

# dNTP Mix

Premix of 25 mM dATP, dCTP, dGTP and dTTP

|  | Cat. No.      | Volume  | Amount       |
|--|---------------|---------|--------------|
|  | DMIX25_100ML  | 100 ml  | 4 x 2.5 mmol |
|  | DMIX25_1000ML | 1000 ml | 4 x 25 mmol  |

For *in vitro* use only

Quality guaranteed for 12 months

Store at -20°C, short term (up to one week) exposure to ambient temperature possible

## dATP

2'-Deoxyadenosine 5'-triphosphate, sodium salt

Molecular formula: C<sub>10</sub>H<sub>13</sub>N<sub>5</sub>O<sub>12</sub>P<sub>3</sub> (Anion)

Molecular weight: 488.16 (Anion)

## dCTP

2'-Deoxycytidine 5'-triphosphate, sodium salt

Molecular formula: C<sub>9</sub>H<sub>13</sub>N<sub>3</sub>O<sub>13</sub>P<sub>3</sub> (Anion)

Molecular weight: 464.13 (Anion)

## dGTP

2'-Deoxyguanosine 5'-triphosphate, sodium salt

Molecular formula: C<sub>10</sub>H<sub>13</sub>N<sub>5</sub>O<sub>13</sub>P<sub>3</sub> (Anion)

Molecular weight: 504.16 (Anion)

## dTTP

2'-Deoxythymidine 5'-triphosphate, sodium salt

Molecular formula: C<sub>10</sub>H<sub>14</sub>N<sub>2</sub>O<sub>14</sub>P<sub>3</sub> (Anion)

Molecular weight: 479.14 (Anion)

## Form

clear aqueous solution

## pH

8.5 ± 0.2 (22 °C)

## Purity

≥ 99 % (HPLC)

## Application for PCR

For standard PCR applications a final concentration of 200 μM each dNTP is recommended.

## Description

dNTP Mix is an equimolar mixture of ultrapure dATP, dCTP, dGTP, and dTTP supplied as clear aqueous solution (pH 8.5) and suitable for all molecular biology applications including PCR/qPCR, reverse transcription, DNA labeling and DNA sequencing

## Quality Control Specifications

Low Copy Long Range PCR (18 kb, lambda DNA, template dilution series): PCR fragment with 50 pg of template or less

RT-PCR (749 bp fragment, human GAPDH gene, template dilution series): PCR fragment with 10 pg of template or less

Contamination with bacterial or human DNA: not detectable

DNases, RNases, Nicking Activity: not detectable

Proteases: not detectable

## Selected References

[1] Erlich *et al.* (1988) Primer-directed enzymatic amplification of DNA with a thermostable DNA polymerase. *Science* **29** (239):487.

[2] Gelfand *et al.* (1991) Detection of specific polymerase chain reaction product by utilizing the 5'-3' exonuclease activity of *Thermus aquaticus* DNA polymerase. *Proc. Natl. Acad. Sci. USA* **88** (16):7276.

[3] Sanger *et al.* (1977) DNA sequencing with chain-terminating inhibitors. *Proc. Natl. Acad. Sci. USA* **74**:5463.