



Galeterone

Catalog No: tcsc0334

Available Sizes
Size: 5mg
Size: 10mg
Size: 50mg
Size: 100mg
Size: 200mg
Size: 500mg
Specifications
CAS No: 851983-85-2
Formula: C ₂₆ H ₃₂ N ₂ O
Pathway: Metabolic Enzyme/Protease
Target: Cytochrome P450
Purity / Grade: >98%
Solubility: DMSO: 18 mg/mL (46.33 mM; Need ultrasonic and warming)
Alternative Names: TOK-001;VN-124-1





Observed Molecular Weight:

388.55

Product Description

TOK-001 is a multifunctional antiandrogen and CYP17 inhibitor (IC_{50} =47 nM) in castration resistant prostate cancer (CRPC).

IC50 & Target: IC50: 47 nM (CYP17)^[1]

In Vitro: TOK-001 affords strong CYP17 lyase inhibition, with IC_{50} of 47 $nM^{[1]}$. TOK-001 is both a CYP17A1 inhibitor and androgen receptor antagonist and the similarity of these binding modes is likely the reason for this dual mechanism of action. This CYP17A1 binds abiraterone and TOK-001 with absorbance decreases at 402 nm and increases at 424 nm, consistent with nitrogen binding to the heme iron (type II interaction) with K_d of [2]. When LNCaP cells are cultured in medium supplemented with charcoal-stripped serum (CSS, T[3].

In Vivo: Mice inoculated with LAPC-4 tumors are treated subcutaneously with 0.15 mmol/kg of TOK-001 twice daily. Mice treated with TOK-001 have smaller average tumor volume on day 31 when compared to control (p= 0.0001). TOK-001 treatment also significantly reduces the growth rate of tumor growth compared to control (p[1].

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