

## H 89 dihydrochloride

Catalogue Number : 1303954

RUO: For Research Use Only. Not for use in diagnostic procedures.

### Product Information

**Synonyms:** H-89 dihydrochloride hydrate

**Chemical Name:** N-[2-[(E)-3-(4-bromophenyl)prop-2-enyl]amino]ethyl]isoquinoline-5-sulfonamide;hydrate;dihydrochloride

**Molecular Formula:** C<sub>20</sub>H<sub>20</sub>BrN<sub>3</sub>O<sub>2</sub>S 2HCl

**Molecular Weight:** 519.3

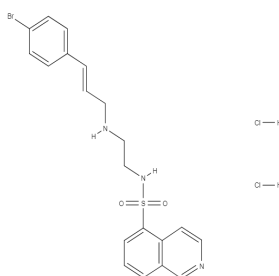
**CAS Number:** 130964-39-5

**Purity:** ≥98%

**Applications:** FA

**Formulation:** Crystalline solid

**Storage:** Product should be kept at -20°C.



### Description

H 89 is a potent PKA inhibitor and an inhibitor for other protein kinases including S6K1, MSK1, ROCKII, PCKalpha, and MAPKAP-K1b. It has been shown to decrease morphine withdrawal symptoms in mice and to inhibit the forskolin-induced neurite outgrowth of PC12D pheochromocytoma cells.

### Preparation & Storage

Soluble in organic solvents such as DMF and DMSO. DMSO up to 25mg/ml.

### References

1. Chijiwa, T., Mishima, A., Hagiwara, M., Sano, M., Hayashi, K., Inoue, T., ... Hidaka, H. (1990). Inhibition of forskolin-induced neurite outgrowth and protein phosphorylation by a newly synthesized selective inhibitor of cyclic AMP-dependent protein kinase, N-[2-(p-bromocinnamylamino)ethyl]-5-isoquinolinesulfonamide (H-89), of PC12D pheochromocytoma cells. *Journal of Biological Chemistry*, 265(9), 5267-5272.
2. Hidaka, H., Kobayashi, R. (1992). Pharmacology of protein kinase inhibitors. *Annual review of pharmacology and toxicology*, 32(1), 377-397.
3. Lochner, A., Moolman, J. A. (2006). The many faces of H89: a review. *Cardiovascular Therapeutics*, 24(34), 261-274.