

## Catalog # 10-1600 CaCCInh-A01

CAS# 407587-33-1

6-(1,1-Dimethylethyl)-2-[(2-furanylcarbonyl)amino]-4,5,6,7-tetrahydro-benzo[b]thiophene-3-carboxylic acid Lot # S103105



A potent inhibitor of TMEM16A ( $IC_{50} = 2.1 \mu M$ ), a calcium-activated chloride channel.<sup>1</sup> Blockade of TMEM16A with CaCCInh-A01 protects against renal fibrosis<sup>2</sup>, reduces blood brain barrier permeability, attenuates brain infarct size and neurological deficits after ischemic stroke<sup>3</sup>, and induces apoptosis and cell cycle arrest in various epithelium-originated cancer cells<sup>4</sup>.

- 1) Namkung et al. (2011) TMEM16A inhibitors reveal TMEM16A as a minor component of calcium-activated chloride channel conductance in airway and intestinal epithelial cells; J. Biol. Chem. **286** 2365
- Li et al. (2022) Blockade of TMEM16A protects against renal fibrosis by reducing intracellular Cl<sup>-</sup> concentration; Br. J. Pharmacol. **179** 3043
- 3) Liu et al. (2019) TMEM16A Inhibition Preserves Blood-Brain Barrier Integrity After Ischemic Stroke; Front. Cell Neurosci. **13** 360
- 4) Guan et al. (2016) Inhibition of calcium-activated chloride channel ANO1 suppresses proliferation and induces apoptosis of epithelium originated cancer cells; Oncotarget **7** 78619

## PHYSICAL DATA

Molecular Weight:	347.43
Molecular Formula:	C <sub>18</sub> H <sub>21</sub> NO <sub>4</sub> S
Purity:	>98% by TLC
	NMR: (Conforms)
Solubility:	DMSO (40 mg/ml)
Physical Description:	White solid
Storage and Stability:	Store as supplied 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 3 months.

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Focus Biomolecules LLC 400 Davis Drive, Suite 600 Plymouth Meeting PA 19462 www.focusbiomolecules.com