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**Product Data Sheet**

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Product Name: Polmacoxib

Cat. No.: GC18271

**Chemical Properties**

Cas. No. 301692-76-2

Chemical Name 4-[3-(3-fluorophenyl)-4,5-dihydro-5,5-dimethyl-4-oxo-2-furanyl]-benzenesulfonamide

SMILES FC1=CC(C2=C(C3=CC=C(S(N)(=O)=O)C=C3)OC(C)(C)C2=O)=CC=C1Formula C<sub>18</sub>H<sub>16</sub>FNO<sub>4</sub>S

M.Wt 361.4

Solubility DMF: 20 mg/ml, DMSO: 20 mg/ml, DMSO:PBS (pH 7.2)(1:8): 0.5 mg/ml, Ethanol: 5 mg/ml  
Store Storage 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 sizes: ship with RT, or blue ice upon request.

Structure **Background**

Polmacoxib is an inhibitor of cyclooxygenase 2 (COX-2) and the carbonic anhydrase subtypes I (CAI) and CAII. It inhibits COX-2 in the absence of carbonic anhydrase II with an IC<sub>50</sub> value of 40 nM, which increases by approximately 4- and 17-fold in the presence of a CAII at a molar ratio of 1:1 and 1:5, respectively. It also inhibits CAI and CAII (IC<sub>50</sub>s = 210 and 95 nM, respectively). Polmacoxib prevents >95 and 90% of prostaglandin E<sub>2</sub> (PGE<sub>2</sub>) production in HCA-7 and HT-29 human colon cancer cells, respectively, using concentrations of 0.01 and 0.001 µg/ml. It inhibits polyp formation in a transgenic mouse model of intestinal polyp formation and tumor growth in human colorectal carcinoma mouse xenograft models when used at a dose of 7 mg/kg. The inhibition of COX-2 and CAII by polmacoxib has the potential for fewer serious systemic adverse effects, including cardiovascular events associated with COX-2 selective inhibitors such as celecoxib. Formulations containing polmacoxib have been used in the treatment of

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

Tel: (909) 407-4943 Fax: (626) 353-8530 E-mail: tech@glpbio.com

Address: 10292 Central Ave. #205, Montclair, CA, USA

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osteoarthritis.

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