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### Syphilis – "The Great Imitator"

Wake AHEC requires all speakers to disclose any relevant financial conflicts of interest.

Victoria Mobley has no relevant financial conflicts of interest to disclose.

#### **Learning Objectives**

- When to suspect syphilis
- Typical & atypical clinical presentations of syphilis
- Diagnosing syphilis
- How to treat syphilis

North Carolina
Public Health

Other clinical considerations



# When should you suspect syphilis infection

### <u>ALWAYS</u>





Warning: The next several slides have some graphic pictures - though, let's be honest, if you are a public health clinician this is nothing you haven't seen before...

# Typical Presentations of Primary Syphilis

# **Atypical Presentations of Primary Syphilis**

### Typical Presentations of Secondary Syphilis

### **Atypical Presentations of Secondary Syphilis**

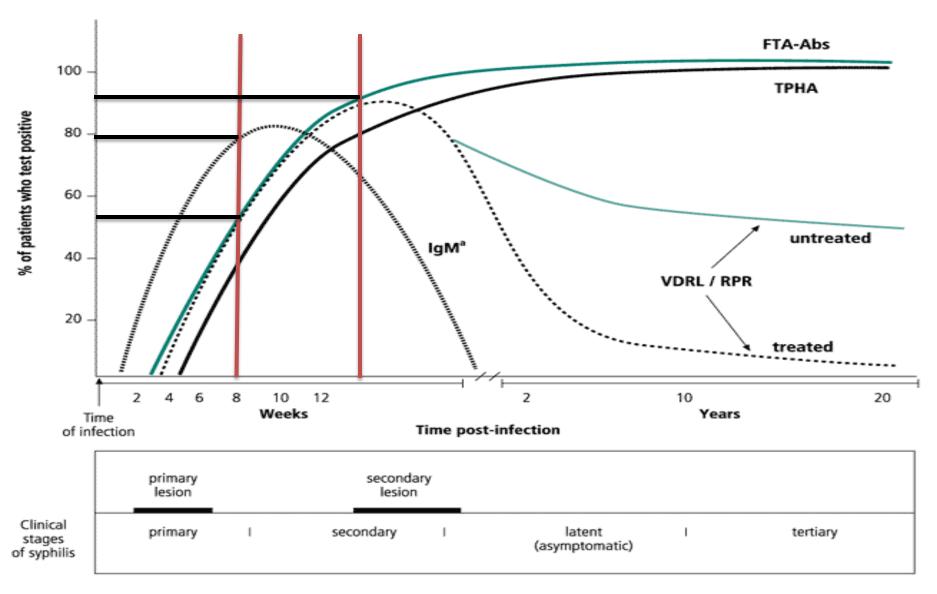


# Other Atypical Presentations of Early Syphilis Infection

Stage	Symptom Onset	Clinical Features HIV negative	Clinical Features HIV positive
Primary	2-3 weeks	Single chancre	Multiple chancres
Secondary	3-6 weeks (after chancre heals)	Rash Condylomata lata, Alopecia LAD Viral-like syndrome	Overlap of P&S symptoms Malignant lues Ocular symptoms
Early Latent Late Latent	<1 year >1 year	Asymptomatic	Asymptomatic
Tertiary Late benign	4-10 years	Gummas: skeleton, spine, mucosa	Gummas may form in months vs. years
Cardiovascular	15-20 years	Endarteritis obliterans CAD involvement Aortic insufficiency	Rapidly progressing aortitis
Neurosyphilis	Anytime	Asymptomatic→ varied neuro deficits	More rapid sxs progression

#### Syphilis Serologies

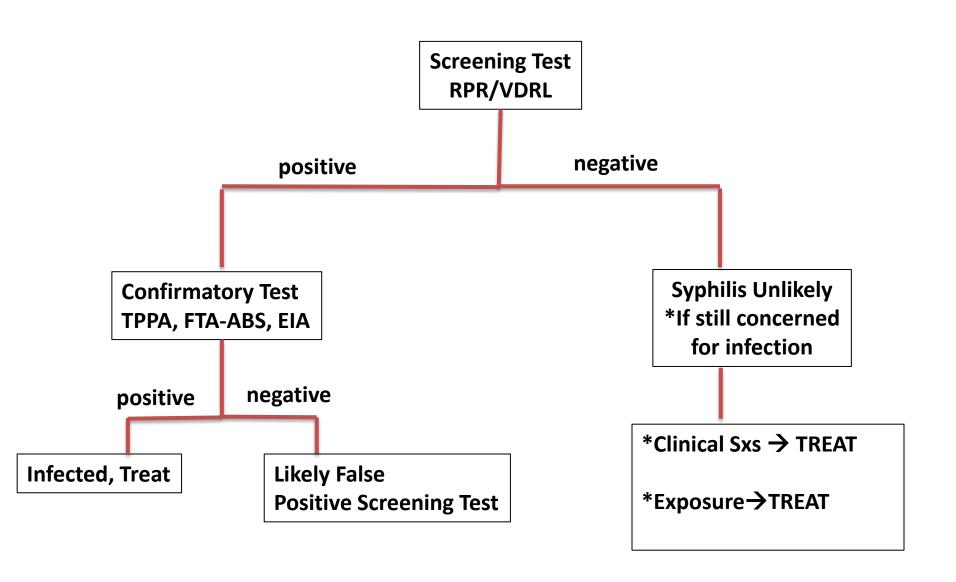
Fig. 1. Common patterns of serological reactivity in syphilis patients

