iFSP1

Cat. No.:	HY-136057		
CAS No.:	150651-39-1		
Molecular Formula:	C ₂₀ H ₁₃ N ₅		
Molecular Weight:	323.35		
Target:	Ferroptosis		
Pathway:	Apoptosis		
Storage:	Powder	-20°C	3 years
		4°C	2 years
	In solvent	-80°C	2 years
		-20°C	1 year

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Preparing Stock Solutions		Mass Solvent Concentration	1 mg	5 mg	10 mg	
	1 mM	3.0926 mL	15.4631 mL	30.9262 mL		
		5 mM	0.6185 mL	3.0926 mL	6.1852 mL	
	10 mM	0.3093 mL	1.5463 mL	3.0926 mL		
	Please refer to the solubility information to select the appropriate solvent.					
n Vivo	1. Add each solvent Solubility: 2 mg/n	one by one: 10% DMSO >> 40% PE(nL (6.19 mM); Suspended solution; N	G300 >> 5% Tween-8 eed ultrasonic	0 >> 45% saline		

Description	iFSP1 is a potent, selective and glutathione-independent inhibitor of ferroptosis suppressor protein 1 (FSP1) (AIFM2) with an EC ₅₀ of 103 nM. iFSP1 selectively induces ferroptosis in GPX4-knockout cells which overexpressed FSP1. iFSP1 is able to sensitize a variety of human cancer cell lines to the ferroptosis inducer, such as (1S,3R)-RSL3 (HY-100218A) ^[1] .			
IC ₅₀ & Target	EC50: 103 nM (FSP1) ^[1]			
In Vitro	iFSP1 (0.001-1 μM; 24 hours) inhibits the Gpx4-knockout cell growth as a dose-dependent manner, but does not inhibit the wild type cell growth. Treatment with the ferroptosis inhibitor Lip-1 protects GPX4- knockout cells from iFSP1-induced ferroptosis ^[1] . iFSP1 (0.001-1 μM; 24 hours) is less efficient than genetic deletion of FSP1, whereas iFSP1 treatment in the FSP1-knockout background had no additive effect to RSL3-induced ferroptosis ^[1] . iFSP1 (3 μM; 24 hours) treatment results in an obvious toxicity of RSL3 in a panel of genetically engineered (FSP1-knockout)			

Product Data Sheet

 $H_2 N$

Ν

\/\ N human cancer cell lines^[1].

AIFM2: the flavoprotein apoptosis-inducing factor mitochondria-associated 2 is a previously unrecognized anti-ferroptotic gene. AIFM2, which is renamed ferroptosis suppressor protein 1 (FSP1)^[1]

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

Cell Proliferation Assay^[1]

Cell Line:	Wild-type and Gpx4-knockout Pfa1 or HT 1080 cells overexpressing FSP1–HA		
Concentration:	0.001-1 μΜ		
Incubation Time:	24 hours		
Result:	Was toxic to cells that depend solely (no GPX4 expression detectable) on FSP1 function.		
Cell Viability Assay ^[1]			
Cell Line:	NCl-H1437, NCl-H1437 FSP1 KO, U-373, U-373 FSP1 KO, MDA-MB-436, MDA-MB-436 FSP1 KO, SW620, SW620 FSP1 KO, MDA-MB-435S, MDA-MB-435S FSP1 KO, A549 and A549 FSP1 KO		
Concentration:	3 μΜ		
Incubation Time:	24 hours		
Result:	Sensitized a variety of human cancer cell lines to the ferroptosis inducer (1S,3R)-RSL3.		

CUSTOMER VALIDATION

- Cell Discov. 2022 May 3;8(1):40.
- Adv Mater. 2023 Jan 13;e2211579.
- Nat Commun. 2023 Oct 30;14(1):6908.
- Small. 2021 Aug;17(32):e2101368.
- Adv Healthc Mater. 2023 Jul 11;e2300994.

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REFERENCES

[1]. Doll S, et al. FSP1 is a glutathione-independent ferroptosis suppressor.Nature. 2019 Nov;575(7784):693-698.

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

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