

# Chelerythrine Chloride

✓ 5 mg

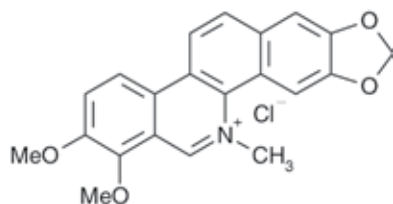
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For Research Use Only. Not For Use In Diagnostic Procedures.

**Background:** Chelerythrine is a benzophenanthridine alkaloid that elicits a wide range of biological responses. It was initially reported to be a potent PKC inhibitor ( $IC_{50} = 660$  nM) through interaction with the catalytic domain, competitive with respect to the phosphate acceptor, noncompetitive with respect to ATP (1), and is often used in research as a means to inhibit PKC activation (2-4). However, subsequent studies have shown that chelerythrine is not a potent PKC inhibitor and its effects are independent of PKC inhibition (5,6). Researchers have demonstrated that chelerythrine activates JNK and p38 through an oxidative stress mechanism (6), inhibits Bcl-xL by preventing Bcl-xL-Bak BH3 peptide binding ( $IC_{50} = 1.5$   $\mu$ M) and disrupting the interaction between Bcl-xL and Bax (7), and induces apoptosis (6-8).

**Molecular Formula:** C<sub>21</sub>H<sub>18</sub>NO<sub>4</sub>Cl

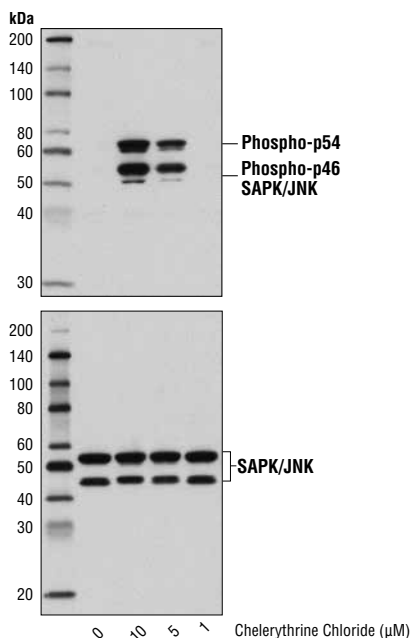


**Molecular Weight:** 383.82 g/mol

**Solubility:** Soluble in DMSO at 5 mg/ml; very poorly soluble in ethanol and water with maximum solubility in water ~10-50  $\mu$ M.

**Purity:** >99%

**Directions for Use:** Chelerythrine Chloride is supplied as a lyophilized powder. For a 10 mM stock, reconstitute the 5 mg in 1.30 ml DMSO. Working concentrations and length of treatment can vary depending on the desired effect, but it is typically used at 1-20  $\mu$ M either as a pretreatment for 15 min-1 hr prior to treating with a stimulator or used alone with varying treatment times lasting up to 24 hr.



Western blot analysis of extracts from HeLa cells, serum-starved overnight and untreated or treated with Chelerythrine Chloride (2 hr) at the indicated concentrations, using Phospho-SAPK/JNK (Thr183/Tyr185) (81E11) Rabbit mAb #4668 (upper) or SAPK/JNK (56G8) Rabbit mAb #9258 (lower).

**Storage:** Store lyophilized or in solution at -20°C, desiccated. Protect from light. In lyophilized form, the chemical is stable for 24 months. Once in solution, use within 3 months to prevent loss of potency. Aliquot to avoid multiple freeze/thaw cycles.

**Background References:**

(1) Herbert, J.M. et al. (1990) *Biochem Biophys Res Commun* 172, 993-9.  
(2) Chao, M.D. et al. (1998) *Planta Med* 64, 662-3.  
(3) Nakai, M. et al. (1999) *J Neurochem* 72, 1179-86.  
(4) Iwabu, A. et al. (2004) *J Biol Chem* 279, 14551-60.  
(5) Lee, S.K. et al. (1998) *J Biol Chem* 273, 19829-33.  
(6) Yu, R. et al. (2000) *J Biol Chem* 275, 9612-9.  
(7) Chan, S.L. et al. (2003) *J Biol Chem* 278, 20453-6.  
(8) Chmura, S.J. et al. (2000) *Clin Cancer Res* 6, 737-42.