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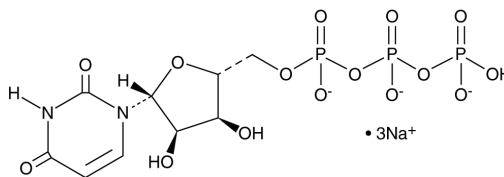
PRODUCT INFORMATION



Uridine 5'-triphosphate (sodium salt)

Item No. 9003530

CAS Registry No.: 19817-92-6
Formal Name: uridine 5'-(tetrahydrogen triphosphate), trisodium salt
Synonyms: NSC 20260, UTP
MF: C₉H₁₂N₂O₁₅P₃ • 3Na
FW: 550.1
Purity: ≥95%
UV/Vis.: λ_{max}: 262 nm
Supplied as: A crystalline solid
Storage: -20°C
Stability: ≥2 years



Information represents the product specifications. Batch specific analytical results are provided on each certificate of analysis.

Laboratory Procedures

Uridine 5'-triphosphate (UTP) (sodium salt) is supplied as a crystalline solid. Aqueous solutions of UTP (sodium salt) can be prepared by directly dissolving the crystalline solid in aqueous buffers. The solubility of UTP (sodium salt) in PBS, pH 7.2, is approximately 10 mg/ml. We do not recommend storing the aqueous solution for more than one day.

Description

UTP is a nucleotide and dual agonist of purinergic P2Y₂ and P2Y₄ receptors (EC₅₀s = 55 and 80 nM, respectively, for stimulation of phospholipase C in 1321N1 cells expressing human receptors).^{1,2} It is selective for P2Y₂ and P2Y₄ receptors over P2Y₆ receptors (EC₅₀ = >10,000 nM).² UTP stimulates proliferation of PANC-1 cells (EC₅₀ = 13.1 μM), an effect that can be prevented by siRNA against the P2Y₂ receptor.¹ It induces vasoconstriction in perfused isolated canine epicardial coronary artery in a concentration-dependent manner.³ UTP is formed from uridine monophosphate (UMP) by two sequential phosphorylations and can be converted to cytidine 5'-triphosphate (CTP; Item No. 18147).⁴ It also reacts with glucose-1-phosphate (Item No. 30566) to form UDP-glucose (Item No. 15602), a precursor in the biosynthesis of glycogen.⁵

References

1. Choi, J.H., Ji, Y.G., and Lee, D.H. Uridine triphosphate increases proliferation of human cancerous pancreatic duct epithelial cells by activating P2Y2 receptor. *Pancreas* **42(4)**, 680-686 (2013).
2. Maruoka, H., Jayasekara, M.P.S., Barrett, M.O., et al. Pyrimidine nucleotides with 4-alkyloxymino and terminal tetraphosphate δ-ester modifications as selective agonists of the P2Y₄ receptor. *J. Med. Chem.* **54(12)**, 4018-4033 (2011).
3. Matsumoto, T., Nakane, T., and Chiba, S. UTP induces vascular responses in the isolated and perfused canine epicardial coronary artery via UTP-preferring P2Y receptors. *Br. J. Pharmacol.* **122(8)**, 1625-1632 (1997).
4. Berg, J.M., Tymoczko, J.L., and Stryer, L. In de novo synthesis, the pyrimidine ring is assembled from bicarbonate, aspartate, and glutamine. *Biochemistry*. Gatto, G., editor, 5th edition, W. H. Freeman (2002).
5. Berg, J.M., Tymoczko, J.L., and Stryer, L. Glycogen is synthesized and degraded by different pathways. *Biochemistry*. Gatto, G., editor, 5th edition, W. H. Freeman (2002).

WARNING

THIS PRODUCT IS FOR RESEARCH ONLY - NOT FOR HUMAN OR VETERINARY DIAGNOSTIC OR THERAPEUTIC USE.

SAFETY DATA

This material should be considered hazardous until further information becomes available. Do not ingest, inhale, get in eyes, on skin, or on clothing. Wash thoroughly after handling. Before use, the user must review the complete Safety Data Sheet, which has been sent via email to your institution.

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