

SAN MATEO COUNTY PUBLIC HEALTH LABORATORY TEST INFORMATION

Section: Syphilis Serology Test Name: Syphilis EIA

Test Includes: Captia Syphilis EIA		
Reporting		
Results Available: 4 days from receipt	Contact Number: (650) 573-2500	
Reference		
Method: Captia Syphilis EIA performed on Bio-Rad Evolis instrument		
Turnaround Time: 4 days if non-reactive	Reference Range: Non-Reactive	
Limitations: May not detect a recent infection,	Interpretation: Non-reactive indicates that the	
or infection in a person with a severely	patient does not have detectable antibody to the	
compromised immune system or an old,	infectious agent. Reactive indicates that the	
successfully cured infection (>10 years for	patient has detectable antibody to the infectious	
example). The test can cannot be used to	agent, and depending on the clinical picture,	
distinguish between active and cured cases nor	may have a current or past infection.	
to determine responses to therapy.	-	
Specimen Requirements		
Specimen Collection: Venipuncture	Sample Type: Blood	
Volume/Amount Required: 7-10 mls whole	Preferred Specimen: Serum, EDTA or citrated	
blood or 5 ml serum or plasma	plasma	
Collection/Preservation: Red top or tiger top	Storage Instructions: Do not freeze or	
tube, EDTA or citrate.	refrigerate whole blood. Separated serum may	
	be held at 2-8° C for up to 5 days. Serum can	
	be stored frozen at -20°. Plasma samples may	
	be stored at 2-8° C for up to 48 hours. Plasma	
	samples should not be frozen.	
Causes for Rejection: Discrepancy in	Sample Container: Red top or tiger top tube;	
specimen identification; insufficient quantity of	EDTA or Citrate	
specimen; gross hemolysis		
Sample Test Kit:	Availability: Performed Tuesday and Friday	
Diagnostic Information: The Captia Syphilis E	IA test is a test for the qualitative detection of	
IgG antibodies to <i>T. pallidum</i> is serum (or plasma	a) specimens, to be used in conjunction with non-	
treponemal testing to provide serological evidence	ee of infection with <i>T. pallidum</i> . Any reactive or	
equivocal results from this test must be supplement	ented with a quantitative non-treponemal test	
(such as the RPR) to distinguish from active dise	ase and assist in ruling out false positives.	
Specimen Submission		
Request Form: Standard Clinical Test Request	Specimen Handling: Use Universal	
form or through computer interface	Precautions	
Transport Temperature: Ambient	Shipping Requirements: Ship on cold packs	
temperature for clotted blood; separated serum	(2-8°C), or frozen (-20°C) and mailed on dry	
at 2-8°C (refrigerated) or -20°C (frozen).	ice.	



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Billing	
CPT Code(s):	Fees: \$10.00
Effective Date: July 1, 2008	