



Human Kidney cDNA

Catalog #: HD-901
Quantity: 30 Reactions

Storage Conditions: Store at -20oC. It is good for one year of the date of purchase if stored

properly.

Applications: The cDNA is primed with oligo dT primer and is ideal for gene expression analysis by PCR, characterization of alternative splicing of mRNA, and Gene cloning and target sequencing.

Quality Control: The PCR-Ready cDNA is functionally tested with the control primers in the recommended PCR reaction.

Description: The PCR-ready first strand cDNA is synthesized from high quality RNA isolated from adult human normal healthy tissues. Total RNA used for cDNA synthesis is isolated by modified guanidine thiocyanate techniques and treated with RNase-free DNase. The total RNA was primed with oligo dT primer and reverse transcribed by a reverse transcriptase enable synthesis of full length cDNA up to 8.9kb. Use μl cDNA for each PCR reaction.

The amplification conditions used for amplification of beta-actin as a positive control in a volume of 100µl:

cDNA	1µl
10X polymerase reaction buffer	10µ1
MgCl ₂ , 25mM (2nM final)	7.8µ1
Nucleotide Mix, 10mM (0.2mM final)	2.0µ1
Upstream beta-Actin primer (100μM)	1µl
Downstream beta-Actin primer (100µM)	1µ1
Taq Polymerase (5 units)	1µ1
Water	<u>76.2µ1</u>
	100µl

Program of amplification:

Denaturation 94°c for 2 minutes

25 cycles:

Denaturation 94°c for 1 minute Annealing 60°c for 1 minute Extension 72°c for 2 minutes

Final extension 72°c for 5 min

Hold 4°c

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Zvagen Laboratories, San Diego, California, USA. Telephone: +1 858 546 0720; Fax: +1 858 546 0736: Technical support e-mail address: zinfo@zyagen.com. For additional product and distributor information, see www.zyagen.com