

Product Name: Lenalidomide (CC-5013) Revision Date: 06/13/2023



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Lenalidomide (CC-5013)

Cat. No.:	A4211	
CAS No.:	191732-72-6	
Formula:	C13H13N3O3	
M.Wt:	259.3	
Synonyms:	Revlimid,IMiD3,CC 5013,CDC-501,CDC 501	
Target:	Apoptosis	
Pathway:	TNF-α	
Storage:	Store at -20°C	

Solvent & Solubility

insoluble in EtOH; insoluble in H2O; \geq 100.8 mg/mL in DMSO Mass Solvent 1mg 5mg 10mg Preparing Concentration In Vitro Stock Solutions 1 mM 3.8565 mL 19.2827 mL 38.5654 mL 5 mM 3.8565 mL 0.7713 mL 7.7131 mL 10 mM 0.3857 mL 1.9283 mL 3.8565 mL

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

Biological Activity

Shortsummary	Antineoplastic agent,inhibits angiogenesis		
IC ₅₀ & Target	13 nM (TNF-α)		
In Vitro	Cell Viability Assay	C El trocarda	
	Cell Line: 1000 Cell	Peripheral blood mononuclear cells (PBMCs)	
	Preparation method:	The solubility of this compound in DMSO is >10 mM. General tips for obtaining	
		a higher concentration: Please warm the tube at 37 $^{\circ}\mathrm{C}$ for 10 minutes and/or	
		shake it in the ultrasonic bath for a while.Stock solution can be stored below	
		-20°C for several months.	
	Reacting conditions:	10 μM, 7 days	
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	Applications:	The cells were incubated with the dye at 37°C for 10 min and treated for 7 days
		in RPMI culture medium with lenalidomide. Cells were surface stained with
		anti-CD4-PerCP and anti-CD25-APC, followed by intracellular staining with
		anti-FOXP3-PE. Lenalidomide inhibited the expression of CD4+CD25high
	Blow	CTLA-4+FOXP3+ cells. Incubation with lenalidomide significantly decreases
	E soore me d.	expression of the T regulatory cell population after 7 days of culture. The drug
	and Paration."	decreased the percentage of CD4+CD25high cells expressing both CTLA-
	Real Providence	and FOXP3 from 25 to 12%.
	Animal experiment	
In Vivo	Animal models:	Male Sprague–Dawley rats
	Dosage form:	Oral administration, 50 mg/kg or 250 mg/kg
	Applications:	In the rat mesenteric window assay (RMWA), representative difference
		between vehicle and 50 or 250 mg/kg lenalidomide-treated rats were visualized
		by staining with an antibody against rat endothelium in bFGF-induced
	E to uncour	angiogenic windows. The induction of angiogenesis by bFGF was significantl
	Redon, Espare	inhibited by oral treatment of lenalidomide in a dose-dependent mannel
	Renew Der	Lenalidomide significantly decreased the percentage of vascularized area from
		5.16% in the control group to 2.58 and 1.69 in the 50 and 250 mg/kg group
		respectively.
	Other notes:	Please test the solubility of all compounds indoor, and the actual solubility ma
		slightly differ with the theoretical value. This is caused by an experimenta
		system error and it is normal.



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References

[1] Galustian C, Meyer B, Labarthe M C, et al. The anti-cancer agents lenalidomide and pomalidomide inhibit the proliferation and function of T regulatory cells. Cancer Immunology, Immunotherapy, 2009, 58(7): 1033-1045.

[2] Dredge K, Horsfall R, Robinson S P, et al. Orally administered lenalidomide (CC-5013) is anti-angiogenic in vivo and inhibits endothelial cell migration and Akt phosphorylation in vitro. Microvascular research, 2005, 69(1): 56-63.

Caution

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NOT FOR HUMAN, VETERINARY DIAGNOSTIC OR THERAPEUTIC USE.

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