

TransScript® One-Step gDNA Removal and cDNA Synthesis SuperMix

Cat. No. AT311

Storage: at -20°C for two years

Description

Unique genomic DNA remover is combined with *TransScript*® First-Strand cDNA Synthesis SuperMix to achieve simultaneous genomic DNA removal and cDNA synthesis. After cDNA synthesis, gDNA remover and reverse transcriptase are inactivated by heating at 85°C for 5 seconds.

Highlights

- Simultaneous genomic DNA removal and cDNA synthesis in one tube to minimize RNA contamination.
- The product obtained from 15 minutes reaction is used for qPCR; the product obtained from 30 minutes reaction is used for PCR.
- cDNA up to 12 kb.

Applications

Multiple copy and low copy gene detection

Kit Contents

| Component | AT311-02 (50 rxns) | AT311-03 (100 rxns) |
|---|--------------------|---------------------|
| <i>TransScript</i> ® RT/RI Enzyme Mix | 50 µl | 100 µl |
| gDNA Remover | 50 µl | 100 µl |
| 2×TS Reaction Mix | 500 µl | 1 ml |
| Random Primer(N9) (0.1 µg/µl) | 50 µl | 100 µl |
| Anchored Oligo(dT) ₁₈ Primer (0.5 µg/µl) | 50 µl | 100 µl |
| RNase-free Water | 500 µl | 1 ml |

First-Strand cDNA synthesis

1. Reaction Components

| Component | Volume |
|---|----------------------|
| Total RNA/mRNA | 50 ng -5 µg/5-500 ng |
| Anchored Oligo(dT) ₁₈ Primer (0.5 µg/µl) | 1 µl |
| or Random Primer (0.1 µg/µl) | 1 µl |
| or GSP | 2 pmol |
| 2×TS Reaction Mix | 10 µl |
| <i>TransScript</i> ® RT/RI Enzyme Mix | 1 µl |
| gDNA Remover | 1 µl |
| RNase-free Water | to 20 µl |

Optional: for higher efficiency, suggest to mix RNA, primer and water first. Incubate the mixture at 65°C for 5 minutes, on ice for 2 minutes. Then add other components.

2. Incubation

- For anchored oligo(dT)₁₈ primer or GSP, incubate at 42°C for 15 minutes (for qPCR) or incubate at 42°C for 30 minutes (for PCR).
- For random primer, incubate at 25°C for 10 minutes. After that, incubate at 42°C for 15 minutes (for qPCR) or incubate at 42°C for 30 minutes (for PCR).

3. Incubate at 85°C for 5 seconds to inactivate enzymes.

RT-PCR

Reaction Components

| Component | Volume | Final Concentration |
|--|----------|---------------------|
| cDNA | 2 µl | as required |
| Forward Primer (10 µM) | 1 µl | 0.2 µM |
| Reverse Primer (10 µM) | 1 µl | 0.2 µM |
| 2× <i>TransTaq</i> [®] HiFi PCR SuperMix II | 25 µl | 1× |
| ddH ₂ O | Variable | - |
| Total volume | 50 µl | - |

Thermal cycling conditions

| | | |
|---------|------------|----------------|
| 94°C | 2-5 min | } 30-35 cycles |
| 94°C | 30 sec | |
| 50-60°C | 30 sec | |
| 72°C | 1-2 kb/min | |
| 72°C | 5-10 min | |

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