## Pociredir

Cat. No.:	HY-139400		
CAS No.:	2490676-18	-9	
Molecular Formula:	C <sub>22</sub> H <sub>18</sub> FN <sub>5</sub> O <sub>2</sub>		
Molecular Weight:	403.41		
Target:	Histone Me	thyltrans	ferase
Pathway:	Epigenetics		
Storage:	Powder	-20°C	3 years
		4°C	2 years
	In solvent	-80°C	6 months
		-20°C	1 month

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## SOLVENT & SOLUBILITY

		Mass Solvent Concentration	1 mg	5 mg	10 mg		
	Preparing Stock Solutions	1 mM	2.4789 mL	12.3943 mL	24.7887 mL		
		5 mM	0.4958 mL	2.4789 mL	4.9577 mL		
		10 mM	0.2479 mL	1.2394 mL	2.4789 mL		
	Please refer to the so	lubility information to select the ap	propriate solvent.				
		t one by one: 10% DMSO >> 90% corn oil mg/mL (5.16 mM); Clear solution					
		d each solvent one by one: 10% DMSO >> 40% PEG300 >> 5% Tween-80 >> 45% saline lubility: ≥ 1 mg/mL (2.48 mM); Clear solution					
		nt one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline) ng/mL (2.48 mM); Clear solution					

BIOLOGICAL ACTIVITY				
Description	Pociredir (FTX-6058) is a potent and orally active inhibitor of Embryonic Ectoderm Development (EED). Pociredir can induce HbF protein expression in cell and murine models. Pociredir can be used for the research of select hemoglobinopathies, including sickle cell disease and β-thalassemia <sup>[1][2]</sup> .			
In Vitro	Pociredir inhibits PRC2 via binding to EED, which induces robust HbF protein expression in both cell and murine models <sup>[2]</sup> . MCE has not independently confirmed the accuracy of these methods. They are for reference only.			

**Product** Data Sheet

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## REFERENCES

[1]. Boerner LK. Virtual meeting delivers first time drug structures. April 12, 2021.

[2]. Fulcrum Therapeutics Presents Published Structure of Investigational Small Molecule FTX-6058 at the American Chemical Society (ACS) Spring 2021 Virtual Conference.

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

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