

***Mycobacterium tuberculosis*, Strain H37Rv, Cytosol Fraction**

**Catalog No. NR-14834**

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**For research use only. Not for use in humans.**

**Contributor:**

BEI Resources or NIH – TB Vaccine Testing and Research Materials Contract

**Manufacturer:**

Karen Dobos, Ph.D., Colorado State University, Fort Collins, Colorado, USA or NIH – TB Vaccine Testing and Research Materials Contract

**Product Description:**

NR-14834 is a preparation of the cytosol fraction of *Mycobacterium tuberculosis* (*M. tuberculosis*), strain H37Rv and contains cytosolic proteins and soluble material released from the cell wall during disruption of the bacilli. The culture was grown to late-log phase in glycerol-alanine-salts medium, washed with PBS and inactivated by gamma irradiation. The bacilli were suspended at a concentration of 2 g/mL in PBS containing 8 mM EDTA, DNase, RNase, and a proteinase inhibitor tablet, and broken in a French Press pressure cell at 4°C. Unbroken cells were removed by low speed (3,000 × g) centrifugation. The cell wall was isolated by centrifugation at 27,000 × g. The supernatant was subjected to a 100,000 × g centrifugation for four hours, then collected and dialyzed against 10 mM ammonium bicarbonate. The protein content was determined using the BCA protein assay.

**Material Provided:**

Each vial contains approximately 1 mg of cytosol fraction from *M. tuberculosis*, strain H37Rv provided in 10 mM ammonium bicarbonate.

**Packaging/Storage:**

NR-14834 was packaged aseptically in cryovials. The product is provided frozen on dry ice and should be stored at -80°C or colder immediately upon arrival. Freeze-thaw cycles should be avoided.

**Citation:**

Acknowledgment for publications should read “The following reagent was obtained through BEI Resources, NIAID, NIH: *Mycobacterium tuberculosis*, Strain H37Rv, Cytosol Fraction, NR-14834.”

**Biosafety Level: 1**

Appropriate safety procedures should always be used with this material. Laboratory safety is discussed in the following publication: U.S. Department of Health and Human Services, Public Health Service, Centers for Disease Control and Prevention, and National Institutes of Health. [Biosafety in Microbiological and Biomedical Laboratories \(BMBL\)](#), 6th ed. Washington, DC: U.S. Government Printing Office, 2020.

**Disclaimers:**

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**References:**

1. Lee, B. Y., S. A. Hefta and P. J. Brennan. “Characterization of the Major Membrane Protein of Virulent *Mycobacterium tuberculosis*.” *Infect. Immun.* 60 (1992): 2066-2074. PubMed: 1563797.
2. Cole, S. T., et al. “Deciphering the Biology of *Mycobacterium tuberculosis* from the Complete Genome Sequence.” *Nature* 393 (1998): 537-544. PubMed: 9634230. Erratum in: *Nature* 396 (1998): 190-198.
3. Hirschfield, G. R., M. McNeil and P. J. Brennan. “Peptidoglycan-Associated Polypeptides of *Mycobacterium tuberculosis*.” *J. Bacteriol.* 172 (1990): 1005-1013. PubMed: 2105289.

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