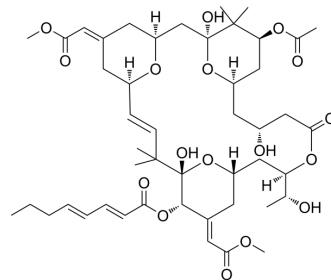


Bryostatin 1

Cat. No.:	HY-105231		
CAS No.:	83314-01-6		
Molecular Formula:	C ₄₇ H ₆₈ O ₁₇		
Molecular Weight:	905.03		
Target:	PKC; HIV; Bacterial		
Pathway:	Epigenetics; TGF-beta/Smad; Anti-infection		
Storage:	Powder	-20°C	3 years
	In solvent	-80°C	6 months
		-20°C	1 month



BIOLOGICAL ACTIVITY

Description	<p>Bryostatin 1 is a natural macrolide isolated from the bryozoan <i>Bugula neritina</i> and is a potent and central nervous system (CNS)-permeable PKC modulator. Bryostatin 1 binds to the isolated C1 domain of Munc13-1 and the full-length Munc13-1 protein with K_Ds of 8.07 nM and 0.45 nM, respectively. Bryostatin 1 has anti-cancer, anti-inflammatory, neuroprotective, anti-HIV-1 infection properties^{[1][2][3][4]}.</p>									
IC₅₀ & Target	HIV-1									
In Vitro	<p>Bryostatin 1 (1 μM; 5 minutes; HT22 cells) treatment successfully recruits Munc13-1 from the cytosol to the plasma membrane. Effects of Bryostatin 1 on the other Munc13 family members, ubMunc13-2 and bMunc13-2, resembled those of Munc13-1 for translocation^[1].</p> <p>The increased level of expression of Munc13-1 following a 24 h incubation with Bryostatin 1 in both HT22 and primary mouse hippocampal cells is observed^[1].</p> <p>Bryostatin 1 can also affect the immune system by modulating dendritic cells (DCs) via toll-like receptor 4 (TLR4) through the MyD88-independent pathway, which favors an anti-inflammatory environment by inducing a type 2 phenotype that promotes the differentiation of CD4⁺ T-helper (Th) lymphocytes into Th2 versus Th1 effector cells^[2].</p> <p>MCE has not independently confirmed the accuracy of these methods. They are for reference only.</p> <p>Western Blot Analysis^[1]</p> <table border="1"> <tr> <td>Cell Line:</td> <td>HT22 cells</td> </tr> <tr> <td>Concentration:</td> <td>1 μM</td> </tr> <tr> <td>Incubation Time:</td> <td>5 minutes</td> </tr> <tr> <td>Result:</td> <td>Caused Munc13-1 to transfer to the membrane fraction.</td> </tr> </table>		Cell Line:	HT22 cells	Concentration:	1 μM	Incubation Time:	5 minutes	Result:	Caused Munc13-1 to transfer to the membrane fraction.
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Concentration:	1 μM									
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Result:	Caused Munc13-1 to transfer to the membrane fraction.									
In Vivo	<p>Bryostatin 1 (30 μg/kg; intraperitoneal injection; 3 d per week; for 2 weeks; C57BL/6J mice) treatment abolishes the onset of EAE^[2].</p> <p>MCE has not independently confirmed the accuracy of these methods. They are for reference only.</p> <table border="1"> <tr> <td>Animal Model:</td> <td>Female C57BL/6J mice (8-12-week-old) with MOG₃₅₋₅₅^[2]</td> </tr> </table>		Animal Model:	Female C57BL/6J mice (8-12-week-old) with MOG ₃₅₋₅₅ ^[2]						
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Dosage:	30 µg/kg
Administration:	Intraperitoneal injection; 3 d per week; for 2 weeks
Result:	Abolished the onset of experimental autoimmune encephalomyelitis (EAE).

CUSTOMER VALIDATION

- Sci Transl Med. 2018 Jul 18;10(450):eaaq1093.

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REFERENCES

- [1]. Blanco FA, et al. Munc13 Is a Molecular Target of Bryostatin 1. *Biochemistry*. 2019 Jul 9;58(27):3016-3030.
- [2]. Kornberg MD, et al. Bryostatin-1 alleviates experimental multiple sclerosis. *Proc Natl Acad Sci U S A*. 2018 Feb 27;115(9):2186-2191.
- [3]. Zeng N, et al. Bryostatin 1 causes attenuation of TPA-mediated tumor promotion in mouse skin. *Mol Med Rep*. 2018 Jan;17(1):1077-1082.
- [4]. Proust A, et al. HIV-1 infection and latency-reversing agents bryostatin-1 and JQ1 disrupt amyloid beta homeostasis in human astrocytes. *Glia*. 2020 Apr 6.

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

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