ARQ 621

MedChemExpress

SOLVENT & SOLUBILITY

In Vitro

Cat. No.:	HY-16062
CAS No.:	1095253-39-6
Molecular Formula:	$C_{28}H_{24}Cl_2FN_5O_2$
Molecular Weight:	552.43
Target:	Kinesin
Pathway:	Cell Cycle/DNA Damage; Cytoskeleton
Storage:	-20°C, protect from light * In solvent : -80°C, 6 months; -20°C, 1 month (protect from light)

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Solvent Concentration

DMSO: 110 mg/mL (199.12 mM; Need ultrasonic)

Preparing Stock Solutions	1 mM	1.8102 mL	9.0509 mL	18.1018 mL
	5 mM	0.3620 mL	1.8102 mL	3.6204 mL
	10 mM	0.1810 mL	0.9051 mL	1.8102 mL

1 mg

Mass

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

BIOLOGICAL ACTIVITY		
Description	ARQ 621 is an allosteric, potent and selective inhibitor of Eg5, a microtubule-based ATPase motor protein involved in cell division. Anti-tumor activity ^[1] . ARQ 621 is a kinesin inhibitor ^[2] . ARQ 621 is a click chemistry reagent, it contains an Alkyne group and can undergo copper-catalyzed azide-alkyne cycloaddition (CuAAc) with molecules containing Azide groups.	
IC ₅₀ & Target	Eg5	
In Vitro	Over-expression of Eg5 causes genomic instability and tumor formation in mice; therefore, Eg5 is a potential anti-cancer target ^[1] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.	

REFERENCES

[1]. L. C. Chen, et al. First-in-human study with ARQ 621, a novel inhibitor of Eg5: Final results from the solid tumors cohort. J Clin Oncol. 2011, May (20): 3076-3076.

[2]. Lindsay S Roberts, et al. Mapping Novel Metabolic Nodes Targeted by Anti-Cancer Drugs that Impair Triple-Negative Breast Cancer Pathogenicity. ACS Chem Biol. 2017

 NH_2

10 mg

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Product Data Sheet

CI

5 mg



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

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