



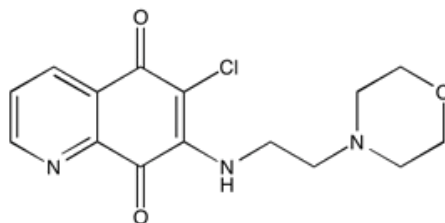
**Catalog # 10-1352**

**NSC-663284**

383907-43-5

6-Chloro-7-[[2-(4-morpholinyl)ethyl]amino]-5,8-quinolinedione

Lot# FBA6029



Potent, selective and irreversible inhibitor of CDC25 phosphatases,  $K_i=29, 95$  and  $89$  nM for human CDC25A, B<sub>2</sub> and C respectively and  $>20$ - and  $>450$ -fold selective over VHR and PTP1B.<sup>1</sup> Arrests cells at G<sub>1</sub> and G<sub>2</sub>/M and inhibits cdk1 and 2 activation.<sup>2</sup> Inhibits the growth of a variety of human tumor cell lines.<sup>3</sup> Acts via irreversible oxidation of the catalytic cysteine of CDC25.<sup>4</sup>

- 1) Lazo *et al.* (2001), *Discovery and biological evaluation of a new family of potent inhibitors of the dual specificity protein phosphatases Cdc25*; J. Med. Chem., **44** 4042
- 2) Pu *et al.* (2002), *Dual G1 and G2 phase inhibition by a novel, selective Cdc25 inhibitor 6-chloro-7-(2-morpholin-4-ylethylamino)-quinoline-3,8-dione*; J. Biol. Chem., **277** 46877
- 3) Han *et al.* (2004), *NAD(P)H:quinone oxidoreductase-1-dependent and -independent cytotoxicity of potent quinone Cdc25 phosphatase inhibitors*; J. Pharmacol. Exp. Ther., **309** 64
- 4) Brisson *et al.* (2005), *Redox regulation of Cdc25B by cell-active quinolonediones*; Mol. Pharmacol., **68** 1810

#### **PHYSICAL DATA**

Molecular Weight:	321.77
Molecular Formula:	C <sub>15</sub> H <sub>16</sub> ClN <sub>3</sub> O <sub>3</sub>
Purity:	>98% by TLC
	NMR: (Conforms)
Solubility:	DMSO (up to 15 mg/ml), Ethanol (up to 55 mg/ml)
Physical Description:	Dark Red solid
Storage and Stability:	Store as supplied desiccated at -20°C for up to 2 years from the date of purchase. Solutions in DMSO or ethanol may be stored at -20°C for up to 1 month.

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