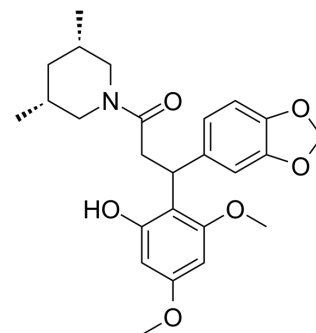


## (±)-ML 209

Cat. No.:	HY-126037		
CAS No.:	1334526-14-5		
Molecular Formula:	C <sub>25</sub> H <sub>31</sub> NO <sub>6</sub>		
Molecular Weight:	441.52		
Target:	ROR		
Pathway:	Metabolic Enzyme/Protease; Vitamin D Related/Nuclear Receptor		
Storage:	Powder	-20°C	3 years
	In solvent	-80°C	6 months
		-20°C	1 month



### SOLVENT & SOLUBILITY

#### In Vitro

DMSO : 100 mg/mL (226.49 mM; Need ultrasonic)

Concentration	Solvent	Mass		
		1 mg	5 mg	10 mg
Preparing Stock Solutions	1 mM	2.2649 mL	11.3245 mL	22.6490 mL
	5 mM	0.4530 mL	2.2649 mL	4.5298 mL
	10 mM	0.2265 mL	1.1325 mL	2.2649 mL

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

### BIOLOGICAL ACTIVITY

#### Description

(±)-ML 209 (compound 4n), a diphenylpropanamide, is a retinoic acid-related orphan receptor ROR $\gamma$  antagonist with an IC<sub>50</sub> of 1.1  $\mu$ M. (±)-ML 209 inhibits ROR $\gamma$ t transcriptional activity with an IC<sub>50</sub> of 300 nM in HEK293t cells. (±)-ML 209 inhibits the transcriptional activity of ROR $\gamma$ t, but not ROR $\alpha$  in cells. (±)-ML 209 selectively inhibits murine Th17 cell differentiation without affecting the differentiation of naïve CD4<sup>+</sup> T cells into other lineages, including Th1 and regulatory T cells<sup>[1]</sup>.

#### IC<sub>50</sub> & Target

ROR $\gamma$   
1.1  $\mu$ M (IC<sub>50</sub>)

### CUSTOMER VALIDATION

- Antiviral Res. 2023 Dec 4:105769.

See more customer validations on [www.MedChemExpress.com](http://www.MedChemExpress.com)

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## REFERENCES

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[1]. Jun R Huh, et al. Identification of Potent and Selective Diphenylpropanamide ROR $\gamma$  Inhibitors. ACS Med Chem Lett. 2013 Jan 10;4(1):79-84.

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

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