

Data Sheet

Product Name: Parecoxib

Cat. No.: CS-1959

CAS No.: 198470-84-7

Molecular Formula: C19H18N2O4S

Molecular Weight: 370.42 Target: COX

Pathway: Immunology/Inflammation

Solubility: DMSO : \geq 50 mg/mL (134.98 mM)

BIOLOGICAL ACTIVITY:

Parecoxib is a potent and selective COX-2 inhibitor. IC50 value: Target: COX-2 in vitro: The prodrug Parecoxib as well as its active metabolite val have a specific affinity to the cannabinoid (CB) receptor measured in CB1-expressing HEK 293 cells and rat brain tissue [1]. in vivo: Adult male Sprague-Dawley rats were administered parecoxib (10 or 30 mg kg(-1), IP) or isotonic saline twice a day starting 24 h after middle cerebral artery occlusion (MCAO) for three consecutive days [2]. The selective COX-2 inhibitor parecoxib was delivered 20 min before or 20 min after the incision by intraperitoneal injection. Pretreatment with parecoxib markedly attenuated the pain hypersensitivity induced by incision [3].

References:

[1]. Ye Z, et al. Delayed administration of parecoxib, a specific COX-2 inhibitor, attenuated postischemic neuronal apoptosis by phosphorylation Akt and GSK-3β. Neurochem Res. 2012 Feb;37(2):321-9.

[2]. Schr?der H, et al. Parecoxib and its metabolite valdecoxib directly interact with cannabinoid binding sites in CB1-expressing HEK 293 cells and rat brain tissue. Neurochem Int. 2011 Jan;58(1):9-13.

[3]. Guo YJ, et al. Analgesic effects of the COX-2 inhibitor parecoxib on surgical pain through suppression of spinal ERK signaling. Exp Ther Med. 2013 Jul;6(1):275-279.

CAIndexNames:

Propanamide, N-[[4-(5-methyl-3-phenyl-4-isoxazolyl)phenyl]sulfonyl]-

SMILES:

 $\mathsf{CCC}(\mathsf{NS}(=\mathsf{O})(\mathsf{C1} = \mathsf{CC} = \mathsf{C}(\mathsf{C2} = \mathsf{C}(\mathsf{C}) \mathsf{ON} = \mathsf{C2C3} = \mathsf{CC} = \mathsf{C3})\mathsf{C} = \mathsf{C1}) = \mathsf{O}) = \mathsf{O}$

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

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