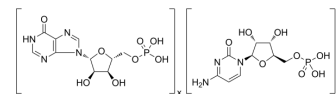


Polyinosinic-polycytidylic acid

Cat. No.:	HY-107202									
CAS No.:	24939-03-5									
Molecular Formula:	$(C_{10}H_{13}N_4O_8P)_x \cdot (C_9H_{14}N_3O_8P)_x$									
Target:	Toll-like Receptor (TLR); PKD; HSP; Bcl-2 Family; Interleukin Related									
Pathway:	Immunology/Inflammation; Apoptosis; Cell Cycle/DNA Damage; Metabolic Enzyme/Protease									
Storage:	<table border="0"> <tr> <td>Powder</td> <td>-20°C</td> <td>3 years</td> </tr> <tr> <td>In solvent</td> <td>-80°C</td> <td>6 months</td> </tr> <tr> <td></td> <td>-20°C</td> <td>1 month</td> </tr> </table>	Powder	-20°C	3 years	In solvent	-80°C	6 months		-20°C	1 month
Powder	-20°C	3 years								
In solvent	-80°C	6 months								
	-20°C	1 month								



SOLVENT & SOLUBILITY

In Vitro	H ₂ O : 50 mg/mL (Need ultrasonic)
In Vivo	1. Add each solvent one by one: PBS Solubility: 100 mg/mL (Infinity mM); Clear solution; Need ultrasonic

BIOLOGICAL ACTIVITY

Description	Polyinosinic-polycytidylic acid (Poly (I:C)), a synthetic analog of double-stranded RNA, is an agonist of toll-like receptor (TLR)-3. Polyinosinic-polycytidylic acid facilitates tumor regression and has a disruptive effect on the airway epithelial barrier. Polyinosinic-polycytidylic acid has protective effects against cerebral ischemia/reperfusion (I/R) injury and can be used as vaccine adjuvant to enhance innate and adaptive immune responses ^{[1][2][3][4][5]} .											
IC₅₀ & Target	Bcl-2	Bax	IL-17A	IL-13								
	HSP70											
In Vitro	<p>Polyinosinic-polycytidylic acid (0.5-5 µg/mL, 3-24 h) induces a dose- and time-dependent increase in paracellular permeability of immortalized airway epithelial cells^[4].</p> <p>Polyinosinic-polycytidylic acid (5 µg/mL, 24 h) does not have cytotoxicity to 16HBE14o- cells^[4].</p> <p>Polyinosinic-polycytidylic acid (5 µg/mL, 6 h) induces disruption of epithelial apical junctional complexes(AJCs) and tight junctions (TJs) in 16HBE14o- cells^[4].</p> <p>MCE has not independently confirmed the accuracy of these methods. They are for reference only.</p> <p>Cell Cytotoxicity Assay^[4]</p> <table border="1" style="width: 100%; border-collapse: collapse;"> <tr> <td style="width: 30%;">Cell Line:</td> <td>16HBE14o- cells</td> </tr> <tr> <td>Concentration:</td> <td>5 µg/mL</td> </tr> <tr> <td>Incubation Time:</td> <td>24 h</td> </tr> <tr> <td>Result:</td> <td>Did not lead to significant accumulation of LDH in cell-culture medium</td> </tr> </table>				Cell Line:	16HBE14o- cells	Concentration:	5 µg/mL	Incubation Time:	24 h	Result:	Did not lead to significant accumulation of LDH in cell-culture medium
Cell Line:	16HBE14o- cells											
Concentration:	5 µg/mL											
Incubation Time:	24 h											
Result:	Did not lead to significant accumulation of LDH in cell-culture medium											

In Vivo

Polyinosinic-polycytidylic acid (2.5-10 mg/mL, Stereotaxic injection, single dose) induces a sustained inflammatory reaction in the substantia nigra (SN) and in the dorsolateral striatum. [2].

Polyinosinic-polycytidylic acid (10 µg/mouse, Intraperitoneal injection, single dose) decreases lung tumor growth in mice^[3].

Polyinosinic-polycytidylic acid (1.25 mg/kg, Intraperitoneal injection, single dose) exerts therapeutic effects against cerebral I/R injury through the downregulation of TLR4/MyD88 signaling via TLR3 in MCAO model mice^[5].

MCE has not independently confirmed the accuracy of these methods. They are for reference only.

Animal Model:	lung tumor-bearing mice ^[3]
Dosage:	10 µg/mouse
Administration:	Intraperitoneal injection (i.p.)
Result:	Induced a significant decrease in the growth of pulmonary metastases in tumor-bearing mice. Reduced the amount of lung foci to ≈ 40%. Significantly increased BAL fluid cell numbers. Increased the level of INF-γ and IL-17A, decreased the levels of IL-13. Increased TLR3 expression.

Animal Model:	Middle cerebral artery occlusion (MCAO) model mice ^[5]
Dosage:	1.25 mg/kg
Administration:	Intraperitoneal injection (i.p.)
Result:	Reduced focal cerebral I/R injury. Increased the expression of Bcl2, Hsp27, and Hsp70, decreased Bax expression, and reduced cellular degeneration and apoptosis. Protected against cerebral ischemia and conferred protection against cerebral I/R injury through the downregulation of TLR4 signaling via TLR3.

CUSTOMER VALIDATION

- Adv Funct Mater. 29 August 2022.
- Chem Eng J. 2021 Aug 15;418:129392.
- Phytomedicine. 2021, 153495.
- Liver Int. 2022 Oct 17.
- Mol Ther Oncolytics. 25 August 2022.

See more customer validations on www.MedChemExpress.com

REFERENCES

[1]. Cheng Y, Xu F. Anticancer function of polyinosinic-polycytidylic acid [J]. Cancer biology & therapy, 2010, 10(12): 1219-1223.

[2]. Deleidi M, Hallett P J, Koprach J B, et al. The Toll-like receptor-3 agonist polyinosinic: polycytidylic acid triggers nigrostriatal dopaminergic degeneration [J]. Journal of Neuroscience, 2010, 30(48): 16091-16101.

[3]. Forte G, Rega A, Morello S, et al. Polyinosinic-polycytidylic acid limits tumor outgrowth in a mouse model of metastatic lung cancer [J]. The Journal of Immunology,

[4]. Rezaee F, Meednu N, Emo J A, et al. Polyinosinic: polycytidylic acid induces protein kinase D-dependent disassembly of apical junctions and barrier dysfunction in airway epithelial cells [J]. *Journal of Allergy and Clinical Immunology*, 2011, 128(6): 1216-1224. e11.

[5]. Wang P F, Fang H, Chen J, et al. Polyinosinic-polycytidylic acid has therapeutic effects against cerebral ischemia/reperfusion injury through the downregulation of TLR4 signaling via TLR3 [J]. *The Journal of Immunology*, 2014, 192(10): 4783-4794.

[6]. Alexopoulou L, Holt AC, Medzhitov R, Flavell RA. Recognition of double-stranded RNA and activation of NF-kappaB by Toll-like receptor 3. *Nature*. 2001;413(6857):732-738.

[7]. Matsumoto M, Kikkawa S, Kohase M, Miyake K, Seya T. Establishment of a monoclonal antibody against human Toll-like receptor 3 that blocks double-stranded RNA-mediated signaling. *Biochem Biophys Res Commun*. 2002;293(5):1364-1369.

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

Tel: 609-228-6898

Fax: 609-228-5909

E-mail: tech@MedChemExpress.com

Address: 1 Deer Park Dr, Suite Q, Monmouth Junction, NJ 08852, USA