

Catalog # 10-1371 iCRT5

CAS# 18623-44-4 4-[5-(3,4-Dimethoxy-benzylidene)-4-oxo-2-thioxo-thiazolidin-3-yl]-butyric acid Lot # X104619



A selective CRT (β -catenin-responsive transcription) inhibitor.¹ IC₅₀=18 nM for Wnt responsive STF16 luciferase. Acts via interfering with the β -catenin, TCF4 interaction.¹ Displays poor inhibition of cell proliferation in triple-negative breast cancer cells.² Increases the surface expression of MHCII, CD80 and CD86 on unstimulated dendritic cells (DCs) with no detrimental effects on immune-phenotype of stimulated DCs.³ Modulators of Wnt signaling show great promise in animal models of several cancers.⁴ Cell permeable.

- 1) Gonsalves et al. (2011), An RNAi-based chemical genetic screen identifies three small-molecule inhibitors of the Wnt/wingless signaling pathway; Proc. Natl. Acad. Sci. USA, **108** 5954
- 2) Bilir et al. (2013), Wnt signaling blockage inhibits cell proliferation and migration, and induces apoptosis in triplenegative breast cancer cells; J. Transl. Med., **11** 280
- 3) Kafer *et al.* (2016), *Inhibitors of β-catenin affect the immune-phenotype and functions of dendritic cells in an inhibitor-specific manner;* Immunopharmacol., **32** 118
- 4) Anastas et al. (2013), WNT signaling pathways as therapeutic targets in cancer, Nat. Rev. Cancer, 13 11

PHYSICAL DATA

Molecular Weight:	367.44
Molecular Formula:	C ₁₆ H ₁₇ NO ₅ S ₂
Purity:	98% by TLC
	NMR: (Conforms)
Solubility:	Soluble in DMSO (up to 30 mg/ml)
Physical Description:	Yellow solid
Storage and Stability:	Store as supplied, desiccated at -20°C for up to 1 year from the date of purchase. Solutions in DMSO may be stored at -20°C for up to 2 months

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