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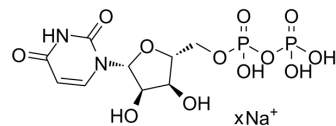
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Uridine 5'-diphosphate sodium salt

Cat. No.:	HY-W010820
CAS No.:	21931-53-3
Molecular Formula:	C ₉ H ₁₄ N ₂ O ₁₂ P ₂ .xNa
Target:	P2Y Receptor; DNA/RNA Synthesis; Endogenous Metabolite
Pathway:	GPCR/G Protein; Cell Cycle/DNA Damage; Metabolic Enzyme/Protease
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.



BIOLOGICAL ACTIVITY

Description	Uridine 5'-diphosphate sodium salt is a potent, selective P2Y ₆ receptor native agonist (EC ₅₀ =300 nM; pEC ₅₀ =6.52) and a potent P2Y ₁₄ antagonist (pEC ₅₀ =7.28). Uridine 5'-diphosphate sodium salt, an endogenous metabolite, catalyzes the glucuronidation of a wide array of substrates and is used in nucleic acid (RNA) biosynthesis ^{[1][2]} .								
IC₅₀ & Target	Human Endogenous Metabolite								
In Vitro	<p>Uridine 5'-diphosphate sodium salt (100 μM; for 15 min) significantly induces microglial CCL2 and CCL3 mRNA expression^[2]. Uridine 5'-diphosphate sodium salt (100 μM; 3 hours) induces chemokine expression in microglia^[2]. Uridine 5'-diphosphate sodium salt (100 μM; 0.5, 1, 3, 6 12 hours) induces expression of mRNA encoding CCL2 and CCL3 within 30 min, and such expression reaches maximal levels at 1 h, returning to basal levels at 3-12 h^[2]. UDP (10, 100, 1000 μM; 3 hours) induces a concentration-dependent increase in the expression of chemokines at both the mRNA and protein level^[2]. Uridine 5'-diphosphate sodium salt (100 μM; for 15 min) induces activation of NFATc1 and NFATc2 in microglia^[2]. MCE has not independently confirmed the accuracy of these methods. They are for reference only.</p> <p>RT-PCR^[2]</p> <table border="1"> <tr> <td>Cell Line:</td> <td>Primary microglia</td> </tr> <tr> <td>Concentration:</td> <td>100 μM</td> </tr> <tr> <td>Incubation Time:</td> <td>For 15 min</td> </tr> <tr> <td>Result:</td> <td>Significantly induced microglial CCL2 and CCL3 mRNA expression.</td> </tr> </table>	Cell Line:	Primary microglia	Concentration:	100 μM	Incubation Time:	For 15 min	Result:	Significantly induced microglial CCL2 and CCL3 mRNA expression.
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REFERENCES

- [1]. Jacobson KA, et al. Development of selective agonists and antagonists of P2Y receptors. *Purinergic Signal*. 2009 Mar;5(1):75-89.
- [2]. Kim B, et al. Uridine 5'-diphosphate induces chemokine expression in microglia and astrocytes through activation of the P2Y6 receptor. *J Immunol*. 2011 Mar 15;186(6):3701-9.

Caution: Product has not been fully validated for medical applications. For research use only.

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