hFSH-β-(33-53) (TFA)

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®

Cat. No.:	НҮ-Р3343А			
Molecular Formula:	C ₁₁₇ H ₁₈₄ F ₃ N ₃₁ O ₃₄ S			
Molecular Weight:	2657.96			
Sequence Shortening:	YTRDLVYKDPARPKIQKTCTF (TFA salt			
Target:	Estrogen Receptor/ERR			
Pathway:	Vitamin D Related/Nuclear Receptor			
Storage:	Sealed storage, away from moisture Powder -80°C 2 years -20°C 1 year * In solvent : -80°C, 6 months; -20°C, 1 month (sealed storage, away from moisture)			

SOLVENT & SOLUBILITY

		Solvent Mass Concentration	1 mg	5 mg	10 mg		
	Preparing Stock Solutions	1 mM	0.3762 mL	1.8811 mL	3.7623 mL		
		5 mM	0.0752 mL	0.3762 mL	0.7525 mL		
		10 mM	0.0376 mL	0.1881 mL	0.3762 mL		
	Please refer to the so	lubility information to select the app	propriate solvent.				
In Vivo		1. Add each solvent one by one: 10% DMSO >> 40% PEG300 >> 5% Tween-80 >> 45% saline Solubility: ≥ 2.5 mg/mL (0.94 mM); Clear solution					
		2. Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline) Solubility: ≥ 2.5 mg/mL (0.94 mM); Clear solution					
		 Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: ≥ 2.5 mg/mL (0.94 mM); Clear solution 					

BIOLOGICAL ACTIVITY			
Description	hFSH-β-(33-53) TFA, a thiol-containing peptide which corresponds to a second FSH receptor-binding domain, is a FSHR (follicle-stimulating hormone receptor) antagonist. hFSH-β-(33-53) TFA inhibits binding of FSH to receptor and is a partial agonist of estradiol synthesis in Sertoli cells ^{[1][2][3]} .		

REFERENCES

[1]. Grasso P, et al. A synthetic peptide corresponding to hFSH-beta-(81-95) has thioredoxin-like activity. Mol Cell Endocrinol. 1991;78(3):163-170.

[2]. Xu Y, et al. Pilot study of a novel (18)F-labeled FSHR probe for tumor imaging. Mol Imaging Biol. 2014;16(4):578-585.

[3]. Santa-Coloma TA, et al. Serine analogues of hFSH-beta-(33-53) and hFSH-beta-(81-95) inhibit hFSH binding to receptor. Biochem Biophys Res Commun. 1992;184(3):1273-1279.

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

 Tel: 609-228-6898
 Fax: 609-228-5909
 E-mail: tech@MedChemExpress.com

 Address: 1 Deer Park Dr, Suite Q, Monmouth Junction, NJ 08852, USA