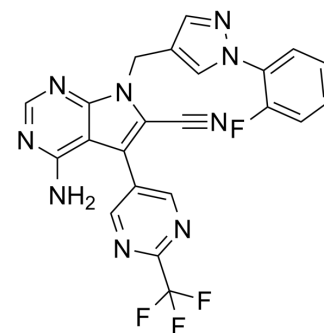


## CFTR corrector 6

<b>Cat. No.:</b>	HY-136939
<b>CAS No.:</b>	2226970-01-8
<b>Molecular Formula:</b>	C <sub>22</sub> H <sub>13</sub> F <sub>4</sub> N <sub>9</sub>
<b>Molecular Weight:</b>	479.39
<b>Target:</b>	CFTR
<b>Pathway:</b>	Membrane Transporter/Ion Channel
<b>Storage:</b>	4°C, protect from light * In solvent : -80°C, 6 months; -20°C, 1 month (protect from light)



### SOLVENT & SOLUBILITY

<b>In Vitro</b>	DMSO : 100 mg/mL (208.60 mM; Need ultrasonic)					
	<b>Preparing Stock Solutions</b>	<b>Solvent</b>	<b>Mass</b>	<b>1 mg</b>	<b>5 mg</b>	<b>10 mg</b>
		<b>Concentration</b>				
		<b>1 mM</b>		2.0860 mL	10.4299 mL	20.8598 mL
		<b>5 mM</b>		0.4172 mL	2.0860 mL	4.1720 mL
<b>10 mM</b>		0.2086 mL	1.0430 mL	2.0860 mL		
Please refer to the solubility information to select the appropriate solvent.						
<b>In Vivo</b>	1. Add each solvent one by one: 10% DMSO >> 40% PEG300 >> 5% Tween-80 >> 45% saline Solubility: 2.5 mg/mL (5.21 mM); Suspended solution; Need ultrasonic					

### BIOLOGICAL ACTIVITY

<b>Description</b>	CFTR corrector 6 is a potent potentiator of Cystic Fibrosis Transmembrane conductance Regulator (CFTR). CFTR corrector 6 has the potential for cystic fibrosis (CF) and other CFTR associated disorders research <sup>[1]</sup> .
<b>In Vitro</b>	CFTR corrector 6 (Example 27) has EC <sub>50</sub> s of 1.25 nM and 1.27 nM for primary cystic fibrosis human bronchial epithelial (CF hBE) cells and fischer rat thyroid (FRT) cell lines <sup>[1]</sup> . MCE has not independently confirmed the accuracy of these methods. They are for reference only.

### REFERENCES

[1]. Joseph Walter Strohbach, et al. Pyrrolopyrimidines as cftr potentiators. WO2018094137A1

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

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