

Product Name: VU 0463271

Catalog No.: 4719

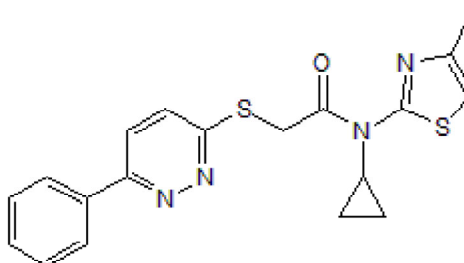
Batch No.: 2

CAS Number: 1391737-01-1

IUPAC Name: *N*-Cyclopropyl-*N*-(4-methyl-2-thiazolyl)-2-[(6-phenyl-3-pyridazinyl)thio]acetamide

1. PHYSICAL AND CHEMICAL PROPERTIES

Batch Molecular Formula: C₁₉H₁₈N₄OS₂
Batch Molecular Weight: 382.5
Physical Appearance: Pale yellow solid
Solubility: DMSO to 50 mM
Storage: Store at +4°C
Batch Molecular Structure:



2. ANALYTICAL DATA

HPLC: Shows 99.6% purity
¹H NMR: Consistent with structure
Mass Spectrum: Consistent with structure

Microanalysis:

	Carbon	Hydrogen	Nitrogen
Theoretical	59.66	4.74	14.65
Found	59.63	4.7	14.65

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

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IUPAC Name: *N*-Cyclopropyl-*N*-(4-methyl-2-thiazolyl)-2-[(6-phenyl-3-pyridazinyl)thio]acetamide

Description:

VU 0463271 is a potent and selective inhibitor of the neuronal K-Cl cotransporter, KCC2 (IC₅₀ = 61 nM); displays >100-fold selectivity versus the Na-K-2Cl cotransporter 1 (NKCC1) and no activity against a panel of 68 GPCRs, ion channels and transporters.

Physical and Chemical Properties:

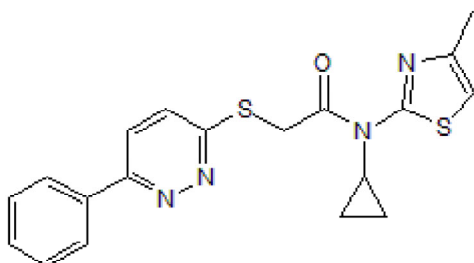
Batch Molecular Formula: C₁₉H₁₈N₄OS₂

Batch Molecular Weight: 382.5

Physical Appearance: Pale yellow solid

Minimum Purity: ≥98%

Batch Molecular Structure:



Storage: Store at +4°C

Solubility & Usage Info:

DMSO to 50 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. *Unless contradicted by product-specific protocols or instructions, our standard recommendations apply:

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:

Sivakumaran et al (2015) Selective inhibition of KCC2 leads to hyperexcitability and epileptiform discharges in hippocampal slices and *in vivo*. *J.Neurosci.* **35** 8291. PMID: 26019342.

Delpire et al (2012) Further optimization of the K-Cl cotransporter KCC2 antagonist ML077: development of a highly selective and more potent *in vitro* probe. *Bioorg.Med.Chem.Lett.* **22** 4532. PMID: 22727639.

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