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Not for use in diagnostic procedures.



# Biotin-16-UTP

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Biotin- $\epsilon$ -aminocaproyl- $\gamma$ -aminobutyryl-[5-(3-aminoallyl)]-uridine-5'-triphosphate] tetralithium salt

**Cat. No. 11 388 908 910**    250 nmol  
25  $\mu$ l, 10 mM

**Store product at  $-15$  to  $-25^{\circ}\text{C}$ .**

<b>1.</b>	<b>General Information .....</b>	<b>3</b>
1.1.	Contents .....	3
1.2.	Storage and Stability .....	3
	Storage Conditions (Product) .....	3
1.3.	Additional Equipment and Reagent required .....	3
1.4.	Application .....	3
<b>2.</b>	<b>How to Use this Product .....</b>	<b>4</b>
2.1.	Before you Begin .....	4
	Sample Materials .....	4
	Working Solution .....	4
	Working solutions for RNA labeling reaction with Biotin-16-UTP .....	4
2.2.	Protocols .....	4
	RNA labeling by <i>in vitro</i> transcription .....	4
	Analysis of labeled RNA .....	5
	Labeling efficiency .....	5
2.3.	Parameters .....	5
	Chemical Formula .....	5
	Chemical Name .....	5
	Structural formula .....	5
	Molecular Weight .....	5
<b>3.</b>	<b>Supplementary Information .....</b>	<b>6</b>
3.1.	Conventions .....	6
3.2.	Changes to previous version .....	6
3.3.	Ordering Information .....	6
3.4.	Trademarks .....	7
3.5.	License Disclaimer .....	7
3.6.	Regulatory Disclaimer .....	7
3.7.	Safety Data Sheet .....	7
3.8.	Contact and Support .....	7

# 1. General Information

## 1.1. Contents

Vial / Bottle	Label	Function / Description	Content
1	Biotin-16-UTP	10 mM tetralithium salt solution.	1 vial, 25 µl

## 1.2. Storage and Stability

### Storage Conditions (Product)

The product is shipped on dry ice.

When stored at  $-15$  to  $-25^{\circ}\text{C}$ , the product is stable through the expiration date printed on the label.

Vial / Bottle	Label	Storage
1	Biotin-16-UTP	Store at $-15$ to $-25^{\circ}\text{C}$ . <b>⚠ A decomposition of approximately 5% may occur within 6 months.</b>

## 1.3. Additional Equipment and Reagent required

### RNA labeling with Biotin-16-UTP

- Biotin/NTP\* mixture, 10x conc.
  - ⓘ *Also available as a Ribonucleoside Triphosphate Set\*.*
- Transcription Buffer, 10x conc., supplied with RNA polymerases\*: 0.4 M Tris-HCl, pH 8.0, 60 mM MgCl<sub>2</sub>, 100 mM Dithiothreitol (DTT), 20 mM spermidine, 100 mM NaCl
- T7, SP6, or T3 RNA Polymerase\*, see label for lot-specific value
- Protector RNase Inhibitor\*, 40 U/µl in Transcription Buffer with glycerol, 50% (v/v)
- Water, PCR Grade\* or water treated with 0.1% diethylpyrocarbonate (v/v)
- EDTA, 0.2 M, pH 8.0
- Water bath

### Analysis of biotin-labeled RNA

- Nylon Membranes\*
- Streptavidin-AP-conjugate for nucleic acid detection\*.

## 1.4. Application

Biotin-16-UTP can be used in the following applications:

- Substrate for T7, SP6, and T3 RNA Polymerases\*; replaces UTP in *in vitro* transcription for RNA labeling.
- Linearized template DNA with T7, SP6, or T3 promoter is *in vitro* transcribed with the corresponding RNA polymerases using ATP, GTP, CTP, UTP, and Biotin-16-UTP, respectively.

Labeled RNA can be subsequently detected with the:

- Streptavidin-AP-conjugate for nucleic acid detection\*.

## 2. How to Use this Product

### 2.1. Before you Begin

#### Sample Materials

Biotin-16-UTP is used with linearized DNA containing an T7, SP6, or T3 promoter.

**i** The amount of synthesized labeled RNA depends on the amount, size (site of linearization), and purity of the template DNA.

**⚠** **Avoid RNase contamination:** After restriction digest, purify the linearized DNA with the High Pure PCR Product Purification Kit\* or via phenol/chloroform extraction, and subsequent ethanol precipitation.

#### Working Solution

##### Working solutions for RNA labeling reaction with Biotin-16-UTP

Reagent/Buffer	Composition/Concentration
Biotin/NTP* mixture, 10x conc.	10 mM ATP
	10 mM GTP
	10 mM CTP
	6.5 mM UTP
	3.5 mM Biotin-16-UTP
	in Tris-neutralized solution, pH 7.5

### 2.2. Protocols

#### RNA labeling by *in vitro* transcription

The following protocol describes a labeling reaction using Biotin-16-UTP and SP6, T7, or T3 RNA Polymerases.

**1** Add the following to a microcentrifuge tube on ice:

Reagent	Volume [μl]
Linearized template DNA, 1 μg	X
Biotin/NTP mixture, 10x conc.	2
Transcription Buffer, 10x conc.	2
Water, PCR Grade	X
Protector RNase Inhibitor	1
T7, SP6, or T3 RNA Polymerase, 40 U	X
<b>Final Volume</b>	<b>20</b>

- Mix and centrifuge briefly.
- Incubate for 2 hours at +37°C.

