1998 Guidelines for Treatment of Sexually Transmitted Diseases

Each year there are an estimated 12 million new STD infections in the United States; about 3 million of these cases occur in teens. By the time they reach adulthood, 1 in 4 teens will have had an STD, which may include herpes and/or human papilloma virus (HPV), two of the most common viral STDs for which there is no known cure.

In its 1997 report, *The Hidden Epidemic: Confronting Sexually Transmitted Diseases,* the Institute of Medicine (IOM) cited a lack of appropriate screening and treatment of STDs as a factor contributing to the STD epidemic in the United States. The IOM panel concluded that patients with diagnosed STDs often receive treatment not commensurate with current standards of care.

The Centers for Disease Control and Prevention (CDC)—in conjunction with a group of nationally recognized STD experts from public health, academia, medical research, and managed care organizations—reviewed scientific literature and clinical practice data to develop the *1998 Guidelines for Treatment of Sexually Transmitted Disease*. The CDC guidelines contain recommendations for quality of care and outcomes of STD therapy: cure, relief of signs and symptoms, prevention of complications, and prevention of further transmission. The 1998 version replaces the 1993 guidelines.

These guidelines are intended for use by health care providers, trainers, educators, researchers, and others in primary care, adolescent care, family medicine, family planning, internal medicine, obstetrics-gynecology, urology, dermatology, emergency care, nursing, and HIV care. Early, effective STD treatment can also substantially reduce HIV transmission. Effective STD detection and treatment provides the greatest health benefits for women, particularly adolescent and young adult women, and their babies. However, people of both sexes and all ages benefit, not only in terms of increased health/ reduced risk, but also in terms of lower health care costs-for the individual as well as the society as a whole.

The CDC guidelines provide basic information for detecting and treating many "silent" STDs which can be difficult to diagnose because they frequently have no symptoms, or symptoms that are very vague or easy to confuse with other disorders. A "silent" STD such as chlamydia can unknowingly be transmitted to partners and can have major consequences in women when not diagnosed and treated. Complications associated with chlamydial infection include pelvic inflammatory disease (PID), potentially fatal tubal pregnancy, infertility, and poor birth outcomes. Chlamydia and certain other STDs can also put patients at greater risk for acquiring and transmitting HIV.

The guidelines include diagnosis and treatment information for all common STDs and are organized by syndrome: STDs characterized by genital ulcers, by urethritis and cervicitis, and by vaginal discharge. Also included are recommendations for STD prevention as well as special considerations for three highrisk populations: women, adolescents, and infants. Finally, the guidelines include sections on other problems that occur among patients with STDs: PID, epididymitis, patients with penicillin allergy, sexual assault issues, and cervical cancer screening.

Significant medical advances have been made since CDC released its 1993 guidelines. Among the most notable advances described in the 1998 guidelines are the following:

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Texas Department of Health

Highly effective single-dose oral therapies have been developed for almost all common curable STDs. Simplifying the treatment for common STDs, such as chlamydia, to one dose of medicine given at the time of examination can significantly increase the number of people who are treated and cured. Single-dose therapies help prevent the serious reproductive outcomes caused by these STDs and shorten the period of infectiousness.

Improved treatments are now available for herpes and HPV. Advances in the treatment of the viral STDs are critical: over 45 million Americans (about 1 in 5) are now infected with herpes type 2, and human papilloma virus (HPV) is a very common STD associated with cervical cancer and genital warts. Patientapplied treatment for HPV makes it easier for patients to comply with prescribed therapy when symptoms occur. More effective treatments for genital herpes help to alleviate symptoms, reduce the emotional stress associated with viral STDs, and possibly reduce transmission.

The introduction of a simple urine test has made it much easier to diagnose and treat chlamydia and gonorrhea in clinical and nonclinical settings. Recent research advances have led to extremely accurate urine tests that make testing of both women and men simple and comfortable. In addition, chlamydia and gonorrhea testing may be conducted in nonclinical settings for large groups of adolescents who are at greatest risk. In high school and other community screening programs, as many as 1 in 8 teenage girls test positive for chlamydia.

