

# 100 mM dNTP Set

Cat. No. ST040-040M Size: 4 x 10 ml Cat. No. ST040-4000 Size: 4 x 1 ml Cat. No. ST046-1000 Size: 4 x 250 µl Cat. No. ST046-0100 Size: 4 x 25 µl

#### Description

100 mM dNTP (2'-deoxynucleoside 5'-triphosphate) Set consists of all four deoxynucleotides (dATP, dCTP, dGTP, dTTP), each at a concentration of 100 mM. The deoxynucleotides are suitable for use in polymerase chain reaction (PCR), sequencing, fill-in, nick translation, cDNA synthesis, and TdT tailing reactions. The product is supplied as ready-to-use solutions.

#### **Features**

- > Compatible with almost DNA polymerases in a variety of applications.
- > ≥ 99% pure as determined by HPLC analysis.
- > Exceptional stability.

## **Applications**

> PCR amplification

## **Kit Contents**

Contents	ST040-040M	ST040-4000	ST046-0100	ST046-1000
dATP	10 ml	1 ml	25 µl	250 µl
dTTP	10 ml	1 ml	25 µl	250 µl
dCTP	10 ml	1 ml	25 µl	250 µl
dGTP	10 ml	1 ml	25 µl	250 µl

#### **Quality Control**

The quality of the 100 mM dNTP Set is tested on a lot-to-lot basis to ensure consistent product quality.

#### **Required Materials**

> PCR equipments Primer PCR tube ➤ PCR grade water

## **Buffer Preparation**

TE Buffer (Tris-EDTA, pH8.0): 10 mM Tris-HCl, pH 8.0 with 0.1mM EDTA

#### Protocol

Add recommended volume of dNTP solution into PCR reaction.

The volume

of dNTP mixture

 $0.16 \, \mu l$ 

 $0.4 \mu l$ 

0.8 µl

1.2 µl

The following in the below table is recommended:

20 µl Final Reaction Volume

Final dNTP

Concentration

0.2 mM

0.5 mM

1.0 mM

1.5 mM

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Final dNTP Concentration	The volume of dNTP mixture
0.2 mM	0.2 µl
0.5 mM	0.5 µl
1.0 mM	1 µl
1.5 mM	1.5 µl

## 25 ul Final Reaction Volume

## 50 µl Final Reaction Volume

Final dNTP Concentration	The volume of dNTP mixture
0.2 mM	0.4 µl
0.5 mM	1 µl
1.0 mM	2 µl
1.5 mM	3 µl

## **Troubleshooting**

Problem	Cause	Solution
Incorrect amplification or PCR inhibition.	Incorrect dNTP concentration.	Check and optimized the dNTP concentration of the PCR reaction.
No amplicon	Error in set up.	Repeat the experiment, checking all reagents are added in correct volumes.  Use master mix to ensure all components added correctly.
Non-specific amplification - smeared product.	Template degraded.	Minimize freeze thawing of DNA. Run template on agarose gel to check integrity.
Wrong size band amplified.	Contamination	Check no template control for bands.

# **Related Ordering Information**

Cat. No.	Description	Size
SM101-0500	Taq DNA Polymerase	500 U
SA001-0500	AGAROSE Tablet, 0.5g	100 Tablets

## Caution

- > Check buffers before use for precipitation.
- > Aliquot reagents to avoid contamination and repeated freeze-thaw cycles.
- > During operation, always wear a lab coat, disposable gloves, and protective equipment.
- > Research Use Only. Not intended for any animal or human therapeutic or diagnostic uses.