**i** *Optional: Remove template DNA by DNase, RNase free\*-treatment (20 U, 15 minutes, +37°C).*

**2** Stop the reaction by adding 2 μl 0.2 M EDTA (pH 8.0) and/or heating to +65°C.

**3** Use the labeled probe immediately or store at -15 to -25°C in aliquots.

## Analysis of labeled RNA

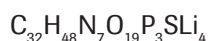
Analyze the transcript by agarose gel electrophoresis and ethidium bromide staining. Estimate the yield of biotin-labeled RNA via spot assay in combination with Nylon Membranes\* and detection with the Streptavidin-AP conjugate for nucleic acid detection\* prior to usage as a hybridization probe.

## Labeling efficiency

Depending on the length, purity, and sequence of the template DNA, approximately 10 µg of biotin-labeled RNA are synthesized under standard conditions. An average of 10 µg RNA per 1 µg template DNA is obtained.

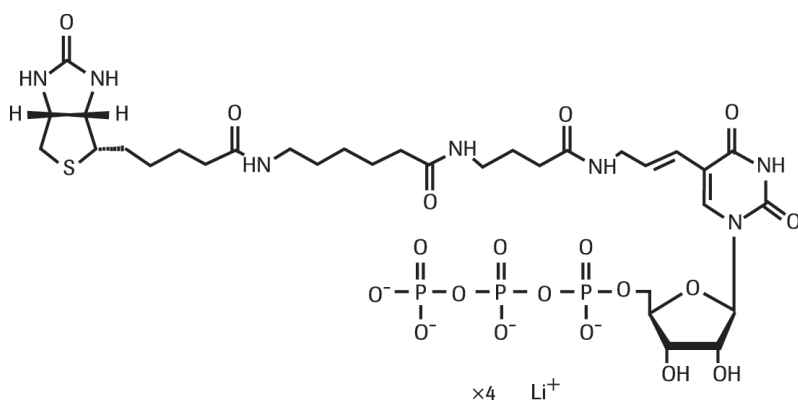
## 2.3. Parameters

### Chemical Formula



### Chemical Name

### Structural formula



**Fig. 1:** Chemical structure of Biotin-16-UTP









### Molecular Weight

987.5 Da

## 3. Supplementary Information

### 3.1. Conventions

To make information consistent and easier to read, the following text conventions and symbols are used in this document to highlight important information:

Text convention and symbols	
 Information Note: Additional information about the current topic or procedure.	
 Important Note: Information critical to the success of the current procedure or use of the product.	
   etc.	Stages in a process that usually occur in the order listed.
   etc.	Steps in a procedure that must be performed in the order listed.
* (Asterisk)	The Asterisk denotes a product available from Roche Diagnostics.

### 3.2. Changes to previous version

Layout changes.

Editorial changes.

### 3.3. Ordering Information

Product	Pack Size	Cat. No.
Reagents, kits		
GTP	400 µl, 40 µmol, 100 mM	11 140 957 001
Nylon Membranes, positively charged	10 sheets, 20 x 30 cm	11 209 272 001
	20 sheets, 10 x 15 cm	11 209 299 001
	1 roll, 0.3 x 3 m	11 417 240 001
	DNase I recombinant, RNase-free	10,000 U, 10 U/µl
CTP	400 µl, 40 µmol, 100 mM	11 140 922 001
Streptavidin-AP-conjugate for nucleic acid detection	150 U, (200 µl)	11 093 266 910
SP6 RNA Polymerase	1,000 U, > 20 U/µl	10 810 274 001
	5,000 U, > 20 U/µl	11 487 671 001
Water, PCR Grade	25 ml, 25 x 1 ml	03 315 932 001
	25 ml, 1 x 25 ml	03 315 959 001
	100 ml, 4 x 25 ml	03 315 843 001
Protector RNase Inhibitor	2,000 U, 40 U/µl	03 335 399 001
	10,000 U, 5 x 2,000 U	03 335 402 001
T7 RNA Polymerase	1,000 U, ≥ 20 U/µl	10 881 767 001
	5,000 U, ≥ 20 U/µl	10 881 775 001
ATP	400 µl, 100 mM 40 µmol	11 140 965 001
T3 RNA Polymerase	1,000 U, ≥ 20 U/µl	11 031 163 001
	5,000 U, ≥ 20 U/µl	11 031 171 001
Ribonucleoside Triphosphate Set	4 x 200 µl, 4 x 20 µmol, 100 mM each	11 277 057 001
UTP	400 µl, 40 µmol, 100 mM	11 140 949 001
High Pure PCR Product Purification Kit	1 kit, up to 50 purifications	11 732 668 001
	1 kit, up to 250 purifications	11 732 676 001

### 3.4. Trademarks

All product names and trademarks are the property of their respective owners.

### 3.5. License Disclaimer

For patent license limitations for individual products please refer to:

**List of biochemical reagent products.**

### 3.6. Regulatory Disclaimer

For life science research only. Not for use in diagnostic procedures.

### 3.7. Safety Data Sheet

Please follow the instructions in the Safety Data Sheet (SDS).

### 3.8. Contact and Support

To ask questions, solve problems, suggest enhancements or report new applications, please visit our **Online Technical Support Site.**

To call, write, fax, or email us, visit **sigma-aldrich.com**, and select your home country. Country-specific contact information will be displayed.

