



**AZ20** 

**Catalog No: tcsc1483** 

3

## **Available Sizes**

Size: 5mg

Size: 10mg

Size: 50mg

Size: 100mg



## **Specifications**

CAS No:

1233339-22-4

Formula:

 $C_{21}H_{24}N_4O_3S$ 

**Pathway:** 

Cell Cycle/DNA Damage;PI3K/Akt/mTOR

**Target:** 

ATM/ATR;ATM/ATR

**Purity / Grade:** 

>98%

**Solubility:** 

10 mM in DMSO

**Observed Molecular Weight:** 

412.51

## **Product Description**

AZ20 is a potent and selective inhibitor of **ATR** with an  $IC_{50}$  of 5 nM, and has 8-fold selectivity against **mTOR** ( $IC_{50}$ =38 nM).





IC50 & Target: IC50: 5 nM (ATR), 38 nM (mTOR) $^{[1]}$ 

In Vitro: AZ20 inhibits ATR immunoprecipitated from HeLa nuclear extracts with an IC $_{50}$  of 5 nM and ATR mediated phosphorylation of Chk1 in HT29 colorectal adenocarcinoma tumor cells with an IC $_{50}$  of 50 nM $^{[1]}$ .

*In Vivo:* AZ20 (25, 50 mg/kg, p.o.) has high permeability combined with good stability to rat hepatocytes and, despite the lack of progress in achieving markedly higher solubility, has respectable bioavailability in a low dose rat PK study. AZ20 (25, 50 mg/kg, p.o.) leads to significant tumor growth inhibition in female nude mice bearing LoVo tumors<sup>[1]</sup>.

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