# Certificate of Compliance 

Corning Incorporated - Life Sciences
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Refer to website for regional contact information.
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Quality Management System - Complies with the current version of the EN ISO 13485 Standard.

Non-Pyrogenic - Tested and met the criteria established in the current version of ANSI/AAMI ST 72, "Bacterial Endotoxins - Test methodologies, routine monitoring, and alternatives to batch testing" and USP <85>, "Bacterial Endotoxins Test". The acceptance level for product is $\leq 0.02 \mathrm{EU} / \mathrm{ml}$ or $\leq 0.8 \mathrm{EU} / \mathrm{device}$.

DNase/RNase Free - Tested by nuclease assay method and found to be free of detectable DNase/RNase contamination. The assay detection limit is $10^{-5}$ Kunitz units/ul for DNase and $10^{-9}$ Kunitz units/ul for RNase.

Human DNA Free - Tested by PCR method and found to be free of detectable human DNA contamination. The assay detection limit is hDNA 5pg.

Sterility - Product has been sterilized and dosimetrically released per the requirements of ANSI/AAMI/ISO 11137, "Sterilization of health care products- Radiation". Products meet a minimum Sterility Assurance Level (SAL) of $10^{-3}$.

Quality Control Testing - Representative production samples are collected and inspected in accordance with current applicable product specifications. Inspection records are reviewed and approved by qualified personnel for product release. Key inspections and inline tests are listed below:

Visual Inspection - Pass
Functional Test - Pass

- This product met Corning Incorporated - Life Sciences' high standards of quality at the time of batch/lot release.


Anthony Sloan
Quality Manager

