Proteins



Tuvusertib

Cat. No.: HY-111451 CAS No.: 1613200-51-3 Molecular Formula: C₁₆H₁₂F₂N₈O Molecular Weight: 370.32 Target: ATM/ATR

Pathway: Cell Cycle/DNA Damage; PI3K/Akt/mTOR

Storage: Powder -20°C 3 years

In solvent

4°C 2 years 6 months -80°C

-20°C 1 month

SOLVENT & SOLUBILITY

In Vitro

DMSO: 5 mg/mL (13.50 mM; Need ultrasonic)

Preparing Stock Solutions	Solvent Mass Concentration	1 mg	5 mg	10 mg
	1 mM	2.7004 mL	13.5018 mL	27.0037 mL
	5 mM	0.5401 mL	2.7004 mL	5.4007 mL
	10 mM	0.2700 mL	1.3502 mL	2.7004 mL

Please refer to the solubility information to select the appropriate solvent.

BIOLOGICAL ACTIVITY

Description Tuvusertib (M1774; ATR inhibitor 1) is a selective and orally active ATR inhibitor extracted from patent W02015187451A1, compound I-l, with a K_i value below 1 $\mu M^{[1]}$.

IC₅₀ & Target Ki: less than 1 uM^[1]

In Vitro Tuvusertib (0-2 μM, 24 h) induces cell death and cell apoptosis in myeloma cells (such as U266 and OPM2 cells)^[2].

 $Tuvus ertib~(0-5~\mu\text{M}, 16~or~24~h)~inhibits~STAT3~p-Y705~phosphorylation~and~down-regulates~STAT3~downstream~targets~(c-1)~color=1.00~color=$ MYC) in U266 and OPM2 cells^[2].

MCE has not independently confirmed the accuracy of these methods. They are for reference only.

Western Blot Analysis^[2]

Cell Line: U266 and OPM2 cells Concentration: 0-2 μΜ

Incubation Time:	24 h
Result:	Increased level of γH2A.X, cleavage of caspase-3, and cleavage of PARP.

REFERENCES

[1]. Lin Li, et al. Non-canonical role for the ataxia-telangiectasia-Rad3 pathway in STAT3 activation in human multiple myeloma cells

[2]. Nadia AHMAD, et al. Radiolabelled derivatives of a 2-amino-6-fluoro-n-[5-fluoro-pyridin-3-yl]- pyrazolo[1,5-a]pyrimidin-3-carboxamide compound useful as atr kinase inhibitor, the preparation of said compound and different solid forms thereof. WO2015187451A1

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

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