

Catalog # 10-1453 HTS01037

CAS# 682741-29-3
4-[(3-Carboxy-1-oxo-2-propen-1-yl)amino]-[2,2'-bithiophene]-5-carboxylic acid, 5-methyl ester
Lot # FBA4180

Inhibits fatty acid binding proteins. Inhibits lipolysis in 3T3-L1 adipocytes and reduces LPS-stimulated inflammation in cultured macrophages. Acts as an antagonist of the protein-protein interaction between AFABP/aP2 and hormone sensitive lipase but does not activate PPAR γ in macrophages¹. Inhibits FABP-dependent and fatty acid-stimulated leukotriene C₄ biosynthesis². Reduces intracellular free fatty acid levels, lowering macrophage inflammation and ER stress³. Reduces LPS-stimulated IL-1 β secretion in a mouse model⁴. Inhibits VLDL-induced foam cell formation⁵.

- 1) Hertzel et al. (2009), Identification and characterization of a small molecule inhibitor of Fatty Acid binding proteins; J. Med. Chem, **52** 6024
- 2) Long et al. (2012), Fatty acids induce leukotriene C4 synthesis in macrophages in a fatty acid binding protein-dependent manner, Biochim. Biophys. Acta, **1831** 1199
- 3) Xu et al. (2015), Uncoupling lipid metabolism from inflammation through fatty acid binding protein-dependent expression of USP2; Mol. Cell. Biol., **35** 1055
- 4) Steen et al. (2016), FABP4/aP2 regulates macrophage redox signaling and inflammasome activation via control of UCP2; Mol. Cell. Biol., Epub ahead of print, Oct. 17
- 5) Boss et al. (2015), FABP4 inhibition suppresses PPARgamma activity and VLDL-induced foam cell formation in IL-4-polarized human macrophages; Atherosclerosis, **240** 424

PHYSICAL DATA

Molecular Weight: 337.37

 $\begin{array}{ll} \text{Molecular Formula:} & C_{14}H_{11}NO_5S_2 \\ \text{Purity:} & 98\% \text{ by HPLC} \end{array}$

NMR: (Conforms)

Solubility: Soluble in DMSO (up to 50 mg/ml)

Physical Description: Yellow solid

Storage and Stability: Store as supplied, desiccated at room temperature 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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