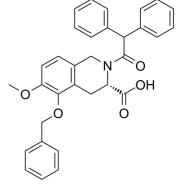


Data Sheet

Product Name: Olodanrigan
Cat. No.: CS-5781
CAS No.: 1316755-16-4
Molecular Formula: C32H29NO5
Molecular Weight: 507.58

Target: Angiotensin Receptor Pathway: GPCR/G Protein

Solubility: DMSO : \geq 34 mg/mL (66.98 mM); H2O : < 0.1 mg/mL (insoluble)



BIOLOGICAL ACTIVITY:

Olodanrigan (EMA401) is a highly selective, orally active, peripherally restricted **angiotensin II type 2 receptor (AT2R)** antagonist. It is under development as a neuropathic pain therapeutic agent. Olodanrigan (EMA401) analgesic action appears to involve inhibition of augmented AngII/AT2R induced p38 and p42/p44 MAPK activation, and hence inhibition of DRG neuron hyperexcitability and sprouting of DRG neurons^{[1][2][3][4]}. **In Vivo:** EMA401 (10 mg/kg; p.o.) results in a significant attenuation of theta power and increase in paw withdrawal latencies (PWL) in rats at day 14 after chronic constriction injury (CCI)^[4].

References:

- [1]. Rice AS et al. EMA401, an orally administered highly selective angiotensin II type 2 receptor antagonist, as a novel treatment for postherpetic neuralgia: a randomised, double-blind, placebo-controlled phase 2 clinical trial. Lancet. 2014 May 10;383(9929):1637-47.
- [2]. Anand U et al. Mechanisms underlying clinical efficacy of Angiotensin II type 2 receptor (AT2R) antagonist EMA401 in neuropathic pain: clinical tissue and in vitro studies. Mol Pain. 2015 Jun 26;11:38.
- [3]. Rice AS et al. EMA401, an orally administered highly selective angiotensin II type 2 receptor antagonist, as a novel treatment for postherpetic neuralgia: a randomised, double-blind, placebo-controlled phase 2 clinical trial. Lancet. 2014 May 10;383(9929):1637-47.
- [4]. Suguru Koyama, et al. An Electroencephalography Bioassay for Preclinical Testing of Analgesic Efficacy.

CAIndexNames:

3-Isoquinolinecarboxylic acid, 2-(2,2-diphenylacetyl)-1,2,3,4-tetrahydro-6-methoxy-5-(phenylmethoxy)-, (3S)-

SMILES:

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Caution: Product has not been fully validated for medical applications. For research use only.

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