

Some patients may be at greater risk of developing TB than others



Tuberculosis (TB) cases in the US continue to increase¹

While anyone can get TB, the USPSTF recommends screening for latent TB infection in populations at increased risk²



18x
higher TB incidence
rate than US-born
persons³

HOMELESS
>8%
of people with TB (aged
15 or older) reported ever
experiencing homelessness,
according to 2023 data⁴

individuals (eg, those with diabetes, HIV, endstage renal disease) are at significantly higher risk of developing active TB1

IMMUNOCOMPROMISED

The CDC also recommends testing patients with diabetes, HIV, cancer, and other conditions⁵



DIABETES

>23%

of TB cases occurred in persons reported to have diabetes⁶

HIV

>4%
of people with TB are coinfected with HIV⁷

CANCER

9 x

more likely for people with certain cancers to develop active TB than to those without cancer⁸



Blood-based IGRA testing is the preferred testing method for TB

Quest Diagnostics is the only national commercial reference lab to offer a choice of TB blood tests.



IGRAs require just 1 tube, 1 draw, and 1 visit

Test information

Test name	Test code	CPT® codes
T-SPOT®.TB	37737	86481
QuantiFERON®-TB Gold Plus, 1 Tube	36970	86480
QuantiFERON®-TB Gold Plus, 4 Tubes, Draw Site Incubated	36971	86480

Count on Quest for guideline-driven screening solutions and a better experience



A simpler lab experience



Lower costs and greater value



Broad test portfolio and expertise you can trust

- 2,400 patient service centers with simple online scheduling
- Test results accessible to patients through MyQuest[®]
- Quest is in-network with the majority of health plans nationwide
- A financial assistance program is available to help qualified patients receive screening at a lower cost
- 3,500 tests, ranging from routine screening to highly specialized testing
- ~850 MDs and PhDs, and dozens of genetic counselors available for test selection and results interpretation



Visit QuestDiagnostics.com/TBTesting to learn more about the importance of testing patients who may be at increased risk for TB

T-SPOT $^{\odot}$. TB and QFT-Plus are approved by the US FDA.

The T-SPOT®. TB test is an in vitro diagnostic test for the detection of effector T cells that respond to stimulation by Mycobacterium tuberculosis antigens ESAT-6 and CFP 10 by capturing interferon gamma (IFN-y) in the vicinity of T cells in human whole blood collected in sodium citrate or sodium or lithium heparin. It is intended for use as an aid in the diagnosis of M tuberculosis infection. The T-SPOT®. TB test is an indirect test for M tuberculosis infection (including disease) and is intended for use in conjunction with risk assessment, radiography, and other medical and diagnostic evaluations. For more information, visit www.tspot.com.

QFT-Plus is approved by the FDA as an in vitro diagnostic aid for detection of Mycobacterium tuberculosis infection. It uses a peptide cocktail simulating ESAT-6 and CFP-10 proteins to stimulate cells in heparinized whole blood. Detection of IFN-y by ELISA is used to identify in vitro responses to these peptide antigens that are associated with M tuberculosis infection. QFT-Plus is an indirect test for M tuberculosis infection (including disease) and is intended for use in conjunction with risk assessment, radiography, and other medical and diagnostic evaluations. Up-to-date licensing information and product-specific disclaimers can be found at www.QuantiFERON.com.

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The CPT® codes provided are based on American Medical Association guidelines and are for informational purposes only. CPT coding is the sole responsibility of the billing party. Please direct any questions regarding coding to the payer being billed.

 $Test\ codes\ may\ vary\ by\ location.\ Please\ contact\ your\ local\ laboratory\ for\ more\ information.$

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