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Zuschläge

- Mindermengenzuschlag
- Trockeneiszuschlag
- Gefahrgutzuschlag
- Expressversand

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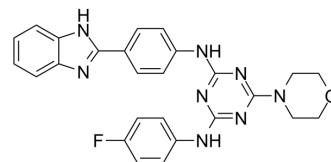
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IITZ-01

Cat. No.:	HY-112897
CAS No.:	1807988-47-1
Molecular Formula:	C ₂₆ H ₂₃ FN ₈ O
Molecular Weight:	482.51
Target:	Autophagy; PI3K; Apoptosis
Pathway:	Autophagy; PI3K/Akt/mTOR; Apoptosis
Storage:	Powder -20°C 3 years 4°C 2 years In solvent -80°C 2 years -20°C 1 year



SOLVENT & SOLUBILITY

In Vitro	DMSO : 150 mg/mL (310.87 mM; Need ultrasonic)					
	Preparing Stock Solutions	Solvent Concentration	Mass	1 mg	5 mg	10 mg
		1 mM		2.0725 mL	10.3625 mL	20.7250 mL
		5 mM		0.4145 mL	2.0725 mL	4.1450 mL
		10 mM		0.2072 mL	1.0362 mL	2.0725 mL
Please refer to the solubility information to select the appropriate solvent.						
In Vivo	1. Add each solvent one by one: 10% DMSO >> 40% PEG300 >> 5% Tween-80 >> 45% saline Solubility: ≥ 2.5 mg/mL (5.18 mM); Clear solution					
	2. Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline) Solubility: 2.5 mg/mL (5.18 mM); Suspended solution; Need ultrasonic					
	3. Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: ≥ 2.5 mg/mL (5.18 mM); Clear solution					

BIOLOGICAL ACTIVITY

Description	IITZ-01 is a potent lysosomotropic autophagy inhibitor with single-agent antitumor activity, with an IC ₅₀ of 2.62 μM for PI3K γ.	
IC ₅₀ & Target	PI3Kγ 2.62 μM (IC ₅₀)	Autophagy
In Vitro	IITZ-01 (0-2 μM, 24 h) enhances autophagosomes formation as indicated by increased expression of LC3-II levels time- and	

dose-dependently in triple-negative breast cancer (TNBC) cell lines (MDA-MB-231 and MDA-MB-453). IITZ-01 also demonstrates potent autophagy inhibitory activity in other breast, lung, and colon cancer cells^[1]. MCE has not independently confirmed the accuracy of these methods. They are for reference only.

Western Blot Analysis^[1]

Cell Line:	Triple-negative breast cancer (TNBC) cell lines (MDA-MB-231 and MDA-MB-453).
Concentration:	0-2 μ M.
Incubation Time:	24 hours.
Result:	Enhanced autophagosomes formation as indicated by increased expression of LC3-II levels.

In Vivo

IITZ-01 (45 mg/kg, i.p. every alternate day for 4 weeks) inhibits average breast tumor growth when compared with control from third day of treatment in triple-negative breast tumor models in mice^[1]. MCE has not independently confirmed the accuracy of these methods. They are for reference only.

Animal Model:	MDA-MB-231 (TNBC)/green fluorescent protein (GFP) orthotopic breast cancer xenografts were developed in CrTac:NCr-Foxn ^{nu} BALB/c female nude mice ^[1] .
Dosage:	45 mg/kg.
Administration:	Intraperitoneal every alternate day for 4 weeks.
Result:	Inhibited average breast tumor growth when compared with control from third day of treatment.

REFERENCES

[1]. Guntuku L, et al. IITZ-01, a novel potent lysosomotropic autophagy inhibitor, has single-agent antitumor efficacy in triple-negative breast cancer in vitro and in vivo. *Oncogene*. 2018 Aug 30.

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

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