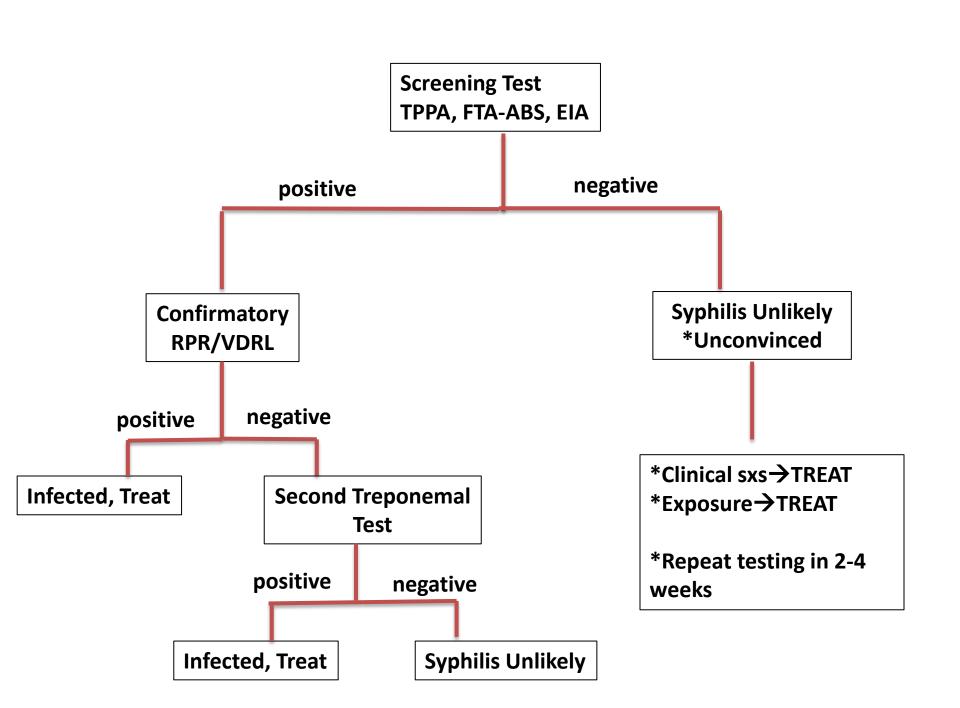




#### **Sensitivity of Syphilis Diagnostic Tests**

Test	Primary	Secondary	Latent
VDRL/RPR	75-85%	100%	95-98%
FTA-ABS	84-86%	100%	100%
TPPA	84-86%	100%	100%
EIA/CIA	93-98%	100%	100%





Stage	Treatment (regardless of HIV status)	Alternative Treatments	Follow-up
Primary, Secondary & Early Latent	Benzathine penicillin G 2.4 million units IM X 1 dose	<ul> <li>Doxycycline 100 BID x         7 days     </li> <li>Tetracycline 500 QID         x 7 days     </li> </ul>	<ul> <li>Clinical f/u 2-4 weeks</li> <li>Lab f/u: 6, 12 and 24 months</li> <li>*HIV positive: 9 and 12 months also</li> </ul>
Latent or unknown duration	Benzathine penicillin G 7.2 million units IM split into 3 weekly doses	<ul> <li>Doxycycline 100 BID x 28 days</li> <li>Tetracycline 500 QID x 28 days</li> <li>Ceftriaxone (CTX)-dose unknown- under specialist's care</li> </ul>	<ul> <li>Clinical f/u 2-4 weeks</li> <li>Lab f/u: 6, 12 and 24 months</li> <li>*HIV positive: 9 and 12 months also</li> </ul>
Neurosyphilis *ocular syphilis	Aqueous crystalline penicillin G 18–24 million units per day X 10–14 days	<ul> <li>Procaine PCN G 2.4         million units IM daily</li> <li>PLUS         Probenecid 500 mg</li> <li>QID x 10-14 days</li> <li>CTX 2 gms IV/IM daily         x 10-14 days</li> </ul>	<ul> <li>Clinical f/u every 6 months up to 2 years until CSF abnl resolve</li> </ul>



#### **Other Considerations**

- Beware of the PROZONE effect
- Always use same testing method for f/u i.e. RPR=RPR; RPR≠VDRL
- True serologic difference is ≥4-fold change in RPR titer
  i.e. 1:2→1:4
  1:2→1:8
- Not all PCN preparations are equal (i.e.Billicin C-R)
- When/Who to retest for syphilis
  - Every 3-6 months in high-risk individuals
- HIV testing should be performed in all patients diagnosed with syphilis, unless they are already known to be positive

### Questions??????