



Data Sheet (26.11.2020)

dNTP Set

Package sizes:

| Cat.-No. | Amount |
|-----------|----------------------------|
| mi-N1005S | 4 x 200 µl (4 x 20 µmol) |
| mi-N1005L | 4 x 1000 µl (4 x 100 µmol) |

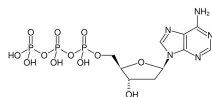
Storage conditions: - 20 ± 5°C

Short term exposure (up to 1 week cumulative) to ambient temperature possible.

Shelf Life: 12 months after date of delivery

For research use only! Only for in vitro use!

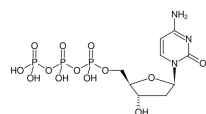
dATP Structure:



Molecular Formula: C₁₀H₁₆N₅O₁₂P₃ (free acid)

Molecular Weight: 491.18 g/mol (free acid)

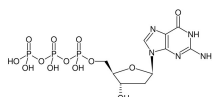
dCTP Structure:



Molecular Formula: C₉H₁₆N₃O₁₃P₃ (free acid)

Molecular Weight: 467.15 g/mol (free acid)

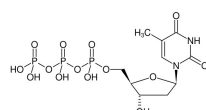
dGTP Structure:



Molecular Formula: C₁₀H₁₆N₅O₁₃P₃ (free acid)

Molecular Weight: 507.18 g/mol (free acid)

dTTP Structure:



Molecular Formula: C₁₀H₁₇N₂O₁₄P₃ (free acid)

Molecular Weight: 482.17 g/mol (free acid)

Purity: ≥ 99 % (HPLC)

Form: clear aqueous solution

Concentration: 100 mM - 110 mM

pH: 8.5 ± 0.2 (22 °C)

Spectroscopic Properties:

dATP: λ_{max} 259 nm, ε 15.4 L mmol⁻¹ cm⁻¹ (Tris-HCl pH 7.0)

dCTP: λ_{max} 271 nm, ε 8.9 L mmol⁻¹ cm⁻¹ (Tris-HCl pH 7.0)

dGTP: λ_{max} 252 nm, ε 13.7 L mmol⁻¹ cm⁻¹ (Tris-HCl pH 7.0)

dTTP: λ_{max} 262 nm, ε 9.6 L mmol⁻¹ cm⁻¹ (Tris-HCl pH 7.0)



Description:

dNTP Set contains four separate solutions of ultrapure dATP, dCTP, dGTP and dTTP supplied as clear aqueous solutions (pH 8.5).

| dNTP | Cat. No. | cap color |
|-------------|-----------------|------------------|
| dATP | mi-N1001 | red |
| dCTP | mi-N1002 | blue |
| dGTP | mi-N1003 | yellow |
| dTTP | mi-N1004 | green |

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:

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

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

Holland *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.