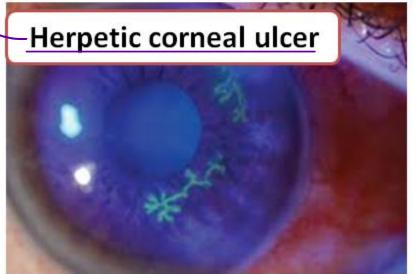
->nerve-shape

□ Fungal : Amphotericin eye drops.

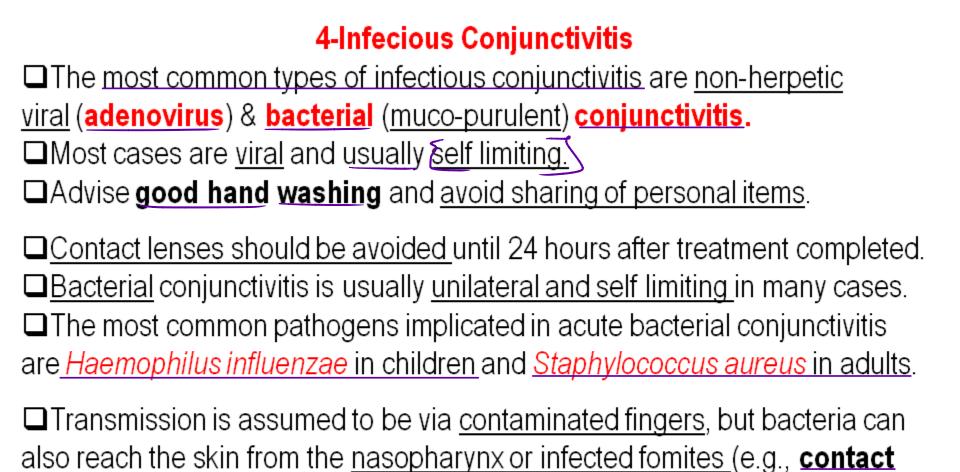




>recurrence > First 48-hours -> Acyclorir-> kill virus and prevent reactivation it







The use of topical steroids for the treatment of bacterial conjunctivitis should be avoided as there is an increased risk of potentiating the infection and prolonging the duration of the disease.

<u>lenses, makeup</u>).

71st line Treatment of bacterial conjunctivities

- (1- Eye hygiene: <u>washing regularly</u> to <u>remove crusts and pu</u>s.
- 2-Chloramphenicol eye drops or ointment applied four times daily.
- For severe infections Chloramphenicol eye drops every 2 hours initially then reduce frequency 4 times daily.
- □Continue treatment for 48 hours after resolution.
- ☐ Treatment should not exceed 5 days, to avoid adverse effects of
- Chloramphenicol (optic neuropathies, blood dyscriasis and aplastic anemia).
- 3- If <u>no response to Chloramphenicol</u> (bacteriological investigation should be requested); one of the followings can be prescribed:
- Fusidic acid eye drops twice daily (for gram positive only).
- Aminoglycoside eye drops or ointment (e.g. tobramycin, neomycin) for gram negative only (like pseudomonas)
- Both <u>Fusidic acid</u> and <u>aminoglycoside</u> causes <u>burning sensation</u> and <u>hypersensitivity</u>.

3-Fluoroquinolone eye drops (ciprofloxacin and gatifloxacin) have a broad spectrum of activity, however they are not first line agents due to the possibility of resistance developing.

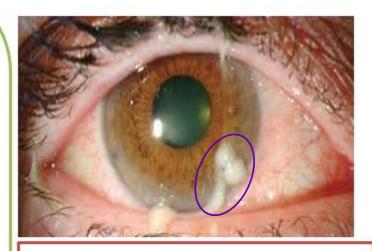
Adverse effects are common and include burning, eye discomfort, conjunctival hyperaemia, crusting, and a bad taste. In rare cases allergic reactions, lid edema, keratitis, light sensitivity, tearing and visual disturbances.

