



SMOBIO[®]

Small Bio, Smart Tool

www.smobio.com

Product Information

EzRNA™ series

T7 High Yield RNA Synthesis Kit (Ψ -UTP)

IT1100 50 RXN

T7 RNA Polymerase Mix	100 μ l
10X T7 Reaction Buffer	100 μ l
NTP (Ψ) Premix (25 mM each)	400 μ l
Control DNA	10 μ l
Lithium Chloride (7.5M)	1 ml
Nuclease-Free Water	1 ml

Storage

-20°C for 12 months

Description

The EzRNA™ T7 High Yield RNA Synthesis Kit (Ψ -UTP) is a user-friendly product for enzymatic RNA production. The enzyme mix contains adequate amount of T7 RNA polymerase, pyrophosphatase, and RNase inhibitors for *in vitro* transcription (IVT). Along with 10X Transcription Buffer and NTP (Ψ) Premix, users can swiftly assemble IVT reactions without compromising RNA yield. The EzRNA™ T7 High Yield RNA Synthesis Kit (Ψ -UTP) allows for the attainment of approximately up to 150 μ g RNA yield within 2 hours at 37°C.

Features

- High yield
- Versatile- suitable for short and long transcripts
- NTP premixed- Minimal pipetting and setup time
- Compatible with CleanCap® Reagent AG
- Lithium chloride included for RNA purification

Application

- Generation of RNA from T7 promoter-driven DNA sequences.
- Suitable for subsequent cap-0 and cap-1 modification.

RNA Synthesis Condition

1. Recommended IVT reaction set-up:

Linearized DNA	1 μ g
10X T7 Reaction Buffer	2 μ l
NTP (Ψ) Premix (25 mM each)	8 μ l
T7 RNA Polymerase Mix	2 μ l
Nuclease-Free Water	to 20 μ l
<hr/>	
Total volume	20 μ l

<Note>

- Except T7 RNA polymerase mix, thaw all of other reaction components fully and assemble reaction at room temperature.
- Reaction can be scaled-up proportionally.
- To avoid nuclease contamination, ensure assemble reaction in a clean environment.

2. Recommended IVT condition:

Incubate reaction at 37°C for 2 hours. To achieve maximum yield, reactions could be optimized and extend reaction time to 4~16 hours based on amount of DNA template and/or the size of IVT transcript.

RNA product purification

There are several options of RNA purification listed below:

1. Lithium chloride (LiCl) purification

Lithium chloride could specifically precipitate RNA molecules without additional ethanol or isopropanol. Along with free nucleotides and proteins in IVT reaction, DNA template would be retained in supernatant when using LiCl. It's important to note that this method requires a certain amount of RNA ($>0.1 \mu\text{g}/\mu\text{l}$) and a minimum length of RNA (>300 bases typically) to be effective.

Brief procedure:

Add 1/2 volume of LiCl into RNA sample and mix thoroughly \rightarrow -20°C incubation for at least 1 hour \rightarrow 4°C , $14000\times g$ centrifugation for at least 15 minutes \rightarrow remove supernatant and ice-cold 70% ethanol wash at least twice \rightarrow air-dry pellet shortly and dissolve RNA pellet with water or buffer for certain time period (depend on pellet size).

2. Phenol-chloroform extraction and ethanol precipitation

By virtue of phase separation, protein and free nucleotides would be brought into organic layer completely. Nucleic acid (RNA and DNA) in water phase could be further purified by ethanol precipitation. Acidic phenol could differentially retain RNA molecules but not DNA in water phase.

3. Spin column purification

Certain buffer compositions enable the specific purification of RNA using silica-based columns. Moreover, oligo-dT column are able to purify poly(A)-containing RNA e.g. mRNA. It is important to adhere to the instructions provided in the column purification kit for proper usage.

RNA product assessment

- **Gel electrophoresis**

Compared to DNA, RNA (especially long RNA) molecules may have numerous of secondary structures. For rigorous evaluation, it is essential to generate denatured condition for RNA gel electrophoresis. This typically involves using denatured RNA samples (RNA samples with denatured loading dye) in either denatured agarose gel or polyacrylamide gel. Due to the properties of different gel mobility, it is necessary to use an RNA size ladder rather than the regular DNA ladder for accurate RNA size determination.

- **RNA quantification**

Standard approach is UV absorbance at 260 nm (A_{260}). Concentration of RNA can be calculated as $A_{260} \times 40 \mu\text{g/ml}$. Note that the presence of any impurity in the samples (such as protein, free nucleotides, or DNA) would affect the absorbance reading, leading to an overestimation of RNA quantity.

Other Information

SMOBIO Technology, Inc. claims all warranties with respect to this document, expressed or implied, including but not limited to those of merchantability or fitness for a particular purpose. In no event shall SMOBIO Technology, Inc. be liable, whether in contract, tort, warranty, or under any statute or any other basis for special, incidental, indirect, punitive, multiple or consequential damages in connection with or arising from this document, including but not limited to the use thereof.

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

Related Products

RC1000	EzRNA RNA Capping System, 50 RXN
IT1000	EzRNA T7 High Yield RNA Synthesis Kit , 50 RXN
IT1200	EzRNA T7 High Yield RNA Synthesis Kit (me ¹ Ψ-UTP), 50 RXN
RI1000	RNAok RNase Inhibitor, 2000 U
RP1400	ExcelRT Reverse Transcription Kit II, 100 RXN
TF1000	SMO-HiFi DNA Polymerase, 100 U
TF3000	G-HiFi DNA Polymerase, 100 U
TP5000	ExcelTaq Hot Start II DNA Polymerase, 500 U
TQ1110	ExcelTaq 2X Q-PCR Master Mix (SYBR, ROX), 200 RXN
TQ2110	ExcelTaq 2X Q-PCR Master Mix (TaqMan, ROX), 200 RXN
DM2100	ExcelBand 100 bp DNA Ladder, 500 μl
NS1000	FluoroVue Nucleic Acid Gel Stain (10,000X), 500 μl

The latest version of the manual can be downloaded from www.smobio.com/shop.

For research use only
2024 ver. 1.0.0