FACT SHEET

Treponemal pallidum Particle Agglutination (TPPA)

Test Description:

The SLO Public Health Laboratory performs the Serodia® -TP• PA test. It is specific for detecting Treponema pallidum antibodies in serum or plasma samples. It does not detect T. pallidum directly. As with all serological tests for syphilis, interpretation of results obtained with the Serodia® -TP•PA syphilis Antibody test must be used in conjunction with the patient's clinical symptoms, medical history and other clinical and/or laboratory finding to produce an overall clinical diagnosis.

Specimens giving inconclusive results in the assay should be re-tested. A repeated inconclusive specimen should be reported as Inconclusive and another specimen drawn in two weeks for testing and/or confirmed by other methods, such as FTA-ABS. The Serodia® -TP• PA is less sensitive than the fluorescent treponemal antibody absorption (FTA-ABS) in primary syphilis but compares favorably in all other stages of syphilis. All treponemal tests end to remain reactive following treponemal infection: therefore they should not be used to evaluate.

Sensitivity:	70-90% primary syphilis
	90-99% secondary syphilis
	70-90% tertiary syphilis
Specificity:	95-99% primary syphilis
	90-99% secondary syphilis
	90% tertiary syphilis

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nfection such as tropical trepanematosis (Yaws, Pinata, Bejel) are likely to result in a positive TP-PA test.

Specimens:

Serum (specimen of choice) or EDTA Plasma sample removed from the red cells, free of hemolysis, bacterial contamination, or lipemia.

Specimens may be stored up to 5 days at refrigerated temperature 2-8C before processing.

CPT Code 86780



San Luis Obispo County Public Health Laboratory

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