

Catalog # 10-1506 SR-8278

1254944-66-5 1,2,3,4-Tetrahydro-2-[[5-(methylthio)-2-thienyl]carbonyl]-3-isoquinolinecarboxylic acid ethyl ester Lot # S102004



Antagonist of the nuclear heme receptor, REV-ERB. Blocks the ability of the REV-ERBα agonist, GSK4112, to enhance REV-ERB-dependent repression in a cotransfection assay.¹ Decreases glucagon secretion and intracellular calcium signals in alphaTC1-9 cells and mouse primary alpha cells.² Enhances the expression of bone morphogenetic protein genes in rat uterus endometrium stromal cells.³ Shortens the period of circadian oscillations in mature granulosa cells stimulated with or without LH.⁴

- 1) Kojetin et al. (2011), Identification of SR8278, a synthetic antagonist of the nuclear heme receptor REV-ERB; ACS Chem. Biol. **6** 131
- 2) Vieira et al. (2013), Involvement of the Clock Gene Rev-erb alpha in the Regulation of Glucagon Secretion in Pancreatic Alpha-cells; PLoS ONE **8** e69939
- 3) Tasaki et al. (2015), Inhibitory role of REV-ERBα in the expression of bone morphogenetic protein gene family in rat uterus endometrium stromal cells; Am. J. Physiol. Cell. Physiol., **308** C528
- 4) Chen et al. (2015), Integration of the nuclear receptor REV-ERBα linked with circadian oscillators in the expressions of Alas1, Pparg1a, and II6 genes in rat granulosa cells; Chronobiol. Int., **32** 739

PHYSICAL DATA

Molecular Weight:	361.49
Molecular Formula:	C18H19NO3S2
Purity:	98% by TLC
	NMR: (Conforms)
Solubility:	DMSO (up to 60 mg/ml), Ethanol (up to 35 mg/ml)
Physical Description:	White waxy solid
Storage and Stability:	Store as supplied desiccated at -20°C for up to 1 year from the date of purchase. Solutions in
	DMSO or ethanol may be stored at -20°C for up to 1 month.

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