Vaccination is recommended as the most effective means of preventing hepatitis A and hepatitis B infections that are sexually transmitted.

Inactivated hepatitis A vaccines, administered as a 2-dose series, are safe and highly immunogenic. Men who have sex with men are at highest risk for sexually transmitted hepatitis A. Sexual transmission accounted for 30% to 60% of the estimated 240,000 new hepatitis B virus (HBV) infections per year in the United States. Multiple age groups must be targeted to effectively prevent HBV transmission and HBV-related chronic liver disease.

Significant advances have been made in STD treatment during pregnancy. New treatments for chlamydia produce fewer side effects. Improved treatments for STDs in pregnancy may produce fewer side effects and reduce the number of infants born prematurely. Also, new recommendations that focus on screening for and treating bacterial vaginosis among women at high risk (those with a previous history of preterm birth) will likely reduce the number of premature births due to this STD.

To improve STD care in the United States, IOM has called for wider dissemination of the CDC treatment guidelines, and for all primary care providers (including managed care organizations and their health plans) to implement these recommendations. The *1998 Guidelines for the Treatment of Sexually Transmitted Diseases* are available at the CDC website: <u>www.cdc.gov/nchstp/dstd/</u>.

Health professionals can also obtain copies (free of charge) from the Texas Department of Health (TDH) by sending a written request for *stock number 6-110* to the TDH Warehouse, Literature and Forms, 1100 W. 49th Street, Austin, Texas 78756.

TDH staff are available to assist health professionals with reporting regulations and confidential sex partner referral, as well as provide consultation for the treatment of patients diagnosed with or exposed to an STD/HIV.

Call (512) 490-2500 for more information about STD diagnosis, treatment, reporting and prevention.

1998 Quick-Reference Treatment Guide: Selected Sexually Transmitted Diseases

The Texas Department of Health Bureau of HIV/STD Prevention prepared the following quick reference chart for sexually transmitted diseases commonly treated in an outpatient setting. These treatment guidelines are excerpted from the Centers for Disease Control and Prevention *1998 Guidelines for Treatment of Sexually Transmitted Diseases*. It is not an exhaustive list of effective treatments, nor are these recommendations to be construed as inflexible standards or rules.

Disease	Recommended Treatment	Alternative	
Bacterial Vaginosis			
	 Metronidazole 500 mg PO BID for 7 days OR Clindamycin cream 2% one full applicator (5g) intravaginally at bedtime for 7 days OR Metronidazole gel 0.75% one full applicator (5g) intravaginally BID for 5 days 	 Metronidazole 2 g PO single dose OR Clindamycin 300 mg PO BID for 7 days 	
High-Risk Preg	nancy (Premature Delivery) • Metronidazole 250 mg PO TID for 7 days	• Metronidazole 2 g PO single dose OR • Clindamycin 300 mg PO BID for 7 days	
Chancroid ^{1,2}			
	 Azithromycin 1 g PO single dose OR Ceftriaxone 250 mg IM single dose OR Ciprofloxacin 500 mg PO BID for 3 days OR Erythromycin base 500 mg PO QID for 7 days 		
Chlamydial Infe	ections		
Adults ^{2,3}	 Azithromycin 1 g PO single dose OR Doxycycline 100 mg PO BID for 7 days 	 Erythromycin base 500 mg PO QID for 7 days OR Erythromycin ethylsuccinate 800 mg PO QID for 7 days OR Ofloxacin 300 mg PO BID for 7 days 	
Children		a series and a series of the s	
<45 kg	• Erythromycin base 50 mg/kg/day PO divided into 4 doses daily for 10-14 days ⁴		
≥45 kg and <8 yrs old • Azithromycin 1 g PO single dose			
\geq 8 yrs old	 Azithromycin 1 g PO single dose OR Doxycycline 100 mg PO BID for 7 days 		
Pregnancy	 Erythromycin base 500 mg PO QID for 7 days OR Amoxicillin 500 mg PO TID for 7 days 	 Erythromycin 250 mg PO QID for 14 days OR Erythromycin ethylsuccinate 800 mg PO QID for 7 days OR Erythromycin ethylsuccinate 400 mg PO QID for 14 days 	
Epididymitis			
	For epididymitis caused by GC or CT: • Ceftriaxone 250 mg IM single dose PLUS • Doxycycline 100 mg PO BID for 10 days	For epididymitis caused by enteric organisms or for patients allergic to recommended treatment: • Ofloxacin 300 mg PO BID for 10 days	

