



25 mM dNTP Mix, PCR Grade

Cat No. ST100-1000
Store at -20°C

Size: 1 ml

Description

25 mM dNTP (2'-deoxynucleoside 5'-triphosphate) Mix consists of a solution of all four nucleotides (dATP, dCTP, dGTP, dTTP), each at a concentration of 25 mM. It is neutralized to pH 8.0 with NaOH, and supplied in purified water. 25 mM dNTP Mix is suitable for use in polymerase chain reaction (PCR), sequencing, fill-in, nick translation, cDNA synthesis, and TdT-tailing reactions.

Features

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

Application

- PCR amplification

Kit Contents

| Contents | ST100-1000 |
|----------------|------------|
| 25 mM dNTP Mix | 1 ml |

Quality Control

The quality of the 25 mM dNTP Mix, PCR Grade is tested on a lot-to-lot basis to ensure consistent product quality.

Required Materials

- PCR equipments
- PCR tube
- Primer
- PCR grade water

Protocol

Add recommended volume of dNTP solution into PCR reaction.

The following in the below table is recommended:

20 µl Final Reaction Volume

| Final dNTP Concentration | dNTP Volume |
|--------------------------|-------------|
| 0.2 mM | 0.16 µl |
| 0.5 mM | 0.4 µl |
| 1.0 mM | 0.8 µl |
| 1.5 mM | 1.2 µl |

25 µl Final Reaction Volume

| Final dNTP Concentration | dNTP Volume |
|--------------------------|-------------|
| 0.2 mM | 0.2 µl |
| 0.5 mM | 0.5 µl |
| 1.0 mM | 1 µl |
| 1.5 mM | 1.5 µl |

50 µl Final Reaction Volume

| Final dNTP Concentration | dNTP Volume |
|--------------------------|-------------|
| 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 |

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.