

Treponema Pallidum Antibody (TP)

Item Number: TP

Introduction

Discover Anti-Treponema pallidum antibody (TP) for reliable Syphilis detection via IFA and immunohistochemistry. High purity, polyclonal antibody with excellent performance. Enquire now!

Learn More

Feature	Description
Product Name	Treponema pallidum antibody
Host Species	Rabbit
Application	IFA, Immunohistochemistry
Form/Appearance	Purified from antiserum
Concentration	5 mg/ml
Isotype	lgG
Clonality	Polyclonal
Purity	≥95%
Buffer	0.01M Phosphate Buffered Saline, pH 7.2, containing 0.1% sodium azide
Cross Reactivity	Antiserum has not been absorbed and may react with related microorganisms
Specificity	All antigens
Shelf Life	Six years from date of manufacture
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Condition	Description
Storage (Short Term)	2-8°C (Less than 6 months)
Storage (Long Term)	-20°C
Shipping	Cold Packs
Indicator	Acceptance Criteria
Indicator Appearance	Acceptance Criteria Liquid component should be clear and transparent; kit components should be complete and intact; liquid should not leak; packaging label should be clear and undamaged.
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Appearance	Liquid component should be clear and transparent; kit components should be complete and intact; liquid should not leak; packaging label should be clear and undamaged.
Appearance Volume Positive Reference	Liquid component should be clear and transparent; kit components should be complete and intact; liquid should not leak; packaging label should be clear and undamaged. Each component should not be less than the indicated value. When tested with Treponema pallidum antibody (TP) national reference material or enterprise positive reference material standardized by national
Appearance Volume Positive Reference Conformity Rate Negative Reference	Liquid component should be clear and transparent; kit components should be complete and intact; liquid should not leak; packaging label should be clear and undamaged. Each component should not be less than the indicated value. When tested with Treponema pallidum antibody (TP) national reference material or enterprise positive reference material standardized by national reference material, no negative results should occur. When tested with TP national reference material or enterprise negative reference material standardized by national reference material, no positive
Appearance Volume Positive Reference Conformity Rate Negative Reference Conformity Rate	Liquid component should be clear and transparent; kit components should be complete and intact; liquid should not leak; packaging label should be clear and undamaged. Each component should not be less than the indicated value. When tested with Treponema pallidum antibody (TP) national reference material or enterprise positive reference material standardized by national reference material, no negative results should occur. When tested with TP national reference material or enterprise negative reference material standardized by national reference material, no positive results should occur. When tested with TP national reference material or enterprise sensitivity reference material standardized by national reference material, the
Appearance Volume Positive Reference Conformity Rate Negative Reference Conformity Rate Minimum Detection Limit	Liquid component should be clear and transparent; kit components should be complete and intact; liquid should not leak; packaging label should be clear and undamaged. Each component should not be less than the indicated value. When tested with Treponema pallidum antibody (TP) national reference material or enterprise positive reference material standardized by national reference material, no negative results should occur. When tested with TP national reference material or enterprise negative reference material standardized by national reference material, no positive results should occur. When tested with TP national reference material or enterprise sensitivity reference material standardized by national reference material, the results should meet L1, L2, and L3 being positive, and L4 being negative. When tested with TP national precision reference material or enterprise precision reference material standardized by national reference material,
Appearance Volume Positive Reference Conformity Rate Negative Reference Conformity Rate Minimum Detection Limit Repeatability	Liquid component should be clear and transparent; kit components should be complete and intact; liquid should not leak; packaging label should be clear and undamaged. Each component should not be less than the indicated value. When tested with Treponema pallidum antibody (TP) national reference material or enterprise positive reference material standardized by national reference material, no negative results should occur. When tested with TP national reference material or enterprise negative reference material standardized by national reference material, no positive results should occur. When tested with TP national reference material or enterprise sensitivity reference material standardized by national reference material, the results should meet L1, L2, and L3 being positive, and L4 being negative. When tested with TP national precision reference material or enterprise precision reference material standardized by national reference material, repeated measurements 10 times, the coefficient of variation (CV) should be no greater than 10%. When tested with TP national precision reference material or enterprise precision reference material standardized by national reference material,
Appearance Volume Positive Reference Conformity Rate Negative Reference Conformity Rate Minimum Detection Limit Repeatability Batch-to-Batch Difference	Liquid component should be clear and transparent; kit components should be complete and intact; liquid should not leak; packaging label should be clear and undamaged. Each component should not be less than the indicated value. When tested with Treponema pallidum antibody (TP) national reference material or enterprise positive reference material standardized by national reference material, no negative results should occur. When tested with TP national reference material or enterprise negative reference material standardized by national reference material, no positive results should occur. When tested with TP national reference material or enterprise sensitivity reference material standardized by national reference material, the results should meet L1, L2, and L3 being positive, and L4 being negative. When tested with TP national precision reference material or enterprise precision reference material standardized by national reference material, repeated measurements 10 times, the coefficient of variation (CV) should be no greater than 10%. When tested with TP national precision reference material or enterprise precision reference material standardized by national reference material, repeated measurements 10 times, the coefficient of variation (CV) should be no greater than 15%.