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Disease	Recommended	Treatment	Alternative	
Gonococcal Info	ections ^{2,3,5}			
Adults and Chi Cervix, Uret	ldren ≥45kg thra, Rectum			
	 Cefixime 400 m Ceftriaxone 123 Ciprofloxacin 5 Ofloxacin 400 n PLUS* Azithromycin 5 	ng PO single dose OR 5 mg IM single dose OR 500 PO single dose OR ng PO single dose 1 g PO single dose OR	• Spectinomycin 2 g IN	1 single dose ⁶
	• Doxycycline 10	0 mg PO BID for 7 days		
Pharynx	• Ceftriaxone 123 • Ciprofloxacin 5 • Ofloxacin 400 PLUS*	5 mg IM single dose OR 600 mg PO single dose O mg PO single dose	R	
	 Azithromycin 1 Doxycycline 10 	l g PO single dose OR 0 mg PO BID for 7 days		
Conjunctiva or	Disseminated Gon	ococcal Infection (DGI)	;	
Children (<45	kg): Vagina, Urethi • Ceftriaxone 123	a , Pharynx, Rectum 5 mg IM once	• Spectinomycin 40mg	∕kg IM single dose (max 2៛
Neonates With	Ophthalmia Neon • Ceftriaxone 25- (max 125 mg)	atorum and Born to Infec 50 mg∕kg IV or IM singl	e dose	
Pregnancy	• Ceftriaxone 12	5 mg IM single dose	• Spectinomycin 2 g IN	⁄I single dose
Herpes Simplex	Virus			
First Clinical E	pisode of Genital F • Acyclovir 400 n • Acyclovir 200 n • Famciclovir 25 • Valacyclovir 1	lerpes ng PO TID for 7-10 days ng PO 5 times a day for 7 0 mg PO TID for 7-10 days g PO BID for 7-10 days C	OR -10 days OR ys DR	
Episodic Recur	rent Infection • Acyclovir 400 n • Acyclovir 200 n • Acyclovir 800 n • Famciclovir 123 • Valacyclovir 50	ng PO TID for 5 days OR ng PO 5 times a day for 5 ng PO BID for 5 days OR 5 mg PO BID for 5 days C 0 mg PO BID for 5 days	days OR DR	
Daily Suppress	ive Therapy			
	 Acyclovir 400 r Valacyclovir 25 Valacyclovir 50 Valacyclovir 10 Famciclovir 250 	ng PO BID OR 50 mg PO BID OR 10 mg PO once a day OR 100 mg PO once a day OI 1 mg PO BID	2	
HIV Infection	Immunocomproi from increased d	nised patients may benef osages of antiviral drugs	it 6	

* Any one of the bulleted treatments listed above an italicized *PLUS* must be **combined** with **any one** listed below.

