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**Product Data Sheet**


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Product Name: Ampiroxicam

Cat. No.: GC14104

**Chemical Properties**

Cas. No. 99464-64-9

Chemical Name ethyl 1-[[2-methyl-1,1-dioxo-3-(pyridin-2-ylcarbamoylethyl]oxy]ethyl carbonate

SMILES CCOC(=O)OC(C)OC1=C(N(S(=O)(=O)C2=CC=CC=C21)C)C(=O)NC3=CC=CC=N3Formula C<sub>20</sub>H<sub>21</sub>N<sub>3</sub>O<sub>7</sub>S

M.Wt 447.46

Solubility DMF: 20 mg/ml, DMF:PBS (pH 7.2)(1:3): 0.25 mg/ml, DMSO: 10 mg/ml

Storage Store at -20°C

General tips For obtaining a higher solubility, please warm the tube at 37 °C and shake it in the ultrasonic bath for a while. Stock solution can be stored below -20°C for several months.

Shipping Condition Evaluation sample solution : ship with blue ice All other available size: ship with RT, or blue ice upon request.

Structure **Background**

Ampiroxicam (CP65703) is a nonselective cyclooxygenase inhibitor used as anti-inflammatory drug. Target: COX. Ampiroxicam is a non-steroidal anti-inflammatory drug. It is a prodrug of piroxicam. Ampiroxicam inhibits the stretching response in mice induced by phenylbenzoquinone (PBQ) with maximum protective effect (MPE) of 2 mg/kg. Ampiroxicam inhibits swelling in a dose-responsive manner in the rat foot edema (RFE) assay with ED<sub>50</sub> of 28 mg/kg at single oral dose and 7.8 mg/kg at 5 daily oral dose. Ampiroxicam blocks primary and secondary lesion development in rat adjuvant arthritis with ED<sub>50</sub> of 2.2 mg/kg and 0.5 mg/kg, respectively. Ampiroxicam (3.2 mg/kg) leads to a plasma concentration of 12 µg/mL at a T<sub>max</sub> of 2 hours for piroxicam derived from ampiroxicam in rats [1]. Ultraviolet-A (UVA)-irradiated 1% Ampiroxicam sensitized in guinea pigs shows positive reaction in the patch testing to UVA-irradiated 1% Ampiroxicam and 1% thiosalicylate (TOS). Concentration of Ampiroxicam is easily

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reduced by the increase in UVA irradiation doses, as compared with that of piroxicam [2].

### References:

- [1]. Aoki T, et al. Premedication with cyclooxygenase-2 inhibitor meloxicam reduced postoperative pain in patients after oral surgery. *Int J Oral Maxillofac Surg.* 2006 Jul;35(7):613-7.
- [2]. Sasaki, T., et al., Antigenic characterization in ampiroxicam-induced photosensitivity using an in vivo model of contact hypersensitivity. *J Dermatol Sci*, 1999. 21(3): p. 170-5.

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