LaboPass[™] cDNA Synthesis Kit

Cat. No CMRTK001 50 reactions
Cat. No CMRTK002 100 reactions

Lot No.

Note: For laboratory use only

Description

- The LaboPassTM cDNA synthesis kit is optimized for efficient and reproducible synthesizing first-strand cDNA from total RNA or poly(A) RNA. This kit provides the all components necessary for synthesis of cDNA including reverse transcriptase, dNTPs, reaction buffer, RNase inhibitor, primer and nuclease-free water.
- Included M-MuLV Reverse Transcriptase in this kit is a recombinant form of the reverse transcriptase
 from the Moloney Murine Leukemia Virus (M-MuLV) which possesses enhanced cDNA synthesis activity
 and reduced RNase H activity. Reduction of RNase H activity enables higher yield of full-length cDNA
 transcripts and increased thermostability.

Contents

	50 rxn	100 rxn
• Reverse Transcriptase (200 U/µI)	50 µl	100 µl
• 5X RT Buffer	200 µl	400 µl
 dNTP (each 10 mM) 	50 µl	100 µl
 RNase inhibitor (40 U/µI) 	50 µl	100 µl
 Oligo (dT)₁₈ (100 μM) 	50 µl	100 µl
• Random hexamer (0.2 $\mu g/\mu I$)	50 µl	100 µl
 Nuclease-free water 	1 ml	1.5 ml
Store at -20°C		

Applications

- cDNA synthesis for RT-PCR and RT-qPCR
- · cDNA synthesis for Cloning

Storage and Stability

LaboPassTM cDNA synthesis kit is stable for 1 year when stored at -20°C.

Quality Control

Each lot of Reverse Transcriptase, reaction buffer, RNase inhibitor and dNTPs is tested for contamination such as nuclease.

General cDNA synthesis protocol

- 1. Add the following components into a sterile, nuclease-free tube for each 20 μ l reaction :
 - RNA 1-8 μ l 0.1 ng \sim 5 μ g total RNA or 1 ng \sim 500 ng poly(A) RNA

• Oligo (dT) or Random primer
• dNTP (each 10 mM)
• Nuclease-free water

1 µl
1 µl
1 µl
Up to 10 µl

- 2. Heat for 5 min at 65°C, quick chill on ice and briefly spin-down.
- 3. Add the following components to the above mixture and gently mix:

5X RT reaction buffer 4 µl
 RNase inhibitor (40 U/µl) 1 µl
 M-MulV Reverse Transcriptase 1 µl
 Nuclease-free water Up to 10 µl

- 4. Incubate at 42° C, for $30 \sim 60$ min.
- 5. Inactivate enzyme by heating at 70°C for 15 min.
- 6. Store at -20°C until ready for use.

