



REPORT

Report No./Day Sr : DRP/HEA/23-24/56 / 4
Name : DEMO
Age / Sex : 34-Years / M
Referring Doctor : Self
Doctor Advised : MANTOUX TEST
Sample Collected on : 22-Jul-2023
Sample Received on : 22-Jul-2023
Report Released on : 22-Jul-2023

MANTOUX TEST

Test Report Status	Result	
0.1 ML INJ OF IN 10,000 P.P D (LLU) GIVEN L/D ON L T /RT FORE ARM	123 H	0.00 - 0.00
GIVEN ON	20-Jul-2023	0.00 - 0.00
OBSERVED AFTER 48 HOURS	22-Jul-2023	0.00 - 0.00
OBSERVATION	On Left Arm	0.00 - 0.00
ERYTHEMA	222 H	0.00 - 0.00
INDURATION	333 H	0.00 - 0.00
VESICLE	Absent	0.00 - 0.00
RESULT : MANTOUX TEST IS	Negative	0.00 - 0.00

The TST is performed by injecting 0.1 ml of tuberculin purified protein derivative (PPD) into the inner surface of the forearm. The injection should be made with a tuberculin syringe, with the needle bevel facing upward. The TST is an intradermal injection. When placed correctly, the injection should produce a pale elevation of the skin (a wheal) 6 to 10 mm in diameter.

The skin test reaction should be read between 48 and 72 hours after administration by a health care worker trained to read TST results. A patient who does not return within 72 hours will need to be rescheduled for another skin test.

The reaction should be measured in millimeters of the induration (firm swelling). The reader should not measure erythema (redness). The diameter of the indurated area should be measured across the forearm (perpendicular to the long axis). Some persons may react to the TST even though they are not infected with M. tuberculosis. The causes of these false-positive reactions may include, but are not limited to, the following:

Previous TB vaccination with the bacille Calmette-Guérin (BCG) vaccine

Infection with nontuberculosis mycobacteria (mycobacteria other than M. tuberculosis)

Incorrect measurement or interpretation of reaction

Incorrect antigen used

A TB blood test is the preferred method of testing for people who have received the BCG vaccine in order to prevent false-positive reactions. TB blood tests are also called interferon-gamma release assays or IGRAs.

**** End of Report****

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If test result are alarming to unexcepted client is advised to contact to laboratory immediately for possible remedical action.

This report is not valid for medico legal purpose.