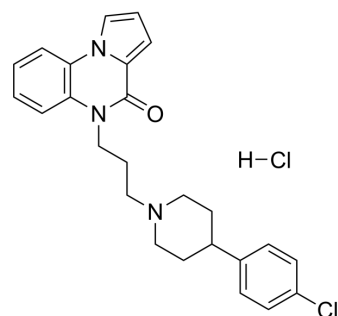


JMS-17-2 hydrochloride

Cat. No.:	HY-123918A
CAS No.:	2341841-07-2
Molecular Formula:	C ₂₅ H ₂₇ Cl ₂ N ₃ O
Molecular Weight:	456.41
Target:	CX3CR1
Pathway:	Immunology/Inflammation
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.



BIOLOGICAL ACTIVITY

Description	JMS-17-2 hydrochloride is a potent and selective CX3CR1 antagonist with an IC ₅₀ of 0.32 nM. JMS-17-2 hydrochloride impairs metastatic seeding and colonization of breast cancer cells ^[1] .								
IC₅₀ & Target	IC50: 0.32 nM (CX3CR1) ^[1]								
In Vivo	<p>JMS-17-2 (10 mg/kg; administered i.p.; twice a day for three weeks) causes a dramatic reduction of tumors in both skeleton and visceral organs in SCID mice^[1].</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>SCID mice (~25g) with MDA-231 xenograft^[1]</td> </tr> <tr> <td>Dosage:</td> <td>10 mg/kg</td> </tr> <tr> <td>Administration:</td> <td>Administered i.p.; twice a day for three weeks</td> </tr> <tr> <td>Result:</td> <td>Caused a dramatic reduction of tumors in both skeleton and visceral organs.</td> </tr> </table>	Animal Model:	SCID mice (~25g) with MDA-231 xenograft ^[1]	Dosage:	10 mg/kg	Administration:	Administered i.p.; twice a day for three weeks	Result:	Caused a dramatic reduction of tumors in both skeleton and visceral organs.
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Dosage:	10 mg/kg								
Administration:	Administered i.p.; twice a day for three weeks								
Result:	Caused a dramatic reduction of tumors in both skeleton and visceral organs.								

CUSTOMER VALIDATION

- Cancer Res. 2022 Sep 14;CAN-22-1199.

See more customer validations on www.MedChemExpress.com

REFERENCES

[1]. Shen F, et al. Novel Small-Molecule CX3CR1 Antagonist Impairs Metastatic Seeding and Colonization of Breast Cancer Cells. Mol Cancer Res. 2016 Jun;14(6):518-27.

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

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