Version: v04

Aptamer Taq HS (Glycerol-Free), 50 U/μL

Catalog No:	MDX015
Lot No:	EN094-B127550
Storage Conditions:	-20°C
Component Lot No:	GFIH-324105A
Expiry date:	June 2026

For Research and Further Manufacturing use only

Quality Control Parameters

Analysis	Specification	Result
Functional	Activity is measured as DNA polymerase units by quantitative PCR analysis against a reference Taq DNA polymerase standard curve. <u>Pass Criteria</u> : Activity must be between 50 and 60 U/µL	59.30 U/μL
Functional	Aptamer specificity is	Passed
Glycerol content	Glycerol concentration is determined by spectrophotometric measurement of a colorimetric product from a coupled enzymatic reaction. Pass Criteria: Glycerol content <0.02 %	Passed
Purity	Purity is measured as a percentage of total protein by quantitative gel electrophoresis on Bioanalyzer (Agilent). <u>Pass Criteria</u> : >50 %	98.6 %
DNA contamination	DNA contamination is measured by quantitative PCR on <i>E. coli</i> and mouse genomic DNA specific targets. <u>Pass Criteria</u> : Amplification traces must overlay with the negative control.	Passed

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COA No: CA_XBE-0068

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	DNase contamination is measured as DNA substrate degradation against a	
DNase contamination	DNase I dilution series by agarose gel electrophoresis.	
	Limit of detection: 6.25×10^{-4} KU DNase I.	Passed
	Pass Criteria:	
	No detectable degradation.	
RNase contamination	RNase contamination is measured by quantitative PCR against RNase standards.	
	Limit of detection: 9.7 x 10^{-3} ng/µL RNase.	Passed
	Pass Criteria:	
	No detectable degradation.	

QA / QC Representative: Ringthen

X. Chen

Date: 22nd May 2024

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Enzyme Dilution Buffer

For research or further manufacturing use only

Catalog No:	MDX015
Lot No:	EN094-B127550
Storage Conditions:	-20°C
Lot number:	TDB-224105A
Expiry date:	June 2026

Quality Control Parameters

Enzyme Dilution Buffer is a glycerol-free 10x buffer used for the dilution of Taq antibody or reverse transcriptase.

Analysis	Specification	Result
Functional	A 3Kb fragment was amplified with a dilution series of Enzyme Dilution Buffer, using standard conditions and 30 cycles.	
	A 3Kb fragment was amplified with a dilution series of <i>Taq polymerase</i> , using standard conditions and 30 cycles.	Passed
	Single distinct bands were observed with agarose gel electrophoresis (ethidium stained).	
Glycerol determination	Glycerol content is < 0.2% determined by spectrophotometric analysis and comparison to a standard curve.	Passed
DNA contamination	Quantitative PCR analysis with no template. Presence of <i>E. coli</i> and mouse genomic DNA checked. Test sample must amplify in line with a reference sample.	Passed
DNase contamination	Incubation of a 1Kb double stranded DNA fragment. Incubation for 4 hours at 37°C with dilution series of DNase I. Analysed by agarose gel electrophoresis. Test sample must show less degradation than the limit of detection 6.25 x 10^{-4} KU/µL.	Passed
RNase contamination	Quantitative PCR analysis with high and low RNase standards. Test sample must show less RNase activity than the limit of detection: $9.7x10^{-3}$ ng/µl RNase.	Passed

QA / QC Representative: Ring has

X. Chen

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