

Product Information	
qScript® Flex cDNA Kit	
Part Number	95049-100
Number of Reactions	100 Reactions
Reaction Size	20 µL
Storage Temperature	-25°C to -15°C
Lot Number	022057
Reference Number	012517
	18817
	014664
	080817
Expiration Date	071417
	02/28/2019

Product Description:

The qScript Flex cDNA Synthesis Kit is configured in separate components for supporting multiple RT priming strategies. The kit provides optimized, flexible components for priming with oligo-dT(20), random hexamers, gene-specific primer (GSP), or any combination thereof. This unique 5X master mix of buffer, magnesium, stabilizers and dNTPs simplifies reaction assembly and ensures robust and reproducible synthesis of first strand product from 10 pg to 4µg of total RNA or purified polyA+ RNA template. The resulting cDNA product is directly compatible with current real-time RT-PCR methods or end-point RT-PCR. Length of cDNA product is dependent upon priming strategy and quality of the RNA template. Oligo dT or GSP can be used for two-step RT-PCR of RNA targets up to 12 kb. Random primer is suitable for RT-PCR of RNA targets less than 1 kb. Any of the three priming methods are suitable for real-time quantitative RT PCR (RT-qPCR).

Component Part Numbers:

- 84002 qScript RT 0.10 mL**
- 84043 5X qScript Flex Reaction Mix 0.40 mL**
- 84044 Oligo dT 0.20 mL**
- 84045 Random Primer 0.20 mL**
- 84046 GSP Enhancer 0.20 mL**
- 84007 Nuclease Free Water 1.50 mL**

Product Specifications				
95049				
Assay	Functional PCR Assay for Flex Kit	β-actin SYBR Green qRT-PCR Assay for qScript Reverse Transcriptase	DNase	RNase
Result	Pass	Pass	Pass	Pass

Quality Control Analysis and Specifications:

Functional PCR Assay for Flex Kit Reaction Mix (5X): Detection of β-actin mRNA from 100 ng to 100 fg of total RNA. Coefficient of determination (R^2) ≥ 0.990 with a slope analysis between -3.20 and -3.70

Single bands visible at 6kb and 8kb from 35 cycles of PCR using 100 ng Oligo-dT cDNA

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.

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