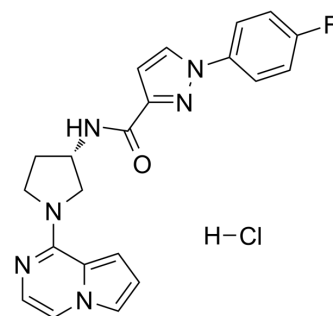


## CXCR7 antagonist-1 hydrochloride

<b>Cat. No.:</b>	HY-139643A
<b>Molecular Formula:</b>	C <sub>21</sub> H <sub>20</sub> ClFN <sub>6</sub> O
<b>Molecular Weight:</b>	426.87
<b>Target:</b>	CXCR
<b>Pathway:</b>	GPCR/G Protein; Immunology/Inflammation
<b>Storage:</b>	4°C, sealed storage, away from moisture * In solvent : -80°C, 6 months; -20°C, 1 month (sealed storage, away from moisture)



### SOLVENT & SOLUBILITY

#### In Vitro

DMSO : 350 mg/mL (819.92 mM; Need ultrasonic)

Concentration	Mass			
	1 mg	5 mg	10 mg	
1 mM	2.3426 mL	11.7132 mL	23.4263 mL	
5 mM	0.4685 mL	2.3426 mL	4.6853 mL	
10 mM	0.2343 mL	1.1713 mL	2.3426 mL	

Please refer to the solubility information to select the appropriate solvent.

### BIOLOGICAL ACTIVITY

#### Description

CXCR7 antagonist-1 hydrochloride is a CXCR7 antagonist that inhibits the binding of the SDF-1 chemokine (also known as the CXCL12 chemokine) or I-TAC (also known as CXCL11) to the chemokine receptor CXCR7. CXCR7 antagonist-1 hydrochloride is useful in preventing tumor cell proliferation, tumor formation, inflammatory diseases, and many other diseases (extracted from patent WO2014085490A1, compound 1.128)<sup>[1]</sup>.

#### IC<sub>50</sub> & Target

CXCR7

### REFERENCES

[1]. Junfa Fan, et al. Cxcr7 antagonists. Patent WO2014085490A1.

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

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