

# Evaluating Patients For Primary Syphilis

## CLINICAL PRESENTATIONS OF PRIMARY SYPHILIS

- Lesion appears 10-90 days after contact at site of exposure; may persist for 2-3 weeks then resolves
- Usually genitorectal but may be extragenital, depending on exposure site
- Clinical presentation, typical or atypical
- Typical: single painless, indurated, clean-based ulcer with rolled edges & bilateral painless adenopathy
- Atypical: can mimic herpes & other genital ulcers
- ~25% present with multiple lesions
- Lesions of primary and secondary syphilis can be present at the same time, especially in HIV positive individuals

## Differential Diagnosis

- Herpes (most common), primary HIV ulcers, chancroid, granuloma inguinale, trauma, and many non-STD infectious and non-infectious causes of genital ulcers
- More than one etiology can be present at the same time



D Syphilitic Ulcer, Shaft



W Syphilitic Ulcer, Shaft



S Multiple Syphilitic Ulcers, Shaft



S Multiple Syphilitic Ulcers Resembling Herpes



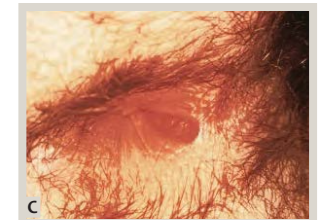
C Syphilitic Ulcer, Vulva



S Multiple Syphilitic Ulcers, Vulva



S Crusted Syphilitic Ulcer, Urethra



C Syphilitic Ulcer, Perianal

## Photo Credits

C - Centers for Disease Control and Prevention; D - With permission from the Denver Metro Health Clinic; S - With permission from San Francisco City Clinic; W - With permission from University of Washington STD Prevention Training Center Washington (photos from UW HSCER Slide Bank)

## To Order Additional Copies

See the online version of the Primary Syphilis Algorithm on the clinical resources page of the CA PTC website: [www.californiaptc.com](http://www.californiaptc.com)

## Acknowledgements

Medical Directors from the National Network of STD Clinical Prevention Training Centers, California STD Controllers Association, Division of STD Prevention of the Centers for Disease Control and Prevention

