# PRODUCT INFORMATION



# PF-04418948

Item No. 15016

CAS Registry No.: 1078166-57-0

Formal Name: 1-(4-fluorobenzoyl)-3-[[(6-

methoxy-2-naphthalenyl)oxy]

methyl]-3-azetidinecarboxylic acid

MF:  $C_{23}H_{20}FNO_5$ 

409.4 FW: **Purity:** ≥98%

UV/Vis.:  $\lambda_{max}$ : 230 nm Supplied as: A crystalline solid

Storage: -20°C Stability: ≥4 years

Information represents the product specifications. Batch specific analytical results are provided on each certificate of analysis.

## **Laboratory Procedures**

PF-04418948 is supplied as a crystalline solid. A stock solution may be made by dissolving the PF-04418948 in the solvent of choice, which should be purged with an inert gas. PF-04418948 is soluble in organic solvents such as ethanol, DMSO, and dimethyl formamide (DMF). The solubility of PF-04418948 in these solvents is approximately 0.5, 16, and 20 mg/ml, respectively.

PF-04418948 is sparingly soluble in aqueous buffers. For maximum solubility in aqueous buffers, PF-04418948 should first be dissolved in DMF and then diluted with the aqueous buffer of choice. PF-04418948 has a solubility of approximately 0.5 mg/ml in a 1:1 solution of DMF:PBS (pH 7.2) using this method. We do not recommend storing the aqueous solution for more than one day.

### Description

Prostaglandin E<sub>2</sub> (PGE<sub>2</sub>) evokes distinct responses through four different 'E prostanoid' (EP) receptors. EP<sub>2</sub> is a G protein-coupled receptor which signals preferentially through Gs proteins. PF-04418948 is an orally active, potent, and selective  $EP_2$  receptor antagonist ( $IC_{50} = 16$  nM).<sup>1</sup> It is over a thousand-fold less active at other prostanoid receptors, including other EP receptors. PF-04418948 can be used in cells and tissues.<sup>2,3</sup> In rats, it significantly reduces the effects of the EP2-selective agonist butaprost on cutaneous blood flow when given orally.1

#### References

- 1. af Forselles, K.J., Root, J., Clarke, T., et al. In vitro and in vivo characterization of PF-04418948, a novel, potent and selective prostaglandin EP2 receptor antagonist. Br. J. Pharmacol. 164(7), 1847-1856 (2011).
- 2. Birrell, M.A., Maher, S.A., Buckley, J., et al. Selectivity profiling of the novel EP2 receptor antagonist, PF-04418948, in functional bioassay systems: Atypical affinity at the guinea pig EP<sub>2</sub> receptor. Br. J. Pharmacol. 168(1), 129-138 (2013).
- 3. Säfholm, J., Dahlén, S.E., Delin, I., et al. PGE2 maintains the tone of the guinea pig trachea through a balance between activation of contractile  $EP_1$  receptors and relaxant  $EP_2$  receptors. Br. J. Pharmacol. 168(4), 794-806 (2013).

WARNING
THIS PRODUCT IS FOR RESEARCH ONLY - NOT FOR HUMAN OR VETERINARY DIAGNOSTIC OR THERAPEUTIC USE.

This material should be considered hazardous until further information becomes available. Do not ingest, inhale, get in eyes, on skin, or on clothing. Wash thoroughly after handling. Before use, the user must review the complete Safety Data Sheet, which has been sent via email to your institution.

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