

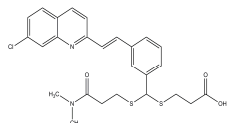
Product: MK-571

ALTERNATE NAME: (E)-3-[[[3-[2-(7-chloro-2-quinolinyl)ethenyl]phenyl]][[3-(dimethylamino)-3-oxopropyl]thio]methyl]thio]-propanoic acid; L-660,711

CATALOG #: 2691-5, 25

AMOUNT: 5 mg, 25 mg

STRUCTURE:



MOLECULAR FORMULA: C₂₆H₂₇ClN₂O₃S₂

MOLECULAR WEIGHT: 515.09

CAS NUMBER: 115104-28-4

APPEARANCE: Yellow solid

SOLUBILITY: DMSO (~40 mg/ml)

PURITY: ≥95%

STORAGE: Store at -20 °C. Protect from air and light

DESCRIPTION: MK 571 is a selective, CysLT₁ receptor antagonist. It blocks the binding of leukotriene LTD₄, but not LTC₄, to human and guinea pig lung membranes with K_i values of 0.22 nM and 2.1 nM, respectively. MK 571 effectively blocks LTD₄ activation of recombinant human and murine CysLT₁ receptors but is ineffective at blocking LTC₄ or LTD₄ activation of the recombinant human or murine CysLT₂ receptors. MK-57 also acts as a specific inhibitor of multi-drug resistance protein-1 (MRP1).

REFERENCES: Vellenga, E., *et al.* (1999). *Br. J. Pharmacol.* **127**, 441-448.

HANDLING: Do not take internally. Wear gloves and mask when handling the product! Avoid contact by all modes of exposure.

RELATED PRODUCTS:

MK-571 (Cat. No. 2691-5, 25)

MK-571, sodium salt (Cat. No. 2692-5, 25)

NSC-23925 (Cat. No. 2686-5,25)

FOR RESEARCH USE ONLY! Not to be used on humans.