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Disease	Recommended	Treatment	Alternative
Human Papillo	ma Virus		
Clinical and S	ubclinical Genital V	Varts ⁶	
Nongonococca	Urethritis ^{2,3}		
	• Azithromycin 1 • Doxycycline 10	g PO single dose OR 0 mg PO BID for 7 days	 Erythromycin base 500 mg PO QID for 7 days OR Erythromycin ethylsuccinate 800 mg PO QID for 7 days OR Ofloxacin 300 mg PO BID for 7 days OR Erythromycin base 250 mg PO QID for 14 days OR Erythromycin ethylsuccinate 400 mg PO QID for 14 days
Pediculosis Pub	is		
	 Permethrin 1% area and washe Lindane 1% sha affected area the Pyrethrins with affected area an 	cream rinse applied to affected d off after 10 minutes OR ampoo applied for 4 minutes to the en washed off ⁷ OR Piperonyl Butoxide applied to d washed off after 10 minutes	
Pelvic Inflamm	atory Disease (O	utpatient Management) ^{2,3}	
Regimen A:	• Ofloxacin 400 n • Metronidazole	ng PO BID for 14 days PLUS 500 mg PO BID for 14 days	
Regimen B:	 Ceftriaxone 250 Cefoxitin 2 g IM Other third gene Pl Doxycycline 10 	omg IM once OR I PLUS Probenecid 1 g PO (concurr eration Cephalosporin L US * 0 mg orally BID for 14 days	ently) OR
Pregnancy	• Patients should with the approp treatments. ⁶	be hospitalized and treated riate recommended parenteral IV	
HIV Infection	•Immunosuppres receive aggressi parenteral antin CDC guidelines	ssed HIV infected women should ve therapy using one of the nicrobials recommended in the	
Scabies			
	• Permethrin 5 % areas of the bod and washed off	cream applied to all y from the neck down after 8-14 hours	 Lindane 1% 1 oz of lotion or 30 g of cream applied thinly to all areas from the neck down and washed off after 8 hours⁷ OR Sulphur 6% precipitated in ointment applied thinly (neck down) nightly for 3 nights (Previous applications should be washed off before new ones are applied and last one washed off after 24 hours.)

* Any one of the bulleted treatments listed above an italicized *PLUS* must be **combined** with **any one** listed below.

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Disease	Recommended	Treatment		Alternative	
Syphilis ^{8,9}					
Primary, Secon Adults	dary, or Early Late • Benzathine pen in a single dose	nt (< 1 year) icillin G 2.4 million unit	s IM	<i>For penicillin allergy only (adults</i> • Doxycycline 100 mg PO BID fo • Tetracycline 500 mg PO OID f	<i>only):</i> or 14 days or 14 days
Children	• Benzathine pen the adult dose o	icillin G 50,000 units/kg l f 2.4 million units in a sir	M, up to Igle dose		or r r duys
Late Latent (>	1 year) or Latent of	Unknown Duration			
Adults	• Benzathine pen for 3 doses, 1 we	icillin G 2.4 million units eek apart (total 7.2 million	s IM 1 units)	 For penicillin allergy only (adult) Doxycycline 100 mg PO BID fo Tetracycline 500 mg PO QID for 	s only): or 28 days or 28 days
Children	• Benzathine pen the adult dose o 3 doses at 1 wee up to 7.2 millior	icillin G 50,000 units/kg f 2.4 million units, admin k intervals (total 150,000 n units)	IM up to istered in units/kg		
Neurosyphilis					
	• Aqueous crysta units a day, adn every 4 hours fo	lline penicillin G 18 - 24 ninistered as 3-4 million u or 10-14 days	million ınits IV	• Procaine penicillin 2.4 million PLUS probenecid 500 mg PO Q 10-14 days (Compliance must be assured.	units IM a day ID, both for)
HIV Infection	For all early syph	ilis treat as above. ⁶			
Pregnancy	Penicillin is the or syphilis during p should be desense Dosages are the s each stage of syph	nly recommended treatm regnancy. Women who a itized and then treated w ame as in nonpregnant p nilis. Erythromycin is NC	ent for re allergic ith penicill atients for V T recomm	in. ended.	
Congenital Syp	hilis ⁶				
с <i>н</i>	 Aqueous crysta units/kg/day, a IV every 12 hour every 8 hours th Procaine penici single dose for 1 	lline penicillin G 100,00 dministered as 50,000 uni rs during the first 7 days ereafter for a total of 10 d llin G 50,000 units/kg/d 0 days	-150,000 ts/kg/dose of life, and ays OR lose IM a da	e ay	
Trichomoniasis					

• Metronidazole 2 g PO single dose⁶

Footnotes @

*For more information, refer to the complete CDC document (*MMWR Recommendations and Reports, January 23, 1998/47[RR-1]: 1-118) *or call TDH at (512) 490-2500.* STD/HIV program staff are also available to assist health care providers with confidential notification of sexual partners exposed to STD and HIV.

