# Quantabio

# qScript® microRNA cDNA Synthesis Kit

Cat. No 95107-025 S 95091-025

Size: 25 x 20 µL reactions 100 x 20 µL reactions Store at -25°C to -15°C

MicroRNAs (miRNAs) are small (approximately 22-24 nucleotide) non-coding RNA molecules that are important regulators of gene expression in a variety of organisms. The distinct expression patterns of miRNAs in different tissues, cell development stages, and disease models suggest that miRNAs play important roles in biological processes. The qScript<sup>™</sup> microRNA cDNA Synthesis Kit provides a convenient, reliable, and highly sensitive system for conversion of miRNAs into cDNA from total RNA samples. The kit is used in concert with pre-validated PerfeCTa<sup>®</sup> microRNA Assays and the PerfeCTa SYBR<sup>®</sup> Green SuperMix family of qPCR products as part of the qScript microRNA Quantification System providing a highly sensitive method for quantification and profiling of miRNAs.

# Description

The qScript microRNA cDNA Synthesis Kit has been designed to convert miRNAs into cDNA starting from total RNA or RNA preparations pre-enriched for miRNAs. MicroRNAs are not polyadenylated in nature. With the qScript microRNA cDNA Synthesis Kit miRNAs are polyadenylated in a poly(A) polymerase reaction. qScript Reverse Transcriptase and other necessary reagents for cDNA synthesis are subsequently added to convert the poly(A) tailed miRNAs into cDNA using an oligo-dT adapter primer. The adapter primer has a unique sequence at its 5' end which allows amplification of cDNAs in real-time qRT-PCR reactions.

The kit includes a Human Positive Control Primer that can be used to quantify the small nucleolar RNA SNORD44 which is ubiquitously expressed in most human tissues. In addition, the kit contains 20% extra Poly(A) Tailing Buffer and microRNA cDNA Reaction Mix to accommodate the use of no poly(A) polymerase and no reverse transcriptase control reactions.

Individual miRNAs are quantified in real-time SYBR Green qRT-PCR amplification reactions with the desired PerfeCta microRNA Assay and the PerfeCta Universal PCR Primer (specific to the unique sequence of the oligo-dT adapter primer). The pre-designed and validated PerfeCta microRNA Assays provide maximum sensitivity and specificity in qRT-PCR amplification and quantification of microRNAs.

For a complete list of available microRNA assays please visit www.quantabio.com/microrna.

# Components

	95107-025	95107-100
Poly(A) Tailing Buffer (5X)	1 x 60 µL	1 x 240 µL
Poly(A) Polymerase	1 x 25 µL	1 x 100 µL
microRNA cDNA Reaction Mix	1 x 270 µL	1 x 1080 µL
qScript Reverse Transcriptase	1 x 25 µL	1 x 100 µL
PerfeCTa Universal PCR Primer	1 x 125 µL	1 x 500 µL
PerfeCTa Human Positive Control Primer	1 x 25 µL	1 x 100 µL
Nuclease-Free Water	1 x 1.5 mL	1 x 1.5 mL

# Storage and Stability

Store components in a constant temperature freezer at -25°C to -15°C upon receipt. For lot specific expiry date, refer to package label, Certificate of Analysis or Product Specification Form.

# **Reaction Protocol**

#### Poly(A) Tailing Reaction

- 1. Thaw all components (except enzyme), mix thoroughly, and centrifuge briefly. Keep on ice before use.
- 2. Add the following components to a 0.2 mL PCR tube or 96-well plate on ice:

Component	Volume
Poly(A) Tailing Buffer (5X)	2 µL
RNA (up to 1 ug total RNA or miRNA-enriched RNA)	up to 7 µL
Nuclease-Free Water	variable
Poly(A) Polymerase	<u>1 µL</u>
Final Volume	10 µL

- 3. Mix components by gentle vortexing and centrifuge briefly to collect the contents.
- Incubate 60 minutes at 37°C followed by 5 minutes at 70°C. Briefly centrifuge to collect the contents. Keep on ice before cDNA synthesis. The
  incubation at 37°C can be shortened to 20 minutes when using 100 ng or less of total RNA.



#### First-Strand cDNA Synthesis Reaction

5. Setup the cDNA Synthesis Reaction as follows:

Component	Volume	
Poly(A) Tailing Reaction (from step 4 above)	10 µL	
microRNA cDNA Reaction Mix	9 µL	
<u>qScript Reverse Transcriptase</u>	<u>1 µL</u>	
Final Volume	20 µL	

- 6. Mix by gentle vortexing and then briefly centrifuge to collect contents.
- 7. Incubate 20 minutes at 42°C followed by 5 minutes at 85°C.

cDNA is now ready to use for qRT-PCR. If desired, the microRNA cDNA product can be diluted with 10 mM Tris-HCl (pH 8.0), 0.1 mM EDTA. Diluted cDNA is stable for several months at +4°C. For long-term storage store the cDNA at -20°C.

