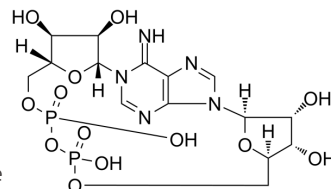


Cyclic ADP-ribose

Cat. No.:	HY-N7395
CAS No.:	119340-53-3
Molecular Formula:	C ₁₅ H ₂₁ N ₅ O ₁₃ P ₂
Molecular Weight:	541.3
Target:	Calcium Channel; TRP Channel; Endogenous Metabolite
Pathway:	Membrane Transporter/Ion Channel; Neuronal Signaling; Metabolic Enzyme/Protease
Storage:	-80°C



SOLVENT & SOLUBILITY

In Vitro

H₂O : 5 mg/mL (9.24 mM; Need ultrasonic and warming)

	Solvent Concentration	Mass	1 mg	5 mg	10 mg
Preparing Stock Solutions	1 mM		1.8474 mL	9.2370 mL	18.4740 mL
	5 mM		0.3695 mL	1.8474 mL	3.6948 mL
	10 mM		---	---	---

Please refer to the solubility information to select the appropriate solvent.

BIOLOGICAL ACTIVITY

Description	Cyclic ADP-ribose (cADPR) is a potent second messenger for calcium mobilization that is synthesized from NAD ⁺ by an ADP-ribosyl cyclase. Cyclic ADP-ribose increases cytosolic calcium mainly by Ryanodine receptor-mediated release from endoplasmic reticulum and also by extracellular influx through the opening of TRPM2 channels ^{[1][2][3]} .
IC ₅₀ & Target	Calcium mobilization ^[1] TRPM2 channels ^[3] Endogenous metabolite ^[1]
In Vitro	cADPR (20 nM) elicits a large rapid Ca ²⁺ release in sea urchin eggs homogenates ^[1] . cADPR (100 μM; 10 min) induces a sustained elevation of intracellular calcium concentration in a subset (64%) of cultured astrocytes ^[4] . cADPR (100 μM) and heat (35-38.5 °C) stimulates oxytocin OT release from the isolated hypothalami of male mice in culture ^[5] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.
In Vivo	cADPR (100 μM; push-pull type of brain microperfusion) elevates OT concentrations in ordinate or subordinate mice ^[5] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.

REFERENCES

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- [4]. Verderio C, et, al. Evidence of a role for cyclic ADP-ribose in calcium signalling and neurotransmitter release in cultured astrocytes. *J Neurochem*. 2001 Aug;78(3):646-57.
- [5]. Zhong J, et, al. Cyclic ADP-Ribose and Heat Regulate Oxytocin Release via CD38 and TRPM2 in the Hypothalamus during Social or Psychological Stress in Mice. *Front Neurosci*. 2016 Jul 22;10:304.
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Caution: Product has not been fully validated for medical applications. For research use only.

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