4- Bacitracin/polymyxin B eye ointment

- Stinging and <u>burning</u> have been reported after administration.
- Continue treatment for 48 hours after resolution.
- Finally: Consider <u>Chlamydia trachomatis</u> or <u>gonococcal infection</u> in <u>sexually active patients</u> (<u>need specific oral treatment</u>).

Instructions for applying eye drops

- □ Place one drop at a time at the **lower** conjunctival sac followed by closure of the eyelids for several seconds.
- □If more than one medication is being applied, wait 3 to 5 minutes between applications.
- Excessive blinking and placing more than two drops at a time, regardless of the medication type, can cause spillage and waste.
- □Contact lenses should be avoided during symptom flare-ups. However, if wear is necessary, drops should be administered 15 minutes before lenses are placed or after lenses are removed.



Mucopurulent conjunctivitis



Chlamydial conjunctivitis (Chlamydia trachomatis)

- 1. Inclusion conjunctivitis (in adults and neonates)
- Irachoma
- □ Trachoma predominantly effects children and occurs mostly in developing countries with low socioeconomic status and poor hygiene.
- □It is caused by Chlamydia trachomatis (subtypes A–C) and can result in blindness if left untreated.
- > Adults: >in adult
- Oral <u>azithromycin</u> (<u>single 1 g oral dose</u>) or <u>Oral doxycycline</u> (21–28 days) or <u>Oral erythromycin for</u> (21–28 days).
- > Children: > = hildren
- Oral azithromycin (20 mg/kg maximum 1 g) single dose.
- ➤ Pregnant women, infants < 6 months or patients allergic to macrolides:
- Topical tetracycline (eye ointment 2 times daily for 6 weeks)

Trachoma







Eyelid scarring

It is a bacterial
eye infection.

It is the main cause
of irreversible
blindness due to
infection that
can however
be prevented.

Normal eye

Eyelid Inflammation

Antibiotics for trachoma



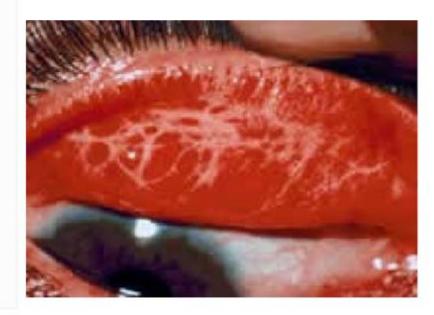
Azithromycin tablets



Azithromycin paediatric oral suspension



Tetracycline eye ointment



Drugs used in treatment of leprosy

1- Dapsone & other sulfones

- Dapsone (diaminodiphenylsulfone) is closely related to the sulfonamides.
- ► It inhibits folate synthesis.
- Resistance can emerge in lepromatous leprosy, if low doses are given.

 inhibit Folic acid synthesis > sultonamide in lepromatous leprosy

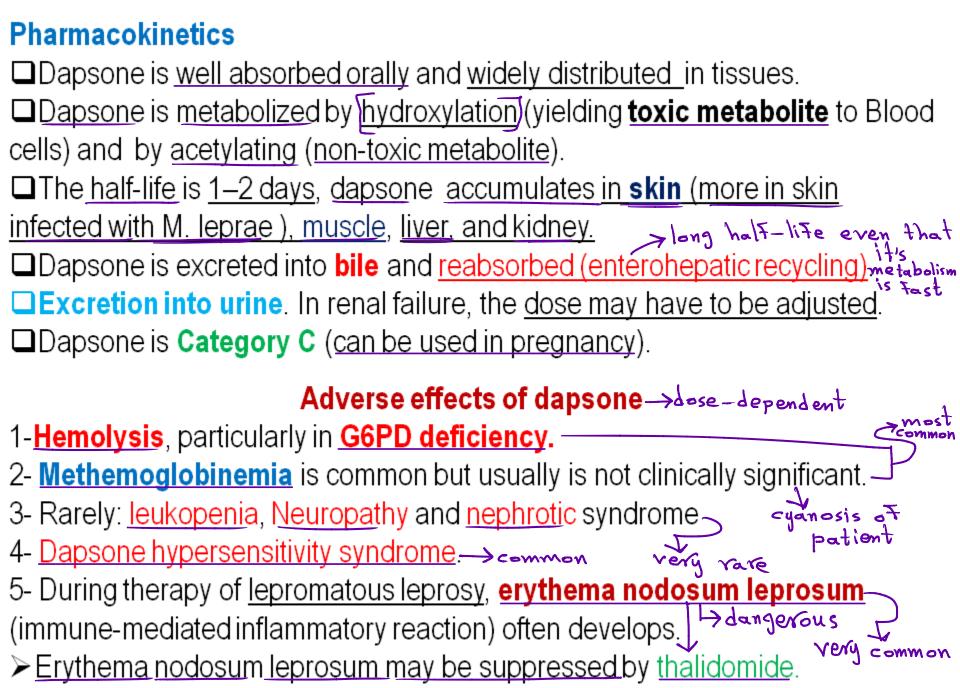
inhibit Folic acid synthesis sulfonamide in lepromatous

