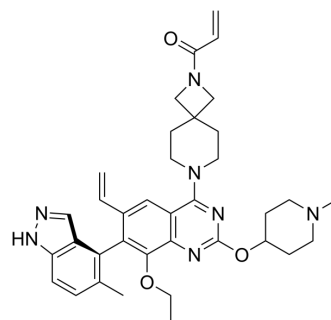


ASP6918

Cat. No.:	HY-162249
Molecular Formula:	C ₃₆ H ₄₃ N ₇ O ₃
Molecular Weight:	621.77
Target:	Ras
Pathway:	GPCR/G Protein; MAPK/ERK Pathway
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.



BIOLOGICAL ACTIVITY

Description	ASP6918 is a potent and orally active KRAS G12C inhibitor with an IC ₅₀ value of 0.028 μM. ASP6918 inhibits cell growth. ASP6918 shows antitumor activity ^[1] .									
IC₅₀ & Target	KRas G12C 0.028 μM (IC ₅₀)									
In Vitro	<p>ASP6918 (compound 29a) (6 days) inhibits cell growth with an IC₅₀ value of 0.0061 μM for NCI-H1373 cells^[1]. MCE has not independently confirmed the accuracy of these methods. They are for reference only.</p> <p>Cell Viability Assay^[1]</p> <table border="1"> <tr> <td>Cell Line:</td> <td>NCI-H1373 cells</td> </tr> <tr> <td>Concentration:</td> <td></td> </tr> <tr> <td>Incubation Time:</td> <td>6 days</td> </tr> <tr> <td>Result:</td> <td>Inhibited cell growth with an IC₅₀ value of 0.0061 μM.</td> </tr> </table>		Cell Line:	NCI-H1373 cells	Concentration:		Incubation Time:	6 days	Result:	Inhibited cell growth with an IC ₅₀ value of 0.0061 μM.
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Concentration:										
Incubation Time:	6 days									
Result:	Inhibited cell growth with an IC ₅₀ value of 0.0061 μM.									
In Vivo	<p>ASP6918 (10, 20, 40, 60 mg/kg; p.o.; daily for 13 days) shows antitumor activity in NCI-H1373 xenograft mouse model^[1]. 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>Four-week-old male nude mice (NCI-H1373 xenograft mouse model)^[1]</td> </tr> <tr> <td>Dosage:</td> <td>10, 20, 40, 60 mg/kg</td> </tr> <tr> <td>Administration:</td> <td>Oral, daily for 13 days</td> </tr> <tr> <td>Result:</td> <td>Inhibited of tumor growth with tumor growth inhibition (TGI) rate of 27%, 68%, 49%, 73% at 10, 20, 40, 60 mg/kg.</td> </tr> </table>		Animal Model:	Four-week-old male nude mice (NCI-H1373 xenograft mouse model) ^[1]	Dosage:	10, 20, 40, 60 mg/kg	Administration:	Oral, daily for 13 days	Result:	Inhibited of tumor growth with tumor growth inhibition (TGI) rate of 27%, 68%, 49%, 73% at 10, 20, 40, 60 mg/kg.
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REFERENCES

[1]. Imaizumi T, et al. Discovery of ASP6918, a KRAS G12C inhibitor: Synthesis and structure-activity relationships of 1-{2,7-diazaspiro[3.5]non-2-yl}prop-2-en-1-one derivatives as covalent inhibitors with good potency and oral activity for the treatment of solid tumors. *Bioorg Med Chem*. 2024 Jan 15;98:117581.

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

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