



Ganetespib

Catalog No: tcsc0697

Available Sizes
Size: 5mg
Size: 10mg
Size: 50mg
Size: 100mg
Size: 200mg
Size: 500mg
Specifications
CAS No: 888216-25-9
Formula: $C_{20}^{H}_{20}^{N}_{4}^{O}_{3}$
Pathway: Metabolic Enzyme/Protease;Cell Cycle/DNA Damage
Target: HSP;HSP
Purity / Grade: >98%
Solubility: DMSO : ≥ 32 mg/mL (87.82 mM)
Alternative Names: STA-9090





Observed Molecular Weight:

364.4

Product Description

Ganetespib is a unique non-geldanamycin heat shock protein 90 (HSP90) inhibitor, with antitumor activity.

IC50 & Target: HSP90^[1]

In Vitro: Ganetespib causes depletion of receptor tyrosine kinases, extinguishing of downstream signaling, inhibition of proliferation and induction of apoptosis with IC₅₀ values ranging 2-30 nM in genomically-defined NSCLC cell lines. Ganetespib is also approximately 20-fold more potent in isogenic Ba/F3 pro-B cells rendered IL-3 independent by expression of EGFR and ERBB2 mutants^[1]. Ganetespib exhibits potent in vitro cytotoxicity in a range of solid and hematologic tumor cell lines, induces the degradation of known Hsp90 client proteins, displays superior potency to the ansamycin inhibitor 17-allylamino-17-demethoxygeldanamycin (17-AAG)^[2]. Ganetespib is a potent HSP90 inhibitor, and shown to kill canine tumor cell lines in vitro^[3]. Ganetespib possesses superior JAK/STAT inhibitory activity to both P6 and 17-AAG in terms of potency or duration of response in the HEL92.1.7 cells^[4].

In Vivo: Ganetespib (125 mg/kg, i.v.) accumulates in tumors relative to normal tissues and displays greater in vivo efficacy than 17-AAG without increased toxicity and inhibits proliferation and induces apoptosis in parallel with EGFR depletion in NCI-H1975 xenografts^[1]. Ganetespib (100, 125, 150 mg/kg, i.v.) shows potent antitumor efficacy in solid and hematologic xenograft models of oncogene addiction, as evidenced by significant growth inhibition and/or regressions^[2].

All products are for RESEARCH USE ONLY. Not for diagnostic & therapeutic purposes!