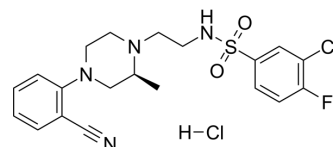


## VU6036720 hydrochloride

**Cat. No.:** HY-148304A  
**Molecular Formula:** C<sub>20</sub>H<sub>23</sub>Cl<sub>2</sub>FN<sub>4</sub>O<sub>2</sub>S  
**Molecular Weight:** 473.39  
**Target:** Potassium Channel  
**Pathway:** Membrane Transporter/Ion Channel  
**Storage:** 4°C, sealed storage, away from moisture and light  
 \* In solvent : -80°C, 6 months; -20°C, 1 month (sealed storage, away from moisture and light)



### SOLVENT & SOLUBILITY

#### In Vitro

DMSO : 40 mg/mL (84.50 mM; ultrasonic and warming and heat to 60°C)

Concentration	Mass			
	1 mg	5 mg	10 mg	
1 mM	2.1124 mL	10.5621 mL	21.1242 mL	
5 mM	0.4225 mL	2.1124 mL	4.2248 mL	
10 mM	0.2112 mL	1.0562 mL	2.1124 mL	

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

### BIOLOGICAL ACTIVITY

#### Description

VU6036720 hydrochloride is a potent and specific in vitro inhibitor of Kir4.1/5.1. VU6036720 hydrochloride can inhibit Kir4.1/5.1 channels with an IC<sub>50</sub> value of 0.24 μM. VU6036720 hydrochloride can be used for the research of brain and kidney [1].

### REFERENCES

[1]. Samantha J McClenahan, et al. VU6036720: The First Potent and Selective In Vitro Inhibitor of Heteromeric Kir4.1/5.1 Inward Rectifier Potassium Channels. Mol Pharmacol. 2022 May;101(5):357-370.

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

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