

## FastGene® Scriptase III

| Cat. No. | Pack Size     |
|----------|---------------|
| LS55     | 100 reactions |
| LS55s    | 10 reactions  |

### Storage:

Store at -20°C.

### Description:

FastGene® Scriptase III is an improved version of M-MLV Reverse Transcriptase, which exhibits high thermal stability (~55°C) and low RNase H activity. This product requires the purchase of primers (oligo dT, random hexamer) separately.

### Characteristics:

- Produces excellent yields of amplifiable cDNA.
- Can synthesize long cDNA targets, up to 12 kb, from small amounts of input RNA.
- Able to reverse transcribe through RNA secondary structures.
- Strict enzyme purity specifications.

### Applications:

- Synthesis of first-strand cDNA
- Array labeling
- cDNA library construction
- 3' and 5' RACE, RT-PCR
- Primer extension

### Quality control:

- Purity: >99% on SDS-PAGE
- Endonuclease- and exonuclease-free
- RNase-free
- Inhibitor-free
- Satisfactory cDNA product yield and length

### Protocol:

1. Prepare the PCR mixture on ice with the following **components included in this kit**:

|                                          |             |
|------------------------------------------|-------------|
| 5× First-Strand Buffer                   | 4 µl        |
| FastGene® Scriptase III (200 units/µl)   | 1 µl        |
| dNTP Mixture (10 mM each)                | 1 µl        |
| 0.1 M DTT                                | 2 µl        |
| Template RNA* ( <b>not included</b> )    | x µl        |
| Primer**                                 | 1 µl        |
| RNase Inhibitor (40 units/µl) - optional | 0.5 µl      |
| Sterile water (RNase free)               | up to 20 µl |

#### \* Concentration of template RNA

Total RNA: 1 pg-5 µg  
Messenger RNA (mRNA): 1 pg-500 ng  
Specific RNA: 0.01 pg-0.5 µg

#### \*\* Prepare one of the following primers

Oligo (dT)<sub>18</sub>: 50-100 µM  
Random hexamer: 50-100 µM  
Specific primer: 15-20 pmol

*An additional annealing step is recommended if using random hexamer. Incubate at 25° for 10 min.*

2. Incubate at 50° for 30-60 min.

3. Incubate at 70° for 15 min to inactivate the reaction.

### Note:

FastGene® Scriptase III excels at 50°C but, can be adjusted within the range of 42°-55°C based on experimental needs. RNAs containing secondary structure and other challenging targets can be synthesized at 55°C without compromising performance. The incubation time for the synthesis can be decreased to a minimum of 30 min.

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