



CERTIFICATE OF ANALYSIS

RSE/EVV RATIO: The potency of this standard in Endotoxin units, (EU) has been determined to be 10 EU/ng by the method formerly described in Appendix C (**Gel-clot Technique**) of the GUIDELINE ON VALIDATION OF THE LIMULUS AMEBOCYTE LYSATE TEST AS AN END PRODUCT ENDOTOXIN TEST FOR HUMAN AND ANIMAL PARENTERAL DRUGS, BIOLOGICAL PRODUCTS, AND MEDICAL DEVICES, published by the U.S. Food and Drug Administration.

EVV Lot: EVV72863 EVV Exp. Date: Oct 2019

LAL Reagent Lot: J3463L LAL Exp. Date: Jun 2021

RSE Lot: H0K354 RSE/EVV Ratio: 10 EU/ng Vial contents: 10000000 EU/vial

Geometric Mean Sensitivity with RSE: 0.125 EU/mL

STORAGE: See package insert.

CAUTION: DO NOT FREEZE ENDOTOXIN SOLUTIONS

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Date: 10 Apr 2018

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