[Cat. No.] Please refer to the Ordering Information

Introduction

AccuPower® RT-PCR PreMix utilizes one-step RT-PCR performing cDNA synthesis and PCR in a single step, in a single tube. It can provide reduced possibility of cross-contamination and errors. This product contains vacuum-dried components including *M-MLV* Reverse Transcriptase, reaction buffer, RNase inhibitor, *Top* DNA Polymerase, stabilizer, and dNTPs. It simplifies preparation of reverse transcription reaction mixture and PCR mixture by adding template RNA and primers without any extra process. Furthermore, by using the RNase H⁺ of *M-MLV* Reverse Transcriptase, template RNA is removed after the cDNA synthesis to minimize the effect of residual template RNA in PCR. After the reaction, samples can be applied directly on agarose gel for analysis due to included tracking dye.

Applications

- First-strand synthesis of cDNA from RNA molecules (RT)
- RT-PCR
- cDNA library construction
- Gene expression analysis

Features & Benefits

- Ease-of-use: Reactants are individually packaged in each of the PCR tubes, it allows any user simply perform cDNA synthesis and PCR in one tube by adding template RNA and its specific primers.
- Stability: Included stabilizer and RNase inhibitor provides high resistance to degradation.
- Reproducibility: Mass production under ISO 9001 quality system allows minimized deviation between lots and reproducible results in replicated tests performed under same conditions and variation.

Composition

Composition	Concentration		
M-MLV Reverse Transcriptase	200 U		
5X Reaction buffer	1X		
DTT	0.25 mM		
dNTPs (dATP, dCTP, dGTP, dTTP)	Each 250 µM		
RNase inhibitor	1 U		
<i>Top</i> DNA Polymerase	1 U		
Stabilizer and tracking dye	1X		

Specifications

Top DNA Polymerase				
5' to 3' exonuclease activity	No			
3' to 5' exonuclease activity	No			
3'–A overhang	Yes			
Fragment size	Up to 5 kb			

Storage

Store at -20°C. If stored in the recommended temperature, this product will be stable until the expiration date printed out on the label.

Online Resources





Visit our product page for additional information and protocols

Ordering Information

Descr	Cat. No.		
	96 tubes	20 µl/rxn	K-2055
0.2 ml thin-wall 8-tube strips with attached cap		50 µl/rxn	K-2057
	480 tubes	20 µl/rxn	K-2055-B
		50 µl/rxn	K-2057-B
0.5 ml thin-wall tubes with attached cap	100 tubes	50 µl/rxn	K-2056
	flat plate	10 µl/rxn	K-2262-1
		20 µl/rxn	K-2262-4
thin-wall 96-well	full-skirted plate	10 µl/rxn	K-2262-2
Inin-wali 96-weli		20 µl/rxn	K-2262-5
	semi-skirted plate	10 µl/rxn	K-2262-3
		20 µl/rxn	K-2262-6
	full-skirted plate	5 µl/rxn	K-2084-1
thin-wall 384-well		10 µl/rxn	K-2084-2
		20 µl/rxn	K-2084-3

Notice

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Explanation of Symbols



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Experimental Procedures

	Steps	Procedure Details				
1	Preparation of reaction mixture	 Add template RNA, primers and nuclease-free water into <i>AccuPower</i>[®] RT-PCR PreMix tubes to make a total volume of 20 μl or 50 μl. Do not include the dried pellet. Amount of template RNA and primer Components 20 μl reaction 				
		Template RNA	Total RNA Poly(A) RNA	0.5-1 μg 0.05-0.1 μg	1-2 μg 0.1-0.2 μg	
		Primers	Gene specific primer	10-30 pmol	20-50 pmol	
		2. Dissolve the vacuum-dried pellet by vortexing or pipetting, and briefly spin down.				
		3. Perform the reac	tion under the following co	Time	Cycles	
		cDNA synthesis	42°C	60 min	1 cycle	
2		Pre-denaturation	42 C 95°C	5 min	1 cycle	
		Denaturation	95°C	10-30 sec	r cyclo	
		Annealing	45-65°C	10-30 sec	30-35 cycles	
		Extension	72°C	1 kb/min	,	
		Final extension	72°C	5 min	1 cycle	
		* Note: Annealing temperature and time need to be optimized for each primer/template combination.				
3	Analyze with gel electrophoresis	 4. After the reaction, maintain the reaction mixture at 4°C. The samples can be stored at -20°C until use. 5. Load samples on agarose gel without adding a loading-dye mixture, and perform gel electrophoresis for analysis. 				

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