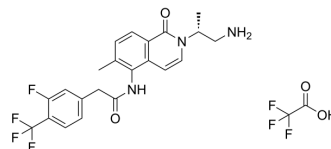


## P2X7-IN-2 TFA

Cat. No.:	HY-18725A
Molecular Formula:	C <sub>24</sub> H <sub>22</sub> F <sub>7</sub> N <sub>3</sub> O <sub>4</sub>
Molecular Weight:	549.44
Target:	P2X Receptor; Interleukin Related
Pathway:	Membrane Transporter/Ion Channel; Immunology/Inflammation
Storage:	4°C, protect from light, stored under nitrogen * In solvent : -80°C, 6 months; -20°C, 1 month (protect from light, stored under nitrogen)



### SOLVENT & SOLUBILITY

#### In Vitro

DMSO : 100 mg/mL (182.00 mM; ultrasonic and warming and heat to 60°C)

Preparing Stock Solutions	Solvent Concentration	Mass		
		1 mg	5 mg	10 mg
	1 mM	1.8200 mL	9.1002 mL	18.2003 mL
	5 mM	0.3640 mL	1.8200 mL	3.6401 mL
	10 mM	0.1820 mL	0.9100 mL	1.8200 mL

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

### BIOLOGICAL ACTIVITY

#### Description

P2X7-IN-2 TFA (compound 58) is a P2X7 receptor inhibitor. P2X7-IN-2 TFA inhibits IL-1 $\beta$  release with an IC<sub>50</sub> value of 0.01 nM. P2X7-IN-2 TFA can be used for the research of autoimmunity, inflammation and cardiovascular disease<sup>[1]</sup>.

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

IC<sub>50</sub>: 0.01 nM (human whole blood IL-1 $\beta$  release)<sup>[1]</sup>

#### In Vitro

P2X7-IN-2 TFA (0-1  $\mu$ M;30 min) inhibits human whole blood IL-1 $\beta$  release with an IC<sub>50</sub> value of 0.01 nM<sup>[1]</sup>. MCE has not independently confirmed the accuracy of these methods. They are for reference only.

### REFERENCES

[1]. Kelly, Michael G, et al. Preparation of isoquinolonecarboxamides as P2X7 purinoceptor modulators and uses thereof. WO2008112205 A1. 2008.

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

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