

Catalog # 10-2575 AS-605240

CAS# 648450-29-7 5-(6-Quinoxalinylmethylene)-2,4-thiazolidine-2,4-dione Lot # S103024

Potent and selective inhibitor of PI 3-kinase γ (PI 3-K γ), IC₅₀ = 8 nM displaying 30-fold selectivity over PI 3-K δ and PI 3-K β and 7.5-fold selectivity over PI 3-K α^1 . Suppresses joint inflammation in two distinct mouse models¹. Ameliorates surgical brain injury², colitis³, intimal hyperplasia-induced inflammatory processes⁴ and renal injury/fibrosis⁵ in rodent models. Orally active.

- 1) Camps et al. (2005), Blockade of PI3Kgamma suppresses joint inflammation and damage in mouse models of rheumatoid arthritis; Nature Med., **11** 936
- 2) Huang et al. (2015), Phosphoinositide 3-Kinase Gamma Contributes to Neuroinflammation ina Rat Model of Surgical Brain Injury; J. Neurosci., **35** 10390
- 3) Peng et al. (2010), Inhibition of phosphoinositide 3-kinase ameliorates dextran sodium sulfate-induced colitis in mice; J. Pharmacol. Exp. Ther., **332** 46
- 4) Smimova et al. (2014), Targeting Pl3Kγ activity decreases vascular trauma-induced intimal hyperplasia through modulation of the Th1 response; J. Exp. Med., **211** 1779
- 5) Yu et al. (2018), A blockade of Pl3Kγ signaling effectively mitigates angiotensin II-induced renal injury and fibrosis in a mouse model; Sci. Rep., **8** 10988

PHYSICAL DATA

Molecular Weight: 257.27

NMR: (Conforms)

Solubility: DMSO (up to 1.5 mg/ml with warming)

Physical Description: Red solid

Storage and Stability: Store as supplied desiccated at -20°C for up to 2 years from the date of purchase. Solutions in

DMSO may be stored at -20°C for up to 1 month.

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