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Product	Size	SKU
VitaScript™ FirstStrand cDNA Synthesis Kit	50 rxn / 20 μl	PCCSKU1301

## **DESCRIPTION**

VitaScript™ First Strand cDNA Synthesis Kit contains:

COMPONENT	VOLUME	DESCRIPTION
VitaScript™ Enzyme Mix	50 µl	contains VitaScript™ Reverse Transcriptase and RNase inhibitor blend
5X VS Reaction Buffer	250 µl	contains dNTPs, MgCl <sub>2</sub> , random 6-mer primers and oligo-dTs in an optimized buffer environment
Nuclease-free water	1 ml	

## GENERAL CONSIDERATIONS

- High purity template RNA is essential for reliable efficient cDNA synthesis. A A<sub>260</sub> / A<sub>280</sub> ratio of 1.7 or higher is strongly recommended.
- The amount of template RNA is depended on the expected copy number of the sequence of interest. In general, 1  $\mu$ g 1 ng of total RNA is recommended, 0.05 100 ng if you are working with isolated mRNA.
- When working with long cDNA synthesis, denaturation of RNA with VS reaction buffer for 5 minutes at 72°C can be applied to remove secondary structures that can impede the reaction.
- This protocol recommends cDNA synthesis for 1 hour at 42°C.
- To enhance the template coverage, the VS reaction buffer also contains random hexamer primers. This provides multiple priming sites along the RNA for the detection of multiple short sequences.

## **REACTION SETUP**

Thaw all components and mix gently. Keep on ice during reaction setup. A control reaction without VitaScript™ Reverse Transcriptase is highly recommended to check for potential DNA contamination.

(Optional) Denature RNA with VS Reaction Buffer for 5 minutes at 72°C. Spin down and instantly put on ice. This can improve transcription for long mRNAs or GC-rich RNA.

- As negative control, replace VitaScript™ Enzyme Mix with 1 µl nuclease-free dH<sub>2</sub>O.
- Mix in a sterile RNase-free tube:

COMPONENT	VOLUME
5X VS Reaction Buffer	4 µl
VitaScript™ Enzyme Mix	1 µl
Total RNA	1-6 µl
Nuclease-free dH <sub>2</sub> O	to 20 µl
total volume	20 µl

STEP	TEMPERATURE	TIME
cDNA Synthesis	42°C	60 minutes
Inactivation of VitaScript™	80°C	10 minutes

• Dilute the reaction with nuclease-free dH<sub>2</sub>O to 200 µl and store at -20°C. Avoid repeated freeze-thaw cycles.

For following PCR applications, the diluted cDNA reaction should represent 10% of the total reaction volume (e.g.  $5 \mu l$  in a 50  $\mu l$  reaction).

