

Last Name: _____		First Name: _____		MI: _____		Patient Phone: _____	Race: <input type="checkbox"/> Caucasian <input type="checkbox"/> African American <input type="checkbox"/> Asian / Pac. Islands <input type="checkbox"/> American Indian <input type="checkbox"/> Other (Specify) _____	Ethnicity: Hispanic Non-Hispanic Unknown	Sex: <input type="checkbox"/> M / <input type="checkbox"/> F	Date of Birth: <input type="checkbox"/> / <input type="checkbox"/> / <input type="checkbox"/> <input type="checkbox"/> M <input type="checkbox"/> D <input type="checkbox"/> Y	
Address: _____											
City: _____			State: _____			Zip Code: _____		County: _____			
<b>SYPHILIS</b>				<b>CHLAMYDIA/GONORRHEA</b> (check all that apply)				Date of Dx: _____			
<input type="checkbox"/> 710 Primary (Initial Lesion Present) <input type="checkbox"/> 720 Secondary (Lesions of Skin or Mucosa) <input type="checkbox"/> 730 Early Latent (Less than One Year) <input type="checkbox"/> 745 Late Latent (More than One Year) <input type="checkbox"/> 750 Late Syphilis (>1 Year, Symptomatic) <input type="checkbox"/> 760 Neurosyphilis <input type="checkbox"/> 790 Congenital Syphilis				<input type="checkbox"/> 100 Chancroid <input type="checkbox"/> 200 Chlamydia <input type="checkbox"/> 300 Gonorrhea <input type="checkbox"/> 350 Gonorrhea, Drug Resistant <input type="checkbox"/> 490 Pelvic Inflammatory				Pregnant? <input type="checkbox"/> yes <input type="checkbox"/> no Sex partner(s) name(s)    Sex partner(s) treatment			
				Source: <input type="checkbox"/> Urogenital <input type="checkbox"/> Urine <input type="checkbox"/> Rectal <input type="checkbox"/> Pharyngeal <input type="checkbox"/> Ophthalmia <input type="checkbox"/> Other				Marital Status: <input type="checkbox"/> Married <input type="checkbox"/> Divorced <input type="checkbox"/> Single <input type="checkbox"/> Co-habiting			
Patient Treated <input type="checkbox"/> Yes <input type="checkbox"/> No If Yes: _____ Date Treated: _____ Treatment: _____ Provider Name: _____ Provider Address: _____				Patient Treated <input type="checkbox"/> Yes <input type="checkbox"/> No If Yes: _____ Date Treated: _____ Treatment: _____ Provider Name: _____ Provider Address: _____				Date of Report: _____			

FORWARD REPORT OF DIAGNOSIS TO: WEST VIRGINIA DEPARTMENT OF HEALTH AND HUMAN RESOURCES, STD PROGRAM  
 350 CAPITOL STREET, ROOM 125 CHARLESTON, WV 25301-3715

Early - Primary, Secondary, Or Latent <1YR <b>BENZATHINE PENICILLIN G</b> - 2.4 MU IM in single dose	<b>AZITHROMYCIN</b> - 1 g orally in a single dose <i>OR</i> <b>DOXYCYCLINE</b> - 100 mg orally twice a day for 7 days	<b>CEFTRIAZONE</b> - 250 mg im in a single dose <i>OR, IF NOT AN OPTION</i> <b>CEFIXIME</b> - 400 mg orally in single dose <i>OR</i> Single-dose injectable <b>CEPHALOSPORIN</b> regimens <b>PLUS</b> <b>AZITHROMYCIN</b> - 1 g orally in a single dose <b>AZITHROMYCIN</b> - 2 g orally can be considered for pregnant women who cannot tolerate a <b>Cephalosporin</b> If <b>Cephalosporin</b> allergy: See 2015 CDC treatment guidelines.
Latent > 1YR, Latent Of Unknown Duration, Late (Cardiovascular, Gumma) <b>BENZATHINE PENICILLIN G</b> 2.4 MU X 3 IM at 1 week intervals (7.2 MU total)	<b>Alternative Regimens</b> <b>ERYTHROMYCIN</b> - base 500 mg orally four times a day for 7 days <i>OR</i> <b>ERYTHROMYCIN ETHYLSUCCINATE</b> - 800 mg orally four times a day for 7 days <i>OR</i> <b>LEVOFLOXACIN</b> - 500 mg orally once daily for 7 days <i>OR</i> <b>OFLOXACIN</b> - 300 mg orally twice a day for 7 days	All contacts of gonorrhea and chlamydia should receive prophylactic treatment using the treatment schedule provided.
<b>Pregnant Women:</b> See 2015 CDC treatment guidelines.	<b>Pregnant Women</b> <b>AZITHROMYCIN</b> - 1 g orally in a single dose <i>OR</i> <b>AMOXICILLIN</b> - 500 mg orally three times a day for 7 days	To view CDC guidelines, go to <a href="http://www.cdc.gov/std/tg2015/default.htm">http://www.cdc.gov/std/tg2015/default.htm</a> or call West Virginia Bureau for Public Health Division of STD/HIV & Hepatitis 800-642-8244 or 304-558-2195. Treatment may also be faxed to the STD Program at 304-558-6478.
Neurosyphilis - <b>AQUEOUS CRYSTALLINE PENICILLIN G</b> - 3 to 4 MU IV every 4 HRS for 10 to 14 days (18-24 MU/day)	<b>Alternative Regimens For Pregnant Women</b> <b>ERYTHROMYCIN</b> - base 500 mg orally four times a day for 7 days <i>OR</i> <b>ERYTHROMYCIN</b> - base 250 mg orally four times a day for 14 days <i>OR</i> <b>ERYTHROMYCIN ETHYLSUCCINATE</b> - 800 mg orally four times a day for 7 days <i>OR</i> <b>ERYTHROMYCIN ETHYLSUCCINATE</b> - 400 mg orally four times a day for 14 days	
Congenital Syphilis - <b>AQUEOUS CRYSTALLINE PENICILLIN G</b> 100,000 - 150,000 UNITS/KG day (50,000 UNITS KG dose IV every 12 HRS) during the first 7 days of life and every 8 HRS thereafter for total of 10 days <i>OR</i> <b>PROCAINE PENICILLIN G</b> - 50,000 UNITS/KG dose IM in single dose for 10 days		
<b>Children:</b> See 2015 CDC treatment guidelines.		

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WEST VIRGINIA DEPARTMENT OF HEALTH AND HUMAN RESOURCES