

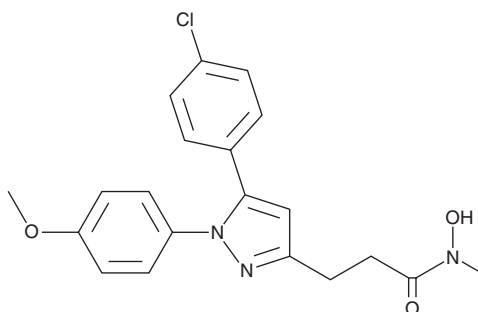
# PRODUCT INFORMATION



## Tepoxalin

Item No. 34849

**CAS Registry No.:** 103475-41-8  
**Formal Name:** 5-(4-chlorophenyl)-N-hydroxy-1-(4-methoxyphenyl)-N-methyl-1H-pyrazole-3-propanamide  
**Synonyms:** ORF 20485, RWJ 20485  
**MF:** C<sub>20</sub>H<sub>20</sub>ClN<sub>3</sub>O<sub>3</sub>  
**FW:** 385.8  
**Purity:** ≥98%  
**UV/Vis.:** λ<sub>max</sub>: 237, 256 nm  
**Supplied as:** A 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

Tepoxalin is supplied as a solid. A stock solution may be made by dissolving the tepoxalin in the solvent of choice, which should be purged with an inert gas. Tepoxalin is soluble in organic solvents such as ethanol, DMSO, and dimethyl formamide (DMF). The solubility of tepoxalin in ethanol is approximately 15 mg/ml and approximately 30 mg/ml in DMSO and DMF.

### Description

Tepoxalin is a dual inhibitor of COX and 5-lipoxygenase (5-LO; IC<sub>50</sub>s = 2.84 and 0.15 μM, respectively, in RBL-1 cell lysates).<sup>1</sup> It also inhibits the activity of 12-, but not 15-, LO (IC<sub>50</sub>s = 4.45 and 146.6 μM for the human platelet and rabbit reticulocyte enzymes, respectively). Tepoxalin reduces the production of thromboxane B<sub>2</sub> (TXB<sub>2</sub>; Item No. 19030) and leukotriene B<sub>4</sub> (LTB<sub>4</sub>; Item No. 20110) induced by the calcium ionophore A23187 (Item Nos. 11016 | 22030) in human peripheral blood mononuclear cells (PBMCs; IC<sub>50</sub>s = 0.01 and 0.07 μM, respectively) and decreases epinephrine-induced platelet aggregation in human platelet-rich plasma (PRP; IC<sub>50</sub> = 0.045 μM). It inhibits paw swelling in a rat model of adjuvant-induced arthritis with an ED<sub>50</sub> value of 3.5 mg/kg.

### Reference

1. Argentieri, D.C., Ritchie, D.M., Ferro, M.P., *et al.* Tepoxalin: A dual cyclooxygenase/5-lipoxygenase inhibitor of arachidonic acid metabolism with potent anti-inflammatory activity and a favorable gastrointestinal profile. *J. Pharmacol. Exp. Ther.* **271**(3), 1399-1408 (1994).

#### WARNING

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

#### SAFETY DATA

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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