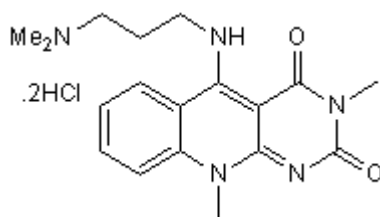


**MATERIAL DATA SHEET****HLI 373****Cat. # I-400**

Inhibitor of Hdm2 ubiquitin ligase (E3). Blocks Hdm2-mediated ubiquitylation and proteasomal degradation of p53; activates p53-dependent transcription. Induces apoptosis in several tumor cell lines that express wild-type p53 such as LOX-IMVI, A549, HT1080 and U2OS.

**Product Information****Quantity:** X mg**Formula:**  $C_{18}H_{23}N_5O_2HCl$  **FW:** 377.87**Structure:****Physical/Chemical Characteristics****Solubility:** Soluble to 100 mM in water and to 10 mM in DMSO**Purity:** > 99%**Use & Storage**

**Use:** Add from DMSO stock directly to in vitro or in vivo assays at desired concentration. Pre-incubation with compound prior to assay is recommended for maximal effect.

**Storage:** Store stock solution at -20°C. Avoid multiple freeze/thaw cycles.

**Literature**

**References:** Kitagaki, *et al* (2008) Mol.Cancer Ther. 7: 2445  
Yang, *et al* (2009) Cancer Sci. 100: 24

*For Laboratory Research Use Only, Not For Use in Humans*