Lyo-Ready Direct RNA-DNA qPCRCatalog No:MDX133Lot No:B134340Saliva, 4xStorage Conditions:-20°CFor research or further manufacturing use onlyExpiry date:January 2027

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. <u>Pass Criteria:</u>	Passed
	Amplification profile of a 1:10 dilution must be consistent for the test and reference sample within ≤ 0.5 Cq difference. The end florescence of the 1:10 dilution must be consistent for the test and reference sample within ≤ 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. <u>Pass Criteria</u> : Amplification traces must overlay with the negative control.	Passed
DNase contamination	DNase contamination is measured as DNA substrate degradation against a DNase I dilution series by agarose gel electrophoresis. Limit of detection: 6.25 x 10 ⁻⁴ KU DNase I. <u>Pass Criteria</u> : No detectable degradation.	Passed
RNase contamination	Quantitative PCR analysis with high and low RNase standards. Limit of detection: 9.7 x 10 ⁻³ ng/µL RNase <u>Pass Criteria</u> : Test sample must show less RNase activity than the limit of detection.	Passed

QA / QC Representative:

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