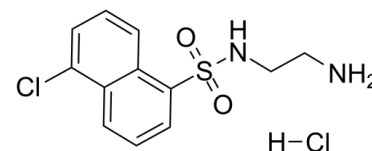


## Data Sheet

<b>Product Name:</b>	A-3 (hydrochloride)
<b>Cat. No.:</b>	CS-0103566
<b>CAS No.:</b>	78957-85-4
<b>Molecular Formula:</b>	C <sub>12</sub> H <sub>14</sub> Cl <sub>2</sub> N <sub>2</sub> O <sub>2</sub> S
<b>Molecular Weight:</b>	321.22
<b>Target:</b>	CaMK; Casein Kinase; PKA; PKC
<b>Pathway:</b>	Cell Cycle/DNA Damage; Epigenetics; Neuronal Signaling; Protein Tyrosine Kinase/RTK; Stem Cell/Wnt; TGF-beta/Smad
<b>Solubility:</b>	DMSO : 125 mg/mL (389.14 mM; Need ultrasonic)



### BIOLOGICAL ACTIVITY:

A-3 hydrochloride is a potent, cell-permeable, reversible, ATP-competitive non-selective antagonist of various **kinases**. It against PKA ( $K_i=4.3 \mu\text{M}$ ), casein kinase II ( $K_i=5.1 \mu\text{M}$ ) and myosin light chain kinase (MLCK) ( $K_i=7.4 \mu\text{M}$ ). A-3 hydrochloride also inhibits PKC and casein kinase I with  $K_i$  values of  $47 \mu\text{M}$  and  $80 \mu\text{M}$ , respectively<sup>[1]</sup>. **In Vitro**: A-3 hydrochloride inhibits MLC-kinase competitively with respect to ATP and that the  $K_i$  value is  $7.4 \mu\text{M}$ . A-3 is also a competitive inhibitor of cAMP-dependent protein kinase, cGMP-dependent protein kinase, protein kinase C, casein kinase I, and casein kinase II, with respect to ATP, exhibits  $K_i$  values of  $4.3 \mu\text{M}$ ,  $3.8 \mu\text{M}$ ,  $47 \mu\text{M}$ ,  $80 \mu\text{M}$ , and  $5.1 \mu\text{M}$ , respectively<sup>[1]</sup>.

### References:

[1]. Inagaki M, et al. Naphthalenesulfonamides as calmodulin antagonists and protein kinase inhibitors. Mol Pharmacol. 1986 Jun;29(6):577-81.

### CAIndexNames:

1-Naphthalenesulfonamide, N-(2-aminoethyl)-5-chloro-, hydrochloride (1:1)

### SMILES:

O=S(C1=C2C=CC=C(Cl)C2=CC=C1)(NCCN)=O.[H]Cl

**Caution: Product has not been fully validated for medical applications. For research use only.**

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