

IMMOLASE [™] DNA Polymerase	Storage Conditions:	-20°C
	Lot number:	IM-314107
For Research Use Only	Expiry date:	August 2016

Quality Control Parameters

Analysis	Specification	Result
Activity	Quantitative PCR analysis amplifying 1 gene from a dilution series of	Passed
	enzyme under standard conditions. Cq and melting profiles must be	
	consistent for the test and reference sample with 0.5+/- Cq variance.	
Sensitivity	Quantitative PCR analysis amplifying 1 gene from a dilution series of	Passed
	mouse cDNA under standard conditions. Cq and melting profiles	
	must be consistent for the test and reference sample with 0.5+/- Cq	
	variance.	
	A 3Kb fragment is amplified with a dilution series of Lambda DNA,	
	using standard conditions and 30 cycles. Single distinct bands were	
	observed with agarose gel electrophoresis (ethidium stained).	
Heat activation	A 125bp fragment is amplified with a dilution series of enzyme, using	Passed
	4 heat activation times and 30 cycles. Single distinct bands were	
	observed, at the appropriate activation time, with agarose gel	
	electrophoresis (ethidium stained).	
DNA contamination	Quantitative PCR analysis with no template. Presence of <i>E. coli</i> and	Passed
	mouse genomic DNA checked. Test sample must amplify in line with	
	a reference sample.	
DNase contamination	Incubation of a 1Kb double stranded DNA fragment. Incubation for	Passed
	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 2.5×10^{-3} U DNase.	

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ImmoBuffer	Storage Conditions:	-20°C
	Lot number:	IB-314107
For Research Use Only	Expiry date:	August 2016

Quality Control Parameters

Analysis	Specification	Result
Functional	Fragment of size 800bp was amplified with a dilution series of IMMOLASE [™] , using standard conditions and 30 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 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 2.5×10^{-3} U DNase.	Passed

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MgCl ₂ Solution, 50mM	Storage Conditions:	-20°C
laigel 2 Solution, Sollina	Lot number:	MG-314107
For Research Use Only	Expiry date:	August 2016

Quality Control Parameters

Analysis	Specification	Result
Functional	Fragments of sizes 800bp and 3000bp are amplified with a dilution series of BIOTAQ [™] DNA Polymerase, using standard conditions and 30 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 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 2.5 x 10 ⁻³ U DNase.	Passed

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