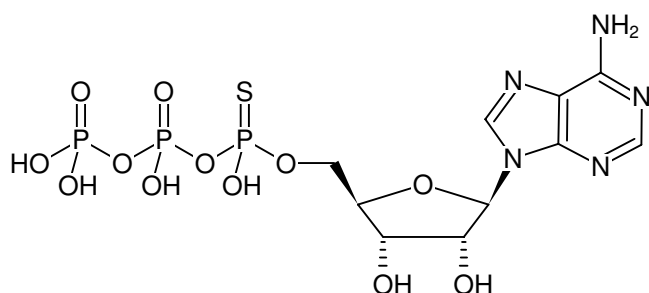


**ATPαS**Adenosine-5'-(α-thio)-triphosphate, Sodium salt; (Mixture of R<sub>p</sub> and S<sub>p</sub> isomers)

Cat. No.	Amount
NU-408S	25 µl (100 mM)
NU-408L	5 x 25 µl (100 mM)



Structural formula of ATPαS

**For general laboratory use.****Shipping:** shipped on gel packs**Storage Conditions:** store at -20 °C

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

**Shelf Life:** 12 months after date of delivery**Molecular Formula:** C<sub>10</sub>H<sub>16</sub>N<sub>5</sub>O<sub>12</sub>P<sub>3</sub>S (free acid)**Molecular Weight:** 523.24 g/mol (free acid)**Exact Mass:** 522.97 g/mol (free acid)**CAS#:** 29220-54-0**Purity:** ≥ 95 % (HPLC)**Form:** solution in water**Color:** colorless to slightly yellow**Concentration:** 100 mM - 110 mM**pH:** 7.5 ± 0.5**Spectroscopic Properties:** λ<sub>max</sub> 259 nm, ε 15.4 L mmol<sup>-1</sup> cm<sup>-1</sup> (Tris-HCl pH 7.5)**Applications:**Crystal structure in complex with adenylyl cyclase<sup>[1]</sup>Activation of beta-cell adenylyl cyclase<sup>[2]</sup>Potent agonist for P2Y<sub>11</sub>-receptor<sup>[3]</sup>Full agonist for rP2Y<sub>1</sub>-receptor<sup>[4]</sup>**Specific Ligands:**Binding to P2Y<sub>2</sub>- and P2Y<sub>6</sub>-receptors<sup>[5]</sup>

Ligand for purinergic receptors:

P2Y<sup>[2,5,7]</sup>P2Y<sub>2</sub><sup>[6]</sup>P2Y<sub>11</sub><sup>[3]</sup>**Related Products:**

γ-[(6-Aminoethyl)-imido]-ATPαS (NU-278) labeled with various fluorescent dyes

**Selected References:**[1] Mou *et al.* (2009) Structural basis for inhibition of mammalian adenylyl cyclase by calcium. *Biochemistry* **48** (15):3387.[2] Chevassus *et al.* (2002) P2Y receptor activation enhances insulin release from pancreatic beta-cells by triggering the cyclic AMP/protein kinase A pathway. *Naunyn Schmiedebergs Arch. Pharmacol.* **366** (5):464.[3] Van der Weyden *et al.* (2000) Pharmacological characterisation of P2Y<sub>11</sub> receptor in stably transfected haematological cell lines. *Mol. Cell. Biochem.* **213** (1):75.[4] Vohringer *et al.* (2000) A chimeric rat brain P2Y<sub>1</sub> receptor tagged with GFP: high-affinity ligand recognition of adenosine diphosphates and triphosphates and selectivity identical to that of wild-type receptor. *Biochem. Pharmacol.* **59** (7):791.[5] Schafer *et al.* (2003) ATP- and UTP-activated P2Y receptors differently regulate proliferation of human lung epithelial tumor cells. *Am. J. Physiol. Lung Cell. Mol. Physiol.* **285** (2):L376.[6] Gorodeski (2002) Regulation of transcervical permeability by two distinct P2 purinergic receptor mechanisms. *Am. J. Physiol. Cell Physiol.* **282** (1):C75.[7] Schäfer *et al.* (1999) ATPαS is a ligand for P2Y receptors in synaptosomal membranes: solubilization of [<sup>35</sup>S]ATPαS binding proteins associated with G-proteins. *Neurochemistry Int.* **34** (4):303.Kim *et al.* (2009) Profiling the selectivity of DNA ligases in an array format with mass spectrometry. *Nucleic Acids Research* **38** (1):e2.Vohringer *et al.* (2001) Pharmacological characterization of the rat brain P2Y<sub>1</sub> receptor expressed in HEK293 cells: Ca<sup>2+</sup> signaling and receptor regulation. *Drug Develop. Res.* **53** (2):172.

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