SUPPORTING INFECTIOUS DISEASE RESEARCH

Anthrax Lethal Factor (LF-HMA), Recombinant from Bacillus anthracis

Catalog No. NR-4367

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

Product Description: Recombinant anthrax LF-HMA was produced in a non-sporulating avirulent strain of *Bacillus anthracis* lacking both of the wild type plasmids, pX01 and pX02. Recombinant LF-HMA was purified using conventional chromatographic techniques. The resulting purified protein lacks all other anthrax virulence factors. Note: NR-4367 was vialed using the same bulk LF-HMA as NR-142 and NR-4368. NR-142, NR-4367 and NR-4368 showed similar cytotoxic activity in side-by-side tests.

Lot: 1722B7

Manufacturing Date¹: 01AUG2002

TEST	SPECIFICATIONS	RESULTS	
Appearance ²	Clear and colorless	Clear and colorless	
SDS-PAGE (Coomassie Blue Densitometer Scan)	 ~ 80–90 kDa band is ≥ 90% of total density 	~ 80–90 kDa band is 96% of total density	
HPLC	Report results (% of protein in full length peak)	14% of protein in full length peak	
Electrospray Mass Spectrometry	Report results (expected MW is 90,496 Da)	5 components: 88,000–91,000 Da	
Concentration by Modified Bradford Assay ^{2,3}	Report results	0.77 mg per mL	
Functional Activity FRET (fluorescence resonance energy transfer) protease assay	Report results (rate of cleavage of MAPKKide™ peptide)	0.377 units/mg	
Cytotoxicity assay [determination of effective concentration (EC ₅₀) by titration of LF-HMA in J774A.1 macrophage cells]	LF-HMA with 1 μ g/mL PA: EC ₅₀ \leq 500 pM 1 μ g/mL LF-HMA alone: EC ₅₀ \geq 10,000 pM 1 μ g/mL PA alone: EC ₅₀ \geq 10,000 pM	33 pM ≥ 10,000 pM ≥ 10,000 pM	
Microbial Content ⁴	No detectable colony-forming units in 0.2 mL final product	No detectable colony-forming units in 0.2 mL final product	
Endotoxin Content (Limulus Amoebocyte Lysate Assay)	< 0.5 EU endotoxin per µg protein	0.003 EU endotoxin per µg protein	
Absorbance Ratio (OD ₂₈₀ /OD ₂₆₀)	≥ 1.7	2.0	
Absorbance Ratio (OD ₂₈₀ /OD ₃₂₀)	≥ 10	23	

¹Stored as frozen bulk LF-HMA from August, 2002 until November, 2006. Frozen bulk LF-HMA was thawed and aliquoted on 27 November, 2006. Lyophilization was completed on 29 November, 2006.

²Prior to lyophilization

³Using BSA as a standard

⁴Performed on bulk LF-HMA prior to being frozen in 2002

Date: 05 JUL 2012

Signature:	Dorothy	C.	you	ng
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Title: Technical Manager, BEI Authentication or designee

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