c-di-AMP

®

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Cat. No.:	HY-12326				
CAS No.:	54447-84-6				
Molecular Formula:	$C_{20}H_{24}N_{10}O_{12}P_{2}$				
Molecular Weight:	658.41				
Target:	STING; Bacterial; Endogenous Metabolite				
Pathway:	Immunology/Inflammation; Anti-infection; Metabolic Enzyme/Protease				
Storage:	Powder	-20°C	3 years		
		4°C	2 years		
	In solvent	-80°C	6 months		
		-20°C	1 month		

SOLVENT & SOLUBILITY

		Mass Solvent	1 mg	5 mg	10 mg
		Concentration			
Preparing Stock Solutions	1 mM	1.5188 mL	7.5941 mL	15.1881 mL	
	5 mM	0.3038 mL	1.5188 mL	3.0376 mL	
		10 mM			

BIOLOGICAL ACTIVITY				
Description	c-di-AMP (Cyclic diadenylate) is a STING agonist, which binds to the transmembrane protein STING thereby activating the TBK3-IRF3 signaling pathway, subsequently triggering the production of type I IFN and TNF. c-di-AMP (Cyclic diadenylate) is also a bacterial second messenger, which regulates cell growth, survival, and virulence, primarily within Gram-positive bacteria, and also regulates host immune response. c-di-AMP (Cyclic diadenylate) acts as a potent mucosal adjuvant stimulating both humoral and cellular responses ^{[1][2][3][4]} .			
IC_{50} & Target	STING ^[3]			
In Vitro	c-di-AMP (Cyclic diadenylate) signaling is a central factor in many Gram-positive bacteria regulating cell wall synthesis, potassium ion channels, DNA repair, and biofilm formation. c-di-AMP is also essential for cell growth, survival, and virulence of several well-known human pathogenic bacteria including S. aureus, L. monocytogenes, S. pyogenes, and Mycobacterium spp ^[1] . c-di-AMP (Cyclic diadenylate) combines with model antigens, such as OVA or β-Gal, acts as a potent mucosal adjuvant stimulating both humoral and cellular responses ^[4] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.			

CUSTOMER VALIDATION

- Gut Microbes. 2022 Jan-Dec;14(1):2119055.
- Cell Death Dis. 2022 Jul 28;13(7):653.

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REFERENCES

[1]. Fahmi T, et al. c-di-AMP: An Essential Molecule in the Signaling Pathways that Regulate the Viability and Virulenceof Gram-Positive Bacteria. Genes (Basel). 2017 Aug 7;8(8).

[2]. Ning H, et al. Recombinant BCG With Bacterial Signaling Molecule Cyclic di-AMP as Endogenous AdjuvantInduces Elevated Immune Responses After Mycobacterium tuberculosis Infection. Front Immunol. 2019 Jul 3;10:1519.

[3]. Ebensen T, et al. The Combination Vaccine Adjuvant System Alum/c-di-AMP Results in Quantitative and QualitativeEnhanced Immune Responses Post Immunization. Front Cell Infect Microbiol. 2019 Feb 19;9:31.

[4]. Sanchez MV, et al. Intranasal delivery of influenza rNP adjuvanted with c-di-AMP induces strong humoral and cellularimmune responses and provides protection against virus challenge. PLoS One. 2014 Aug 20;9(8):e104824.

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

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