## Lyo-Ready Direct RNA-DNA qPCRCatalog No:MDX133Lot No:B129320Saliva, 4xStorage Conditions:-20°CFor research or further manufacturing use onlyExpiry date:August 2026

## **Quality Control Parameters**

Analysis	Specification	Result
Functional	Quantitative real-time PCR analysis amplifying a target gene from a dilution series of mouse RNA under standard cycling conditions. Pass Criteria:	Passed
	Amplification profile of a 1:10 dilution must be consistent for the test and reference sample within $\leq$ 0.5 Cq difference.	
	The end florescence of the 1:10 dilution must be consistent for the test and reference sample within $\leq$ 0.10 difference.	
DNA contamination	Quantitative PCR analysis with no template. Presence of <i>E. coli</i> and mouse genomic DNA checked. Test sample must amplify in concordance with control sample.	Passed
	Pass Criteria: Amplification traces must overlay with the negative control.	
DNase contamination	DNase contamination is measured as DNA substrate degradation against a DNase I dilution series by agarose gel electrophoresis.	Passed
	Limit of detection: 6.25 x 10 <sup>-4</sup> KU DNase I.	
	Pass Criteria: No detectable degradation.	
RNase contamination	Quantitative PCR analysis with high and low RNase standards.	Passed
	Limit of detection: 9.7 x 10 <sup>-3</sup> ng/μL RNase <u>Pass Criteria</u> :	
	Test sample must show less RNase activity than the limit of detection.	

QA / QC Representative:

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Date: 24<sup>th</sup> July 2024

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