## Etavopivat

Cat. No.:	HY-139573		
CAS No.:	2245053-57	-8	
Molecular Formula:	C <sub>22</sub> H <sub>23</sub> N <sub>3</sub> O <sub>6</sub> S	5	
Molecular Weight:	457.5		
Target:	Pyruvate K	nase	
Pathway:	Metabolic E	nzyme/F	Protease
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

	Preparing Stock Solutions	Solvent Mass Concentration	1 mg	5 mg	10 mg
		1 mM	2.1858 mL	10.9290 mL	21.8579 mL
		5 mM	0.4372 mL	2.1858 mL	4.3716 mL
		10 mM	0.2186 mL	1.0929 mL	2.1858 mL

Description			
	Etavopivat is a potent, selective, and orally active erythrocyte pyruvate kinase (PKR) activator. Etavopivat has potent antisickling effects that can be used in studies of sickle cell disease and other haemoglobinopathies <sup>[1][2]</sup> .		
		oves haemoglobin-oxygen affinity and reduces the sickle point (PoS) in human red blood cells <sup>[1]</sup> . confirmed the accuracy of these methods. They are for reference only.	
	Etavopivat (3-22 mg/kg, p.o., mg/kg and 22 mg/kg <sup>[2]</sup> .	p.o., daily, 2 weeks) improves RBC survival and Hb levels in SCA mice <sup>[1]</sup> . , daily, 5 days) causes an increase in 2,3-DPG and ATP in crab-eating monkeys at doses of 8 confirmed the accuracy of these methods. They are for reference only.	
	Dosage:	500-1000 mg/kg	

## Product Data Sheet

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Administration:	p.o., daily, 2 weeks
Result:	Decreased the levels of 2,3-DPG.
	Increased ATP levels.
	Reduced sickling in vivo.

## REFERENCES

[1]. Schroeder P, et al. Etavopivat, a Pyruvate Kinase Activator in Red Blood Cells, for the Treatment of Sickle Cell Disease. J Pharmacol Exp Ther. 2022 Mar; 380(3):210-219.

[2]. Shrestha A, et al. FT-4202, an oral PKR activator, has potent antisickling effects and improves RBC survival and Hb levels in SCA mice. Blood Adv. 2021 May 11;5(9):2385-2390.

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

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