

Product Introduction

BIO

BIO (6-bromoindirubin-3'-oxime) is a specific inhibitor of **GSK-3** with **IC50** of 5 nM for GSK-3a/ β , shows >16-fold selectivity over CDK5, also a pan-**JAK** inhibitor.

Technical Data:

Molecular Weight (MW):	356.17	
Formula:	C ₁₆ H ₁₀ BrN ₃ O ₂	HO N Br
Solubility (25°C)	DMSO 71 mg/mL	
* <1 mg/ml means slightly	Water <1 mg/mL	
soluble or insoluble:	Ethanol 21 mg/mL	
Purity:	>98%	
Storage:	3 years -20°C Powder 6 months-80°C in DMSO	
CAS No.:	667463-62-9	

Biological Activity

BIO (6-bromoindirubin-3'-oxime) is a specific inhibitor of glycogen synthase kinase-3 (GSK-3), with IC50 of 5 nM for GSK-3 α / β , shows >16-fold selectivity over CDK5. BIO interacts within the ATP binding pocket of these kinases, reduces β -catenin phosphorylation on a GSK-3-specific site in cellular models, closely mimicks Wnt signaling in Xenopus embryos. [1] In human and mouse embryonic stem cells, BIO maintains the undifferentiated phenotype and sustains expression of the pluripotent state-specific transcription factors Oct-3/4, Rex-1 and Nanog. BIO-mediated Wnt activation is functionally reversible, as withdrawal of Note: Products protected by valid patents are not offered for sale in countries where the sale of such products constitutes a patent infringement and its liability is at buyer's risk. This item is only

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the compound leads to normal multidifferentiation programs in both human and mouse embryonic stem cells. $^{[2]}$ BIO promotes proliferation in mammalian cardiomyocytes. $^{[3]}6BIO$ is also a pan-JAK inhibitor, with IC50 values of 0.03, 1.5, 8.0, 0.5 μM for TYK2, JAK1, JAK2 and JAK3. BIO selectively inhibits phosphorylation of STAT3 and induces apoptosis of human melanoma cells. $^{[4]}$

BIO suppresses melanoma tumor growth in a mouse xenograft model. [4]

The first pharmacological agent shown to maintain self-renewal in human and mouse embryonic stem cells.

References

- [1] Meijer L, et al. Chem Biol, 2003, 10(12), 1255-1266.
- [2] Sato N, et al. Nat Med, 2004, 10(1), 55-63.
- [3] Tseng AS, et al. Chem Biol, 2006, 13(9), 957-963.
- [4] Liu L, et al. Cancer Res. 2011 Jun 1;71(11):3972-3279.

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