## 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
I
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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