Therefore, the combination of dapsone, rifampin and clofazimine is recommended for initial therapy of lepromatous leprosy. >multi-bacillary

> Can't given alone > resistance

> A combination of dapsone plus rifampin is commonly used for leprosy

- with a lower organism burden (pausibacillary).
- ➤ Dapsone may also be used to <u>prevent and treat Pneumocystis jiroveci</u> pneumonia in persons with AIDS and other types of immunocompromise.
- Dapsone is used also for treating other conditions including dermatitis herpitiformis, acne vulgaris and neutrophilic dermatoses including behcet's disease.



2- Rifampin

- Rifampin (U.S) or <u>rifampicin</u> (<u>Europe</u>) is one of the Rifamycins (drug group which includes <u>Rifampin</u> and <u>rifabutin</u>)
- Rifamycins <u>Inhibit</u> <u>RNA polymerase</u> in the <u>bacteria</u> leading to inhibition of RNA formation.
- Clinical uses of rifampin
- 1- Treatment of Mycobacterium tuberculosis infection (used in combination with other drugs). Monotherapy rapidly leads to resistance.
- 2- Treatment of <u>leprosy</u> (Rifampin delays resistance to dapsone when used for leprosy).
- 3-Used for meningococcal prophylaxis and chemoprophylaxis in contacts of children with H. influenzae type b.

Adverse effects of rifampin

- 1-Minor hepatotoxicity. However, serious hepatic injury rarely occur.
- 2-Drug interactions: Rifampin <u>induces</u> <u>cytochrome</u> <u>P-450</u> <u>leading</u> to <u>accelerated metabolism</u> of <u>other drugs including warfarin</u>, <u>hormonal contraceptives</u>, theophyllin and others.

N.B: Rifampin causes nonhazardous orange body fluids.

- ➤ Rifampin may change the color of urine, sweat, saliva, or tears to (yellow, orange, red, or brown).
- This <u>effect</u> is <u>harmless</u> and <u>will disappear</u> when the <u>medication</u> is <u>stopped</u>. However, teeth and contact lens staining may be permanent.



Remember Rifampin's 4 R's

- 1. RNA polymerase inhibitor.
- 2. Ramps up microsomal cytochrome P-450.
- 3. Red/orange body fluids
- 4. Rapid resistance if used alone needs combination

N.B. Rifapentine is an antibacterial drug similar to Rifampin but has longer half-life than

Clofazimine

- □Clofazimine is a <u>bhenazine dye</u> used for treating <u>multibacillary leprosy.</u>
- □Its mechanism of action has not been clearly established.
- □ Absorption of clofazimine from the gut is variable, and a major portion
- of the drug is excreted in feces. > no risk in kidney or liver
- □Clofazimine is stored widely in reticuloendothelial tissues and skin, and
- its <u>crystals</u> can be seen inside <u>phagocytic reticuloendothelial cells</u>
- □ It is slowly released from these deposits, so the serum half-life may be

2 months.

☐ The most prominent adverse effect is discoloration of the skin and conjunctivae. Gastrointestinal side effects are common.

