

Certificate of Analysis for NR-14855

Ag85 Complex, Purified Native Protein from Mycobacterium tuberculosis, Strain H37Rv

Catalog No. NR-14855

This reagent is the tangible property of the U.S. Government.

Product Description: The Antigen 85 Complex was isolated from *Mycobacterium tuberculosis* (*M. tuberculosis*), strain H37Rv culture filtrate proteins and purified. This complex of fibronectin-binding proteins Ag85A (FbpA), Ag85B (FbpB), and Ag85C (FbpC) is the major secreted protein component of *M. tuberculosis* culture fluids and plays a key role in the pathogenesis of tuberculosis. This protein complex is highly immunogenic and plays a role in cell wall assembly via a mycolyltransferase exchange process. All three purified antigens have shown extensive cross-reactivity and stimulate complement-mediated phagocytosis by host macrophages.

Lot: 63291509 Manufacturing Date: 03DEC2015

Production and QC testing were performed by Colorado State University (CSU). The Colorado State University documentation for lot 15.Rv.2.12.3.Ag85 is attached.

ATCC[®], on behalf of BEI Resources, hereby represents and warrants that the material provided under this certificate has been subjected by the contractor to the tests and procedures specified and that the results described, along with any other data provided in this certificate, are true and accurate to the best of ATCC[®]'s knowledge.

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WORK SHEET FOR PURIFIED NATIVE PROTEINS

General Information

BEI Catalog Number: NR-14855

Product Description: Ag85 Complex (mixture of Rv3804c, Rv1886c, Rv0129c)

CSU Lot Number: 15.Rv.2.12.3.Ag85

Species: M. tuberculosis

Strain: H37Rv

Purification Information

Starting material: ConA flow through of 40% ammonium sulfate cut (163.92mg)

Starting material (select one):

X CFP: Filter Sterilized Yes

Cells: Irradiated N/A

Viability Test Performed: N/A

Protocol used (SOP #'s): PP020.3 Date started: 11 / 16 / 2015 Date completed: 12 / 3 / 2015

Notebook; page(s): Megan Lucas, BEI Production #2, Starting material: pg. 30-34, Purification: pg. 70-73

Additional notes (if applicable): Phenyl sepharose binding buffer was changed to 50 mM, pH 6.8 and a larger

60mL column was used.

Quality Control Information

Clarity of product/suspension after dialysis: clear

BCA: 5.997 μg/μl Notebook and page(s): _

MCL, BEI Production #2; pg 73

Total Protein: 42 mg

Silver Stain Date: 12 / 2 / 2015 Notebook and page(s):

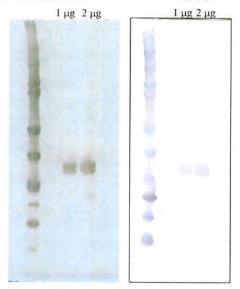
MCL, BEI Production #2; pg 73 Western blot Date: 12 / 2 / 2015 Antibody used: CS-90

Notebook and page(s): MCL, BEI Production #2; pg 73

Mass Spectrometry information/file (if applicable): N/A

QC Gel and Blot:

CS-90



Aliquot Information:

(Research Associate)

84 x 500 µg

12/3/15

(date)