

Product Name: SC 51089

Catalog No.: 3758

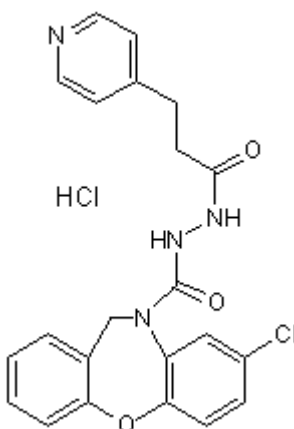
Batch No.: 1

CAS Number: 146033-02-5

IUPAC Name: 8-Chlorodibenz(Z)[b,f][1,4]oxazepine-10(11H)-carboxylic acid 2-[1-oxo-3-(4-pyridinyl)propyl]hydrazide hydrochloride

1. PHYSICAL AND CHEMICAL PROPERTIES

Batch Molecular Formula: C₂₂H₁₉ClN₄O₃.HCl.H₂O
Batch Molecular Weight: 477.35
Physical Appearance: White solid
Solubility: DMSO to 100 mM
Storage: Store at -20°C
Batch Molecular Structure:



2. ANALYTICAL DATA

HPLC: Shows 100% purity
¹H NMR: Consistent with structure
Mass Spectrum: Consistent with structure

Microanalysis:

	Carbon	Hydrogen	Nitrogen
Theoretical	55.36	4.65	11.74
Found	55.09	4.42	11.66

Caution - Not Fully Tested • Research Use Only • Not For Human or Veterinary Use

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Description:

Selective EP₁ prostanoid receptor antagonist (K_i values are 1.3, 11.2, 17.5, 61.1, > 100, > 100, > 100, >100 and > 100 μM for EP₁, TP, EP₃, EP₂, EP₄, FP and DP receptors respectively). Attenuates PGE₂-induced neuronal cell death in vitro and slows tumor growth in vivo.

Physical and Chemical Properties:

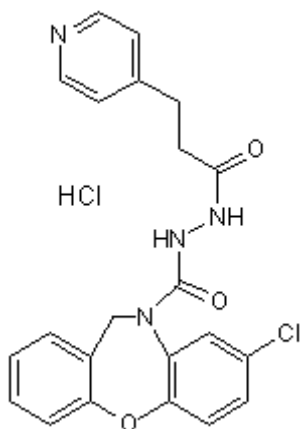
Batch Molecular Formula: C₂₂H₁₉ClN₄O₃·HCl·H₂O

Batch Molecular Weight: 477.35

Physical Appearance: White solid

Minimum Purity: >99%

Batch Molecular Structure:



Storage: Store at -20°C

Solubility & Usage Info:

DMSO to 100 mM

Stability and Solubility Advice:

Some solutions can be difficult to obtain and can be encouraged by rapid stirring, sonication or gentle warming (in a 45-60°C water bath).

Information concerning product stability, particularly in solution, has rarely been reported and in most cases we can only offer a general guide. Our standard recommendations are:

SOLIDS: Provided storage is as stated on the product label and the vial is kept tightly sealed, the product can be stored for up to 6 months from date of receipt.

SOLUTIONS: We recommend that stock solutions, once prepared, are stored aliquoted in tightly sealed vials at -20°C or below and used within 1 month. Wherever possible solutions should be made up and used on the same day.

References:

Abramovitz et al (2000) The utilization of recombinant prostanoid receptors determine the affinities and selectivities of prostaglandins and related analogs. *Biochim.Biophys.Acta* **1483** 285. PMID: 10634944.

Matsuo et al (2004) Inhibition of human glioma cell growth by a PHS-2 inhibitor, NS398, and a prostaglandin E receptor subtype EP₁-selective antagonist. *J.Neurochem.* **66** 285.

Saleem et al (2007) Effect of EP₁ receptor on cerebral blood flow in the middle cerebral artery occlusion model of stroke in mice. *J.Neurosci.Res.* **85** 2433. PMID: 17600836.

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bio-techne.com

info@bio-techne.com

techsupport@bio-techne.com

North America

Tel: (800) 343 7475

China

info.cn@bio-techne.com

Tel: +86 (21) 52380373

Europe Middle East Africa

Tel: +44 (0)1235 529449

Rest of World

www.tocris.com/distributors

Tel: +1 612 379 2956