

Product Information	
qScript® MicroRNA cDNA Synthesis	
Part Number	95107-100
Number of Reactions	100 Reactions
Reaction Size	20 µL
Storage Temperature	-25°C to -15°C
Lot Number	66170557
Reference Number	081120, 020420, 101419, 071020, 317894469, 321076298
Expiration Date	10/31/2022

Product Description:

The qScript microRNA cDNA Synthesis Kit is an optimized reagent system that reverse transcribes small single-stranded RNA into 5'-labeled cDNA using either total RNA or miRNA enriched samples.

Single-stranded RNA is first polyadenylated by poly(A) polymerase before reverse transcription into universal cDNA using high performance qScript RT with a proprietary adapter oligo(dT) primer. The universal cDNA template enables simple and cost-effective microRNA profiling when used together with wet-lab validated PerfeCta microRNA Assays, PerfeCta Universal PCR Primer and PerfeCta SYBR Green SuperMix.

Component Part Numbers:

84002 qScript RT 0.10 mL

- 84173 Poly (A) Tailing Buffer (5X)
- 84175 Poly (A) Polymerase
- 84177 MicroRNA cDNA Reaction Mix
- 84179 PerfeCta Universal PCR Primer
- 84181 PerfeCta Human Positive Control Primer
- 84007 Nuclease Filled Water 1.50 mL

Product Specifications				
95107				
Assay	qScript microRNA cDNA Synthesis Kit	β-actin SYBR Green qRT-PCR Assay for qScript Reverse Transcriptase	DNase	RNase
Result	Pass	Pass	Pass	Pass

Quality Control Analysis and Specifications:

Nuclease Assay

DNase: DNase activity must be below the detectable limits of 100 pg DNase I equivalent as assayed using a fluorogenic substrate following a 1 hour incubation at 37°C with each kit component at 1X concentration.

RNase: RNase activity must be below the detectable limits of 1 pg RNase A equivalent as assayed using a fluorogenic substrate following a 1 hour incubation at 37°C with each kit component at 1X concentration.

qScript microRNA cDNA Synthesis Kit Must have detection of microRNA from 100 ng to 10 pg. Correlation of determination (R^2) \geq 0.990 from Ct standard curve analysis. Slope from Ct standard curve analysis between -3.20 and -3.75. The +PAP reactions must show a \geq 11 Ct difference from the “no-PAP” reactions. There must be at least three negative template controls with “undetermined” values.

β-actin SYBR Green qRT-PCR Assay for qScript Reverse Transcriptase: Detection of β-actin mRNA from 100ng to 100fg of total RNA. Coefficient of determination (R^2) \geq 0.990 with a slope analysis between -3.20 and -3.70

Limitations of Use

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