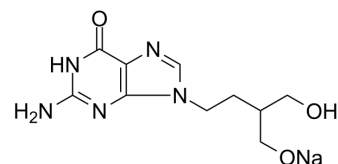


## Penciclovir sodium

Cat. No.:	HY-107739
CAS No.:	97845-62-0
Molecular Formula:	C <sub>10</sub> H <sub>14</sub> N <sub>5</sub> NaO <sub>3</sub>
Molecular Weight:	275.24
Target:	HSV
Pathway:	Anti-infection
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.



### BIOLOGICAL ACTIVITY

<b>Description</b>	Penciclovir (VSA 671) sodium is a potent and selective anti-herpesvirus agent with EC <sub>50</sub> values of 0.5, 0.8 µg/ml for HSV-1 (HFEM), HSV-2 (MS), respectively. Penciclovir sodium shows anti-herpesvirus activity with no-toxic. Penciclovir sodium prevents mortality in mouse <sup>[1][2]</sup> .									
<b>IC<sub>50</sub> &amp; Target</b>	HSV-1 0.5 µg/mL (EC50)	HSV-2 0.8 µg/mL (EC50)								
<b>In Vitro</b>	<p>Penciclovir sodium (0-100 µg/ml) shows anti-herpesvirus activity with EC<sub>50</sub>s of 0.5, 0.8, 2.4, 52, 100, 1.2, 1.6, 0.9, &gt;100 µg/ml for HSV-1 (HFEM), HSV-2 (MS), VZV (Ellen), CMV (AD-169), BHV-1 (Oxford 1964), BHV-2 (New York 1), EHV-1(Quai Hals), FHV-1 (B927), SVV, respectively<sup>[1]</sup>.</p> <p>Penciclovir sodium (0-100 µg/ml) shows no-toxic to uninfected human cells<sup>[1]</sup>.</p> <p>MCE has not independently confirmed the accuracy of these methods. They are for reference only.</p> <p>Cell Cytotoxicity Assay<sup>[1]</sup></p> <table border="1"> <tr> <td>Cell Line:</td> <td>MRC-5,WISH, Flow 4000, Flow 12000, RPMI 2650, WI-38, WI-38 VA13, Hs68, HEp-2, RD, SCC-13,Primary cells</td> </tr> <tr> <td>Concentration:</td> <td>0-100 µg/ml</td> </tr> <tr> <td>Incubation Time:</td> <td></td> </tr> <tr> <td>Result:</td> <td>Showed no-toxic with IC<sub>50</sub>s of &gt;100, &gt;100, 100, &gt;100, 60, &gt;100, &gt;100, &gt;100, &gt;100, &gt;100, &gt;100 for MRC-5,WISH, Flow 4000, Flow 12000, RPMI 2650, WI-38, WI-38 VA13, Hs68, HEp-2, RD, SCC-13,Primary cells, respectively.</td> </tr> </table>		Cell Line:	MRC-5,WISH, Flow 4000, Flow 12000, RPMI 2650, WI-38, WI-38 VA13, Hs68, HEp-2, RD, SCC-13,Primary cells	Concentration:	0-100 µg/ml	Incubation Time:		Result:	Showed no-toxic with IC <sub>50</sub> s of >100, >100, 100, >100, 60, >100, >100, >100, >100, >100, >100 for MRC-5,WISH, Flow 4000, Flow 12000, RPMI 2650, WI-38, WI-38 VA13, Hs68, HEp-2, RD, SCC-13,Primary cells, respectively.
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Concentration:	0-100 µg/ml									
Incubation Time:										
Result:	Showed no-toxic with IC <sub>50</sub> s of >100, >100, 100, >100, 60, >100, >100, >100, >100, >100, >100 for MRC-5,WISH, Flow 4000, Flow 12000, RPMI 2650, WI-38, WI-38 VA13, Hs68, HEp-2, RD, SCC-13,Primary cells, respectively.									
<b>In Vivo</b>	<p>Penciclovir sodium (100 mg/kg; s.c.; daily for 5 days) prevents mortality in mouse<sup>[2]</sup>.</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>Three-week-old female Balb/c mice<sup>[2]</sup></td> </tr> <tr> <td>Dosage:</td> <td>100 mg/kg</td> </tr> <tr> <td>Administration:</td> <td>S.c.; daily for 5 days</td> </tr> </table>		Animal Model:	Three-week-old female Balb/c mice <sup>[2]</sup>	Dosage:	100 mg/kg	Administration:	S.c.; daily for 5 days		
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Result:

Reduced virus titres in the respiratory organs of both w/t- and PR3-infected mice.

## CUSTOMER VALIDATION

- Microchem J. 2021, 106587.

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## REFERENCES

- [1]. M.R. Boyd, et al. Penciclovir: a review of its spectrum of activity, selectivity, and cross-resistance pattern. *Antiviral Chemistry and Chemotherapy*, 1993, (1): 3-11.
- [2]. de la Fuente R, et al. The acyclic nucleoside analogue penciclovir is a potent inhibitor of equine herpesvirus type 1 (EHV-1) in tissue culture and in a murine model. *Antiviral Res.* 1992 May;18(1):77-89.

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

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