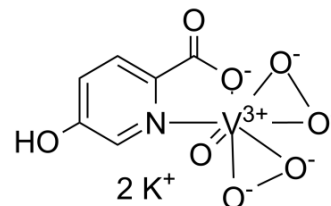


## BpV(HOPic)

Cat. No.:	HY-128693		
CAS No.:	722494-26-0		
Molecular Formula:	C <sub>6</sub> H <sub>4</sub> K <sub>2</sub> NO <sub>8</sub> V		
Molecular Weight:	347.24		
Target:	PTEN		
Pathway:	PI3K/Akt/mTOR		
Storage:	Powder	-20°C	3 years
	In solvent	-80°C	6 months
		-20°C	1 month



### BIOLOGICAL ACTIVITY

<b>Description</b>	BpV(HOPic) is a potent and selective inhibitor of PTEN with an IC <sub>50</sub> of 14 nM. Nanocarrier-BpV(HOPic) has neuroprotective activity <sup>[1][2]</sup> .															
<b>IC<sub>50</sub> &amp; Target</b>	IC <sub>50</sub> : 14 nM (PTEN) <sup>[1]</sup>															
<b>In Vitro</b>	<p>BpV(HOPic) (1 μM) treatment increases cell proliferation and decreases apoptotic rate in MG63 cells received Cisplatin treatment<sup>[3]</sup>.</p> <p>Bpv(HOPic) (1 μM) enhances migration of C2C12 myoblasts and is associated with activation of PI3K/AKT and MAPK/ERK signalling pathways<sup>[4]</sup>.</p> <p>BpV(HOPic) (1 μM; 48 hours) promotes the initiation of swine follicle growth and development, similar as in rodent species and humans<sup>[5]</sup>.</p> <p>Nanocarrier-BpV(HOPic) enhances axonal outgrowth of neurons<sup>[2]</sup>.</p>															
<b>In Vivo</b>	<p>BpV(HOPic) (0.05 mg/kg; i.p.) at reperfusion ameliorates liver ischemia/reperfusion (I/R) injury in vivo<sup>[6]</sup>.</p> <p>BpV(HOPic) (200 μg/kg; i.p.) exacerbates renal dysfunction and promotes tubular damage in mice with ischemia/reperfusion injury (IRI)<sup>[7]</sup>.</p> <table border="1" style="width: 100%; border-collapse: collapse;"> <tr> <td style="width: 20%;"><b>Animal Model:</b></td> <td>Male Wistar rats are subjected to partial hepatic ischemia<sup>[6]</sup></td> </tr> <tr> <td><b>Dosage:</b></td> <td>0.05 mg/kg</td> </tr> <tr> <td><b>Administration:</b></td> <td>I.p. injections at the start of reperfusion</td> </tr> <tr> <td><b>Result:</b></td> <td>Ameliorated reoxygenation injury and reproduced the hepatoprotective effects obtained by adenosine A2A receptor stimulation.</td> </tr> </table> <table border="1" style="width: 100%; border-collapse: collapse;"> <tr> <td style="width: 20%;"><b>Animal Model:</b></td> <td>Male C57BL/6 mice (8-12 weeks old; 20-30 g ) are subjected to renal ischemia<sup>[7]</sup></td> </tr> <tr> <td><b>Dosage:</b></td> <td>200 μg/kg</td> </tr> <tr> <td><b>Administration:</b></td> <td>I.p. injections 1 h before ischemia and then administers every 6 h after ischemia for 24</td> </tr> </table>		<b>Animal Model:</b>	Male Wistar rats are subjected to partial hepatic ischemia <sup>[6]</sup>	<b>Dosage:</b>	0.05 mg/kg	<b>Administration:</b>	I.p. injections at the start of reperfusion	<b>Result:</b>	Ameliorated reoxygenation injury and reproduced the hepatoprotective effects obtained by adenosine A2A receptor stimulation.	<b>Animal Model:</b>	Male C57BL/6 mice (8-12 weeks old; 20-30 g ) are subjected to renal ischemia <sup>[7]</sup>	<b>Dosage:</b>	200 μg/kg	<b>Administration:</b>	I.p. injections 1 h before ischemia and then administers every 6 h after ischemia for 24
<b>Animal Model:</b>	Male Wistar rats are subjected to partial hepatic ischemia <sup>[6]</sup>															
<b>Dosage:</b>	0.05 mg/kg															
<b>Administration:</b>	I.p. injections at the start of reperfusion															
<b>Result:</b>	Ameliorated reoxygenation injury and reproduced the hepatoprotective effects obtained by adenosine A2A receptor stimulation.															
<b>Animal Model:</b>	Male C57BL/6 mice (8-12 weeks old; 20-30 g ) are subjected to renal ischemia <sup>[7]</sup>															
<b>Dosage:</b>	200 μg/kg															
<b>Administration:</b>	I.p. injections 1 h before ischemia and then administers every 6 h after ischemia for 24															

---

	hr
<b>Result:</b>	Raised the level of serum creatinine and blood serum urea nitrogen.

---

## REFERENCES

- [1]. Schmid AC, et, al. Bisperoxovanadium compounds are potent PTEN inhibitors. *FEBS Lett.* 2004 May 21; 566(1-3): 35-8.
- [2]. Zhang B, et, al. Silencing of miR-19a-3p enhances osteosarcoma cells chemosensitivity by elevating the expression of tumor suppressor PTEN. *Oncol Lett.* 2019 Jan; 17(1): 414-421.
- [3]. Dimchev GA, et, al. Phospho-tyrosine phosphatase inhibitor Bpv(Hopic) enhances C2C12 myoblast migration in vitro. Requirement of PI3K/AKT and MAPK/ERK pathways. *J Muscle Res Cell Motil.* 2013 May; 34(2): 125-36.
- [4]. Raffel N, et, al. The effect of bpV(HOpic) on in vitro activation of primordial follicles in cultured swine ovarian cortical strips. *Reprod Domest Anim.* 2019 Aug; 54(8): 1057-1063.
- [5]. Ponte CD, et, al. Pharmacological postconditioning protects against hepatic ischemia/reperfusion injury. *Liver Transpl.* 2011 Apr; 17(4): 474-82.
- [6]. Zhou J, et, al. Pharmacological Inhibition of PTEN Aggravates Acute Kidney Injury. *Sci Rep.* 2017 Aug 25; 7(1): 9503.
- [7]. Kim MS, et, al. Nanotherapeutics of PTEN Inhibitor with Mesoporous Silica Nanocarrier Effective for Axonal Outgrowth of Adult Neurons. *ACS Appl Mater Interfaces.* 2016 Jul 27; 8(29): 18741-53.
- 

**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](mailto:tech@MedChemExpress.com)

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