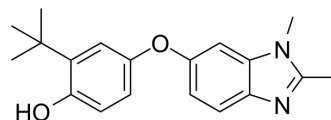


DS20362725

Cat. No.:	HY-143201		
CAS No.:	2735803-20-8		
Molecular Formula:	C ₁₉ H ₂₂ N ₂ O ₂		
Molecular Weight:	310.39		
Target:	Estrogen Receptor/ERR		
Pathway:	Vitamin D Related/Nuclear Receptor		
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 : 10 mg/mL (32.22 mM); ultrasonic and warming and heat to 60°C)				
		Solvent Concentration	Mass 1 mg	5 mg	10 mg
	Preparing Stock Solutions	1 mM	3.2218 mL	16.1088 mL	32.2175 mL
		5 mM	0.6444 mL	3.2218 mL	6.4435 mL
10 mM		0.3222 mL	1.6109 mL	3.2218 mL	
Please refer to the solubility information to select the appropriate solvent.					
In Vivo	1. Add each solvent one by one: 10% DMSO >> 40% PEG300 >> 5% Tween-80 >> 45% saline Solubility: ≥ 1 mg/mL (3.22 mM); Clear solution 2. Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: ≥ 1 mg/mL (3.22 mM); Clear solution				

BIOLOGICAL ACTIVITY

Description	DS20362725 is an estrogen-related receptor α (ERR α) agonist. DS20362725 inhibits the binding between receptor-interacting protein 140 (RIP140) corepressor peptide (10 nM) and GST-ERR α ligand-binding domain (LBD; 1.2 μ M) with an IC ₅₀ value of 0.6 μ M. DS20362725 can be used for the research of metabolic disorders, including type 2 diabetes mellitus (T2DM) ^[1] .
IC₅₀ & Target	ERR α
In Vitro	DS20362725 (compound 3d; 0.002, 0.006, 0.017, 0.051, 0.015, 0.046, 1.4, 4.2, 12.5 μ g/mL; 18 h) activates the transcriptional activity of full-length ERR α in MG63 cells with an EC ₅₀ of 1.1 μ M ^[1] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.

REFERENCES

[1]. Shinozuka T, et al. Discovery of a Novel Class of ERR α Agonists. ACS Med Chem Lett. 2021 Apr 21;12(5):817-821.

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

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