

**Product Information**  
**Reverse Transcription Kit II**

**ExcelRT™ series**

**RP1400     100 RXN**

|                               |         |
|-------------------------------|---------|
| RTase/RI Enzyme Mix           | 100 µl  |
| 5X RT Buffer (DTT/dNTPs)      | 500 µl  |
| Oligo (dT)/Random Primer Mix  | 100 µl  |
| DEPC-Treated H <sub>2</sub> O | 1 ml x2 |

**Storage**

-20°C for 24 months

**Description**

ExcelRT™ Reverse Transcription Kit II is a complete, efficient and convenient kit to synthesize high quality first strand cDNA. This kit contains ExcelRT™ Reverse Transcriptase, which is able to synthesize the first strand cDNA at 37~50°C. The ExcelRT™ Reverse Transcriptase is a recombinant Moloney Murine Leukemia Virus (M-MLV) reverse transcriptase, which is designed to reduce RNase H activity and create better thermal stability. This kit also contains RNAok™ RNase Inhibitor, which is active against RNase A, RNase B, and RNase C. This product is supplied with optimized RT Buffer and Oligo (dT)/Random Primer Mix for highly efficient synthesis of short chain cDNA suitable for real-time PCR.

**Features**

- Contains all components for reverse transcription
- High yield
- Thermostable, up to 50°C
- Reduced RNase H ribonuclease activity
- Suitable for real-time PCR

## Application

- Generation of first strand cDNA from total RNA or mRNA.
- Suitable for generating cDNA from RNA with strong secondary structure which can be reduced at temperature up to 50°C.

## RTase/RI Enzyme Mix

ExcelRT™ Reverse Transcriptase (200 U/μl), RNAok™ RNase Inhibitor (20 U/μl), 20 mM Tris-HCl (pH 7.5), 200 mM NaCl, 0.1 mM EDTA, 1 mM DTT, stabilizer, and 50% (v/v) glycerol

## 5X RT Buffer (DTT/dNTPs)

250 mM Tris-HCl (pH 8.3 at 25°C), 375 mM KCl, 15 mM MgCl<sub>2</sub>, 50 mM DTT and 2.5 mM dNTPs (each)

## Oligo (dT)/Random Primer Mix

50 μM oligo (dT)<sub>20</sub> and 100 μM random hexamers

## First Strand cDNA Synthesis Condition

### 1. Denature (Mixture A):

|                               |                     |
|-------------------------------|---------------------|
| Total RNA                     | X μl (1 ng~2 μg)    |
| Oligo (dT)/Random Primer Mix  | 1 μl                |
| DEPC-Treated H <sub>2</sub> O | to 10 μl final vol. |

Mix well; incubate at 70°C/5 minutes

Place on ice for at least 1 minute

### 2. First strand cDNA buffer (Mixture B) per reaction:

(Master Mix can be prepared before or during the denaturing step)

|                               |      |
|-------------------------------|------|
| 5X RT Buffer (DTT/dNTPs)      | 4 μl |
| DEPC-Treated H <sub>2</sub> O | 5 μl |
| RTase/RI Enzyme Mix           | 1 μl |

Final volume 10 μl

### 3. First strand cDNA synthesis:

|                                      |       |
|--------------------------------------|-------|
| Mixture A (RNA + Primers)            | 10 μl |
| Mixture B (First strand cDNA buffer) | 10 μl |

Final volume 20 μl

Incubate 25°C/10 minutes  
37~50°C/50 minutes

### 4. Termination: 85°C/5 minutes

Store cDNA at -20°C or for immediate qPCR reaction

## Recommended real-time PCR Condition

(SMOBIO's TQ1110 ExcelTaq™ 2X Q-PCR Master Mix (SYBR, ROX))

