

## Chlamydia & Warts Treatment

Drugs	MOA	Dosage	Uses	Side effect
<b>Treatment of Chlamydia</b>  <b>a. Doxycycline (Vibramycin) (cure rate: 98%)</b>	A tetracycline antibiotic. <b>MOA:</b> protein synthesis inhibitor.  <b>1st choice.</b>	<b>Dosage:</b> 100 mg po twice a day for 7 days for urogenital chlamydial infection.	<b>Uses:</b> chronic prostatitis, sinusitis, syphilis, Chlamydia, PID, acne, & Rickettsial infections.	Nausea vomiting & diarrhea. Permanent teeth staining & bone abnormalities Overgrowth of fungal infections in the GIT due to inhibition of normal flora. <b>contraindicated</b> during <b>pregnancy</b> & <b>breast feeding</b> .
<b>b. Azithromycin (Zithromax) 97% extensive Bacterostatic</b>	Macrolide, inhibits RNA-dependent <b>protein synthesis</b> of the bacteria by binding to the ribosomal subunit <b>50S rRNA</b> . <b>Uses:</b> middle ear infections, tonsillitis, throat infections, laryngitis, bronchitis, pneumonia, typhoid & sinusitis. Urethritis (STD), Chlamydia & gonorrhea & cervicitis. Chronic persistent genital chlamydia.	<b>Orally</b> easy absorption → need 2-3 hrs to peak conc. The conc. in tissues can be 50 times higher than in plasma. This is due to ion trapping & the high lipid solubility (Volume of distribution is too low). Following a single 500 mg dose → $T_{1/2}$ → 11-14 hrs (long). 35% of <b>azithromycin</b> is metabolized in the liver, excreted in bile & unchanged in urine <b>Dosage:</b> 1 g in a single dose..	<b>Uses:</b> chronic prostatitis, sinusitis, syphilis, Chlamydia, PID, acne, & Rickettsial infections.	<b>Contraindication:</b> hypersensitivity to macrolide antibiotics.  <b>Pseudomembranous colitis</b> occur during or after therapy. Interfere with control pills. Has bad taste → children. I.V over dose → heart block → encephalopathy.
<b>Alternative regimens:</b> <b>Erythromycin (95% efficacy):</b> <b>Cheap, used in pregnancy disadvantages: 4 times daily &amp; causes GIT disturbances.</b>	Macrolide antibiotic. <b>Used</b> for people who have an allergy to penicillins. <b>Dosage:</b> 500 mg 3 times daily for 7 days.	Displays bacteriocidal activity, in particular, at higher concentrations. <b>Inhibits protein synthesis</b> & subsequent structure & function processes critical for life or replication of the bacteria by binding to the <b>50S subunit</b> of the bacterial rRNA complex. This interferes with the production of functionally useful proteins.	Orally will be inactivated by gastric acid, so given as enteric-coated or more-stable salts or esters, such as erythromycin ethyl succinate. Very <b>rapidly absorbed</b> , & diffuses into <b>most tissues</b> & <b>phagocytes</b> . Due to the high conc. in phagocytes, <b>erythromycin</b> is actively transported to the site of infection, where, during active phagocytosis, large conc. of <b>erythromycin</b> are released. Most of erythromycin is metabolized by demethylation in the liver. Elimination in bile. $T_{1/2}$ → 1.5 hours.	<b>Contraindication:</b> hypersensitivity to macrolide antibiotics.  <b>Pseudomembranous colitis</b> occur during or after therapy. Interfere with control pills. Has bad taste → children. I.V over dose → heart block → encephalopathy.

<b>Treatment of warts</b> <b>Acyclovir (Aciclovir)</b>	A guanosine analogue antiviral drug used for <b>HSV 1 &amp; 2</b> infections, as well as in the treatment of herpes zoster (shingles). extremely selective & low in cytotoxicity.	MOA: It is selectively converted into acyclo-guanosine monophosphate ( <b>acyclo-GMP</b> ) by viral thymidine kinase. Acyclo-GTP is a very potent inhibitor of viral DNA polymerase; acyclo-GTP is incorporated into viral DNA, resulting in chain termination.	Has poor oral bioavailability (15-30%), hence IV administration is necessary if high conc. are required. When orally administered, peak plasma conc. occurs after 1-2 hours. Aciclovir has a high distribution rate; only 30% is protein-bound in plasma. Administered as oral tablet, cream & IV injection. The elimination half-life of aciclovir is approximately 3 hours. It is renally excreted.	NSV, diarrhea & headache. In high doses, <b>hallucinations</b> .
<b>Famciclovir</b>	A guanine analogue antiviral drug used for the treatment of various <b>herpesvirus infections</b> , most commonly for herpes zoster. It is a prodrug form of penciclovir with improved oral bioavailability.	Used for the treatment of herpes simplex virus 2 (genital herpes) & orolabial herpes (cold sores). Suppression of recurring episodes of HSV2. <b>Treatment of recurrent</b> episodes of HSV in HIV patients.	Mild to extreme stomach upset, headaches, mild fever.	

Medical Treatment of Warts	Preparation & MOA	ADRs
<b>Podophyllin resin (Podofin) &amp; Podofilox (condylox)</b>	both applied topically <b>Have antimitotic activity, destroy warts, &amp; inexpensive.</b>	<b>Local irritation &amp; pain.</b>
<b>Trichloroacetic acid (TCA) or bichloroacetic acid</b>	topically applied:	<b>Incomplete response, recurrence is high &amp; may cause pain &amp; burning</b>
<b>5-Fluorouracil</b>	applied as a cream ( <b>antimetabolite; anticancer</b> )	<b>Has long treatment time, can cause burning &amp; irritation.</b>
<b>Imiquimod</b>	Cream. It is an <b>immune enhancer</b> , stimulate production of interferon & other cytokines.	<b>Local skin irritation.</b>
<b>Other Type of Treatment</b> <b>Cryotherapy</b>	Freezing the wart by liquid nitrogen or cryoprobe. Used as <b>first line</b> treatment because response rates are high with <b>few side effects</b> .	It is not <b>recommended to treat vaginal</b> warts because of the risk of vaginal perforation.
<b>Laser</b>	Used for <b>larger</b> & extensive or <b>recurrent</b> genital warts. The laser destroys HPV-induced lesion.	Disadvantages: high cost, ↑healing time, & scarring.
<b>Surgery</b>	Cutting the warts away with local anesthesia. Usually done when the warts are small in size & number.	

#### Treatment of warts in pregnancy:

Liquid nitrogen cryosurgery is safe for pregnancy.

Topical TCA (**Trichloroacetic acid**) for vagina can be applied.