# SB239063

p38 inhibitor

A potent, cell permeable inhibitor of p38 MAP kinase ( $IC_{50}$ =44nM for recombinant purified human p38 $\alpha$ ). Inhibits IL-1 and TNF- $\alpha$  production in LPS-stimulated human peripheral blood monocytes ( $IC_{50}$ =120nM and 350nM, respectively). Reported to protect against mild excitotoxic neuronal injury caused by NMDA and provides substantial protection against cell death induced by either oxygen glucose deprivation (OGD) or magnesium deprivation in cultured neurons. Also, reduces myocardial reperfusion injury via inhibition of endothelial adhesion molecule expression and blockade of polymorphonuclear (PMN) accumulation.

Citations: 9

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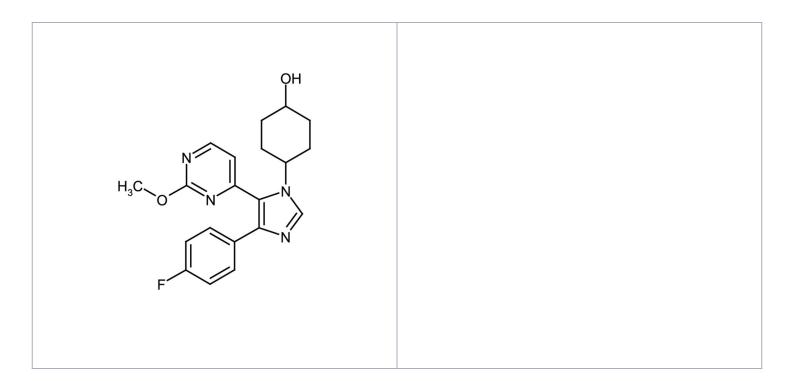
## **Ordering Information**

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ALX-270-351-M001	1mg
ALX-270-351-M005	5mg
ALX-270-351-C500	500µg

Manuals, SDS & CofA

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#### **Handling & Storage**

**Use/Stability** As indicated on product label or CoA when stored as recommended.

**Handling** Protect from light. Packaged under inert gas.

Long Term Storage -20°C

**Shipping** Ambient Temperature

### Regulatory Status RUO - Research Use Only

#### **Product Details**

Alternative Name trans-1-(4-Hydroxycyclohexyl)-4-(fluorophenyl)-5-(2-methoxypyrimidin-4-yl) imidazole

**Appearance** White to off-white solid.

**CAS** 193551-21-2

Couple Target p38

Couple Type Inhibitor

Formula  $C_{20}H_{21}FN_4O_2$ 

**Identity** Identity determined by 1H-NMR.

**MW** 368.4

Purity ≥97% (HPLC)

**Solubility** Soluble in DMSO, 100% ethanol or dichlormethane (may require slight warming).