### Recommended reaction mixture set up for real-time PCR

|                     |                         |
|---------------------|-------------------------|
| cDNA                | 2 µl <sup>*</sup>       |
| Forward primer      | 50~400 nM <sup>**</sup> |
| Reverse primer      | 50~400 nM <sup>**</sup> |
| 2X Q-PCR Master Mix | 10 µl                   |
| H <sub>2</sub> O    | to 20 µl                |
| Total volume        | 20 µl                   |

## Recommended real-time PCR Program

### Two-step cycle

| Steps                                   | Temp.                      | Time                | Cycles |
|---|----------------------------|---------------------|--------|
| Template denature and enzyme activation | 95°C                       | 10 min <sup>#</sup> | 1      |
| Denature                                | 95°C                       | 15 sec              | 40     |
| Annealing/Extension                     | 60°C                       | 60 sec              |        |
| Melting curve analysis                  | Refer to instrument manual |                     |        |

<sup>\*</sup> The recommended amount of cDNA is 100 fg -100 ng. The volume of cDNA should be less than 10% of the total qPCR reaction volume.

<sup>\*\*</sup> The PCR primer concentration for an optimal qPCR reaction may vary according to the primers' properties and template conditions.

<sup>#</sup>10 minutes is suggested in the first step to thoroughly denature DNA and activate enzymes.

## Recommended PCR Condition

(SMOBIO's TP1000 ExcelTaq™ Taq DNA polymerase)

|                    |                      |
|--------------------|----------------------|
| cDNA               | 2~10 µl              |
| Forward primer     | 0.1 – 0.5 µM         |
| Reverse primer     | 0.1 – 0.5 µM         |
| 10X Taq Buffer     | 5 µl                 |
| dNTPs              | 0.2 mM each          |
| Taq DNA polymerase | 0.25 µl (1.25 units) |
| H <sub>2</sub> O   | to 50 µl             |
| Total volume       | 50 µl                |

## Recommended PCR Program

|                       |           |                  |
|-----------------------|-----------|------------------|
| 94°C                  | 2 min     | } 25 ~ 40 cycles |
| 94°C                  | 30 sec    |                  |
| 50~68°C <sup>**</sup> | 30 sec    |                  |
| 72°C                  | 30 sec/kb |                  |
| 72°C                  | 1 min     |                  |

<sup>\*\*</sup>Optimal PCR conditions vary according to primers' thermodynamic properties.

## Other Information

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Caution: Not intended for human or animal diagnostic or therapeutic uses.

## Related Products

|        |  |
|--------|--|
| TQ1100 | ExcelTaq 2X Q-PCR Master Mix (SYBR, no ROX), 200 RXN |
| TQ1110 | ExcelTaq 2X Q-PCR Master Mix (SYBR, ROX), 200 RXN    |
| TQ2110 | ExcelTaq 2X Q-PCR Master Mix (TaqMan, ROX), 200 RXN  |
| TF1000 | SMO-HiFi DNA Polymerase, 100 U                       |
| TF3000 | G-HiFi DNA Polymerase, 100 U                         |
| TP1000 | ExcelTaq Taq DNA Polymerase, 500 U × 1               |
| TP1200 | ExcelTaq 5X PCR Master Dye Mix, 200 RXN              |
| TP5000 | ExcelTaq Hot Start II DNA Polymerase, 500 U          |
| RI1000 | RNAok RNase Inhibitor, 2000 U                        |
| RP1000 | ExcelRT Reverse Transcriptase, 20,000 U              |
| RP1100 | ExcelRT One-step RT-PCR Kit, 50 RXN                  |
| RP1300 | ExcelRT Reverse Transcription Kit, 100 RXN           |
| CK1000 | Champion E. coli Transformation Kit                  |
| DM2100 | ExcelBand 100 bp DNA Ladder, 500 µl                  |
| DM3100 | ExcelBand 1 KB (0.25-10 kb) DNA Ladder, 500 µl       |
| NS1000 | FluoroVue Nucleic Acid Gel Stain (10,000X), 500 µl   |

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