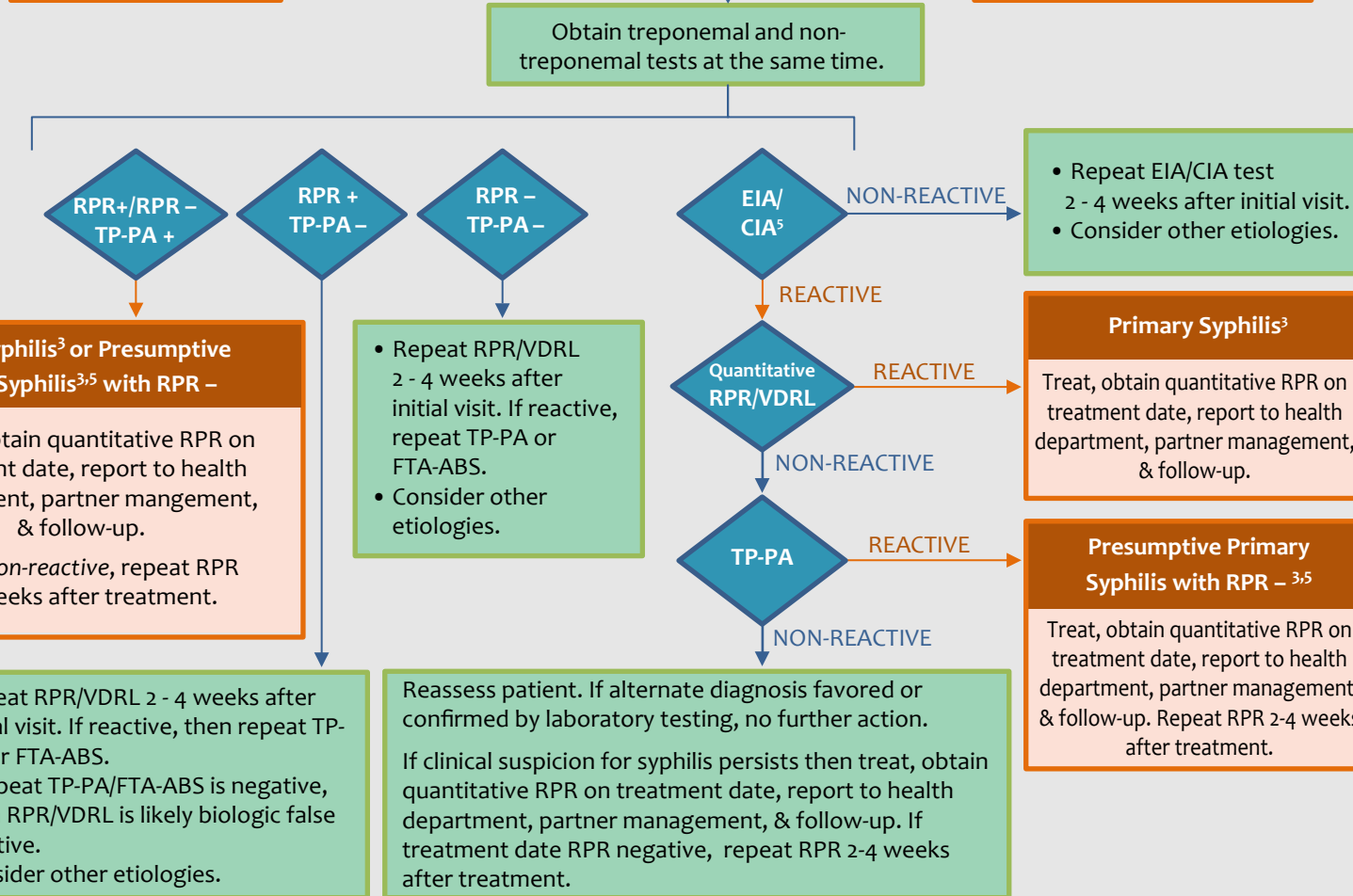
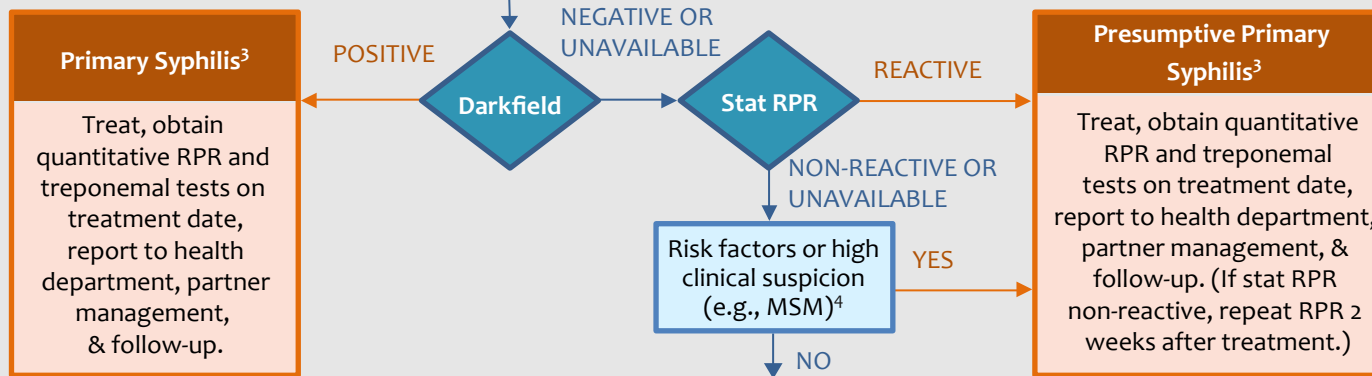
Revised 7/2018

Patient with new genital lesion or suspicious genital ulcer

## SEXUAL HISTORY, RISK ASSESSMENT, & PHYSICAL EXAM

### DIAGNOSTIC WORK-UP

- Darkfield (if available)
- Stat RPR (if available)
- RPR or VDRL serology (quantitative)
- Treponemal test<sup>1</sup> (TP-PA/FTA-ABS/EIA/CIA)
- Herpes culture or PCR<sup>2</sup>
- HIV Test



## SEXUAL HISTORY, RISK ASSESSMENT, & PHYSICAL EXAM

### Sexual History, Risk Assessment (past year)

- Gender of partners, number of partners (new, anonymous, serodiscordant HIV status, exchange of sex for drugs or money)
- Types of sexual exposure
- Recent STDs; HIV serostatus
- Substance abuse
- Condom use