#### Real-Time SYBR Green qRT-PCR Amplification of MicroRNAs

Real-time SYBR Green qRT-PCR is performed using 200 nM of each PerfeCta microRNA Assay Primer and PerfeCta Universal PCR Primer along with the appropriate PerfeCta SYBR Green SuperMix product depending on the instrument platform being used.

For each qRT-PCR reaction add the following components:

Component	Volume	
PerfeC⊤a SYBR Green SuperMix (2X)	25 µL	
PerfeC⊤a microRNA Assay Primer (10 µM)	1.0 µL	
PerfeC⊤a Universal PCR Primer (10 µM)	1.0 µL	
MicroRNA cDNA	up to 23 μL	
Nuclease-Free Water	variable	
Final Volume	50 µL	

The amount of microRNA cDNA can be varied depending on the expression level of the miRNA. As a starting point use about 1ng of total RNA equivalent per qRT-PCR reaction. For miRNAs expressed at low levels you may use 10 ng of total RNA equivalent per qRT-PCR reaction. For most applications 20 to 25 µL qRT-PCR reaction volumes are suitable but reaction volumes can be scaled according to your needs.

2-Step Cycling Protocol	
Pre-incubation/activation:	95°C, 2 minutes
PCR (40 cycles)	
Denature:	95°C, 5 seconds
Anneal:	60°C, 30 seconds (collect fluorescence data)

#### 3-Step Cycling Protocol (Optional)

Pre-incubation/activation:	95°C, 2 minutes
PCR (40 cycles)	
Denature:	95°C, 5 seconds
Anneal:	60°C, 15 seconds
Extend:	70°C, 15 seconds (collect fluorescence data)
	· · · · · · · · · · · · · · · · · · ·

Use of a 3-step cycling protocol or slightly higher annealing temperature may improve the specificity of some assays. Melt curve analysis is optional. Most microRNA qRT-PCR reactions will produce a single, slightly broader first-derivative melt peak compared to reactions using two gene-specific primers due to slight heterogeneity in the poly(A) tail length.

#### **Relevant Products**

- PerfeCta microRNA Assays (<u>www.quantabio.com/microrna</u>)
- PerfeCTa Universal PCR Primer (Cat. No. 95109)
- PerfeCta SYBR Green SuperMix family of qPCR products (<u>www.quantabio.com/group/qpcr\_sybr.php</u>)

# Quantabio

# Limited Label Licenses

Use of this product signifies the agreement of any purchaser or user of the product to the following terms:

- 1. The product may be used solely in accordance with the protocols provided with the product and this manual and for use with components contained in the kit only. QIAGEN Beverly, Inc. grants no license under any of its intellectual property to use or incorporate the enclosed components of this kit with any components not included within this kit except as described in the protocols provided with the product, this manual, and additional protocols available at <u>www.guantabio.com</u>. Some of these additional protocols have been provided by Quantabio product users. These protocols have not been thoroughly tested or optimized by QIAGEN Beverly, Inc.. QIAGEN Beverly, Inc. neither guarantees them nor warrants that they do not infringe the rights of third-parties.
- 2. Other than expressly stated licenses, QIAGEN Beverly, Inc. makes no warranty that this kit and/or its use(s) do not infringe the rights of third-parties.
- 3. This kit and its components are licensed for one-time use and may not be reused, refurbished, or resold.
- 4. QIAGEN Beverly, Inc. specifically disclaims any other licenses, expressed or implied other than those expressly stated.
- 5. The purchaser and user of the kit agree not to take or permit anyone else to take any steps that could lead to or facilitate any acts prohibited above. QIAGEN Beverly, Inc. may enforce the prohibitions of this Limited License Agreement in any Court, and shall recover all its investigative and Court costs, including attorney fees, in any action to enforce this Limited License Agreement or any of its intellectual property rights relating to the kit and/or its components.

©2018 QIAGEN Beverly Inc. 100 Cummings Center Suite 407J Beverly, MA 01915

Quantabio brand products are manufactured by QIAGEN, Beverly Inc.

Intended for molecular biology applications. This product is not intended for the diagnosis, prevention or treatment of a disease.

qScript and PerfeCTa are registered trademarsk of QIAGEN Beverly, Inc. SYBR is a registered trademark of Molecular Probes, Inc.