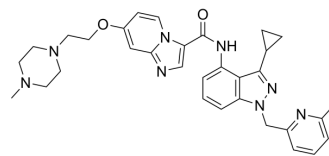


ARRY-382

Cat. No.:	HY-136362		
CAS No.:	1313407-95-2		
Molecular Formula:	C ₃₂ H ₃₆ N ₈ O ₂		
Molecular Weight:	564.68		
Target:	c-Fms		
Pathway:	Protein Tyrosine Kinase/RTK		
Storage:	Powder	-20°C	3 years
		4°C	2 years
	In solvent	-80°C	6 months
		-20°C	1 month



SOLVENT & SOLUBILITY

In Vitro	DMSO : 8.33 mg/mL (14.75 mM; Need ultrasonic)			
		Solvent Concentration	Mass	
			1 mg	5 mg
	Preparing Stock Solutions	1 mM	1.7709 mL	8.8546 mL
		5 mM	0.3542 mL	1.7709 mL
		10 mM	0.1771 mL	0.8855 mL
	Please refer to the solubility information to select the appropriate solvent.			
In Vivo	<ol style="list-style-type: none"> Add each solvent one by one: 10% DMSO >> 40% PEG300 >> 5% Tween-80 >> 45% saline Solubility: ≥ 0.83 mg/mL (1.47 mM); Clear solution Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline) Solubility: ≥ 0.83 mg/mL (1.47 mM); Clear solution Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: ≥ 0.83 mg/mL (1.47 mM); Clear solution 			

BIOLOGICAL ACTIVITY

Description	ARRY-382 is a potent, oral and highly selective inhibitor of CSF1R/c-Fms with an IC ₅₀ of 9 nM. ARRY-382 can be used for the research of advanced or metastatic cancers ^[1] .
IC₅₀ & Target	IC ₅₀ : 9 nM (CSF1R/c-Fms) ^[1]
In Vitro	ARRY-382 (1 μM) dramatically decreases the number of nurse-like cells (NLCs) compared to untreated cells ^[1] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.

REFERENCES

[1]. David K Edwards V, et al. Targeting of colony-stimulating factor 1 receptor (CSF1R) in the CLL microenvironment yields antineoplastic activity in primary patient samples. *Oncotarget*. 2018 May 15;9(37):24576-24589.

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

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