### History of Syphilis

- Prior syphilis (last serologic test & last treatment)

### Physical Exam

- Oral cavity
- Lymph nodes
- Skin
- Palms & soles
- Neurologic
- Eyes
- Genitalia/pelvic
- Perianal

## DIAGNOSTIC ISSUES IN PRIMARY SYPHILIS

- **Darkfield** ~ 80% sensitive, varies with skill of examiner; decreased sensitivity as lesion ages
- A negative RPR/VDRL does not exclude syphilis diagnosis; ~75-85% sensitive in primary syphilis
- Use same test (RPR or VDRL) in sequential testing; titers are not interchangeable
- Need both non-treponemal (RPR or VDRL) and treponemal test (TP-PA, FTA-ABS, EIA, CIA) to make syphilis diagnosis
- Treponemal tests can remain positive for life; utility limited in patients with history of prior syphilis, comparison of non-treponemal titers needed

### For more details on Treponemal Immunoassays:

[www.cdph.ca.gov/Programs/CID/DCDC/CDPH%20Document%20Library/UseofTreponemalImmunoassays\\_Syphilis.pdf](http://www.cdph.ca.gov/Programs/CID/DCDC/CDPH%20Document%20Library/UseofTreponemalImmunoassays_Syphilis.pdf)

**Note:** Evaluate for neurosyphilis (assess if neurologic, ophthalmic or otic symptoms present, as neurosyphilis can occur at any stage of syphilis)

## TREATMENT & FOLLOW-UP

### Treatment of Primary Syphilis

#### Recommended Regimen

- Benzathine Penicillin G 2.4 million units IM x 1

#### Alternative Regimens for Penicillin Allergic Non-Pregnant Patients:

Efficacy not well established & not studied in HIV+ patients; close follow-up essential:

- Doxycycline 100 mg po bid x 2 weeks or
- Tetracycline 500 mg po qid x 2 weeks

\*Pregnant patients with penicillin allergy should be desensitized and treated with penicillin

See CDC STD Treatment Guidelines: [www.cdc.gov/std/treatment](http://www.cdc.gov/std/treatment)

#### California STD Treatment Guidelines Grid:

[www.cdph.ca.gov/Programs/CID/DCDC/CDPH%20Document%20Library/STD-Treatment-Guidelines-Color.pdf](http://www.cdph.ca.gov/Programs/CID/DCDC/CDPH%20Document%20Library/STD-Treatment-Guidelines-Color.pdf)

#### \*\*Additional Testing and Follow-up

**Note:** Also test for HIV, GC/CT, and pregnancy (if female of reproductive age)

- 1-2 weeks: clinical follow-up
- 3, 6, 9, 12, 24 months: serologic follow-up for HIV+ patients
- 6, 12 months: serologic follow-up for HIV- patients
- Failure of titer to decline fourfold (e.g. 1:64 to ≤ 1:16) within 6-12 months from titer at time of treatment may indicate treatment failure. Titer decline may be slower in HIV+ patients.
- Consider retreatment and CSF evaluation if titer fails to decline appropriately

## REPORTING & PARTNER MANAGEMENT

- All syphilis cases and presumptive cases must be reported to the local health department within one working day of diagnosis
- Local health departments will assist in partner notification & management
- Contact Number at Local Health Department: \_\_\_\_\_

<sup>1</sup> Treponemal tests may be more sensitive than non-treponemal tests during primary syphilis.

<sup>2</sup> Also consider culture for Haemophilus ducreyi (chancroid) if exposure in endemic areas or if lesion does not respond to syphilis treatment.

<sup>3</sup> All patients with suspected syphilis should be tested for HIV infection and screened for other STDs. Repeat HIV testing of patients with primary syphilis 3 months after the first HIV test, if the first test is negative.

<sup>4</sup> If the patient is a man who has sex with men (MSM) or has high risk sexual behavior or clinical exam with classic features of a syphilitic ulcer, then standard of care includes presumptive treatment at the time of the initial visit before diagnostic test results are available. Presumptive treatment is also recommended if patient follow-up is a concern.

<sup>5</sup> If the patient does not respond to treatment, repeat RPR/VDRL after treatment and consider other etiologies.