



Certificate of Conformance

February 2, 2021

P/N	Lot Number	Description
56695-01	702156	TRGH CLR 10X200 FIXINS LID IND

Hamilton Company confirms that the products referenced are manufactured to conform to the claims listed in this certification.

- **Produced under Hamilton Production Specification**

The Hamilton Company production process includes a 100% automated inspections. Additionally Quality Control Representatives measure and assess samples pulled from each production lot.

- **Produced in Clean Environment**

The Hamilton Company Reno cleanroom is certified to ISO 14644-1 Class 8 for clean environment particulate load limits.

- **RNase-free and DNase-free**

Analysis and test method used:

For each test, a substrate is reconstituted with a buffer and distributed to a 96-well plate. The test samples, negative control and positive controls are added and the plate is incubated. The results are then read on a fluorometer. Samples that noticeably fluoresce compared to the negative control are considered contaminated. The test can detect as little as 3.5×10^{-7} units (~ 0.5 pg) or RNase A and 0.0005 U (~ 10 pg) of DNase I.

- **Endotoxin/Pyrogen-free**

Analysis and test method used:

FDA Guideline – USP <85>, AAMI ST72: 2011

Samples are extracted for an hour, they are tested in duplicate against a standard curve ranging from 5.0 to 0.005 EU/mL. There are two wells of sample and two wells of sample spiked with the 5.0 EU/mL standard endotoxin. A chromogen is used which changes color if endotoxin is present in the sample. The plate reader determines the amount of endotoxin based off the samples chromogen color and amount used to extract each device/product.

- **Human DNA-Free and PCR-Inhibitor-Free**

Analysis and test method used:

Samples are extracted in sterile DNA-free water. Part of the extracted sample is exposed to PCR reaction reagents containing primers for human genomic DNA to test for the presence of human

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DNA. Part of the extracted sample is added to a PCR reaction to test for PCR inhibition. PCR reactors contain Taq DNA polymerase, 10X PCR buffer, MgCl₂, dNTPs and human primers. Total reaction volume is 30µL. 30 PCR cycles are run. The sample reactions are run on an agarose gel and each product result was compared to both positive and negative controls. Test sensitivity is 1pg – 2pg.

Facility and product testing is performed on a regular basis by a third party testing facility.



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