Footnotes

- 1. All sex partners within 10 days preceding the onset of symptoms should be examined and prophylactically treated.
- 2. Quinolones are contraindicated in pregnant and lactating women and not approved for children <18 years old. (See CDC guidelines about safety profile for children.)
- 3. All sex partners within 60 days preceeding onset of symptoms should be examined and prophylactically treated.
- 4. The effectiveness of treatment with erythromycin is approximately 80%; a second course of therapy may be required.
- 5. Patients with gonococcal infections should also receive an adequate treatment for *Chlamydia trachomatis* since coinfection can occur in up to 40% of cases.
- 6. See CDC guidelines.
- 7. Lindane is not recommended for pregnant or lactating women or for children aged <2 years.
- 8. All sex/needle sharing partners should be examined as follows: 3 months plus duration of symptoms for primary syphilis, 6.5 months plus duration of symptoms for secondary syphilis, and 1 year for early latent syphilis.
- 9. All sex/needle sharing partners with an exposure during the preceding 3 months should be prophylactically treated.

Mumps Outbreak in Gillespie County

A worker in a turkey processing plant in Fredricksburg experienced onset of parotitis on January 8, 1999. He was most likely the source of infection for 2 coworkers who became ill on January 28 and February 1. The three men, aged 30, 50, and 52 years, worked in close proximity to one another. All experienced orchitis as well as parotitis. The immunization status of the 3 men is not known. Other workers in the plant have been vaccinated since it is likely that many were susceptible to mumps.

Mumps is a viral illness with acute onset of uni- or bilateral tender, self-limited swelling of the parotid or other salivary glands, lasting 2 days or more and without other apparent cause. Parotitis, however, can also be caused by influenza, parainfluenza type 3, and cytomegaloviruses, as well as numerous other noninfectious diseases. Approximately 30% of sporadic parotitis are not caused by mumps virus, and 20% to 40% of mumps patients may not have parotid swelling. Orchitis may accompany chlamydia, gonorrhea, chickenpox, brucellosis, tuberculosis, lymphocytic choriomeningitis, relapsing fever, leptospirosis, meliodosis, and pleurodynia. Although single cases of parotitis and orchitis are often of nonmumps etiology, epidemiologically linked cases are much more likely to be due to mumps.

As mumps incidence continues to decline due to increasing implementation of 2 doses of measles/mumps/rubella vaccine, laboratory confirmation is critical to documentation of progress toward eliminating indigenous mumps in the United States. Mumps can be confirmed only through mumps-specific laboratory testing (ie, mumps IgM antibody). (See next page for specimen submission guidelines.)

Normally, suspected mumps cases are reported on a weekly basis by phone: (800) 252-8239. Due to this recent outbreak, the Texas Department of Health (TDH) requests that suspected mumps cases be reported as soon as feasible. *Contact the TDH Immunization Division at* (800) 252-9152.

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Laboratory Confirmation of Mumps: Specimen Collection and Submission

- Collect a single 5 ml specimen of blood \geq 3 days following onset of symptoms and up to 30 days after parotid swelling.
- Collect this specimen in any collection tube without anticoagulant (eg, a red-top tube).
- Separate serum from blood and store serum in a sterile container at 2º-3ºC.
- Freeze serum if there will be more than 3 days from collection to receipt in the laboratory. Whole blood may be sent if specimen is shipped on day of collection. Do not freeze whole blood.
- Label blood tubes or serum containers with the patient's name and date of birth or social security number. Make sure this information on the tube label matches exactly what is written on the laboratory submission form (G-1).
- Specify "Mumps IgM Test" on the label.
- Mail IgM specimens to Viral Diagnostics, Inc., 670 West Arapaho #9, Richardson, TX 75080.

The TDH Immunization Division will pay for mumps IgM testing submitted to Viral Diagnostics provided that the suspected case is reported to the division prior to submitting the specimen.