



## Certificate of Analysis

COA No: CA\_BEM-0010

Version: 01

# Reverse Transcriptase

For Research Use Only

Storage Conditions:	-20°C
Lot number:	RTS-313110
Expiry date:	October 2015

## Quality Control Parameters

Analysis	Specification	Result
Functional	Quantitative PCR analysis amplifying 6 genes from a dilution series of mouse RNA under standard conditions. Cq and melt profiles must be consistent for the test and reference sample with 0.5+/- Cq variance.	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 control 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 $2.5 \times 10^{-3}$ U DNase I.	Passed
RNase contamination	Quantitative PCR analysis with high and low RNase standards. Test sample must show less RNase activity than the limit of detection $9.7 \times 10^{-3}$ ng/ $\mu$ l RNase.	Passed

Authorised by Jade James

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## Certificate of Analysis

COA No: CA\_BMM-0010

Version: 01

### MyTaq One-Step RT-PCR Mix

For Research Use Only

Storage Conditions: -20°C

Lot number: MTOS-213110

Expiry date: October 2015

### Quality Control Parameters

Analysis	Specification	Result
Functional	Fragments of sizes 1000bp and 1400bp were amplified with a dilution series of mouse RNA, using standard conditions and 45 cycles. Single distinct bands were observed with agarose gel electrophoresis (ethidium stained).	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 control 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 $2.5 \times 10^{-3}$ U DNase I.	Passed
RNase contamination	Quantitative PCR analysis with high and low RNase standards. Test sample must show less RNase activity than the limit of detection $9.7 \times 10^{-3}$ ng/ $\mu$ l RNase.	Passed

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## Certificate of Analysis

COA No: CA BE-0031

Version: 01

### RNase Inhibitor

For Research Use Only

Storage Conditions: -20°C

Lot number: RI-113110

Expiry date: October 2015

### Quality Control Parameters

Analysis	Specification	Result
Inhibition	Test level of inhibition by incubating total RNA with concentration gradient of RNase A. Bands were observed with agarose gel electrophoresis (ethidium stained).	Passed

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## Certificate of Analysis

COA No: CA\_BS-0020

Version: 01

### DEPC Water

For Research Use Only

Storage  
Conditions:

-20°C

Lot number:

DWT-213110

Expiry date:

October 2015

### Quality Control Parameters

Analysis	Specification	Result
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 control 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 $2.5 \times 10^{-3}$ U DNase I.	Passed
RNase contamination	Quantitative PCR analysis with high and low RNase standards. Test sample must show less RNase activity than the limit of detection $9.7 \times 10^{-3}$ ng/ $\mu$ l RNase.	Passed

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