## GGTI-286 TFA

**MedChemExpress** 

Cat. No.:	HY-115489A	
Molecular Formula:	$C_{25}H_{32}F_{3}N_{3}O_{5}S$	
Molecular Weight:	543.6	Ţ
Target:	Ras	
Pathway:	GPCR/G Protein	
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.	1972 V

Inhibitors

BIOLOGICAL ACTIVITY				
Description	GGTI-286 TFA, a potent and cell-permeable GGTase I inhibitor, is 25-fold more potent (IC <sub>50</sub> =2 μM) than the corresponding methyl ester of FTI-276 (HY-15873A). GGTI-286 TFA selectively inhibits geranylgeranylation of Rap1A over farnesylation of H-Ras in NIH3T3 cells (IC <sub>50</sub> s=2 and >30 μM, respectively). GGTI-286 TFA also potently inhibits oncogenic K-Ras4B stimulation with an IC <sub>50</sub> of 1 μM <sup>[1][2]</sup> .			
In Vitro	GGTI-286 (10 μM; 2 and 4 h; CHO cells) TFA reduces nuclear localization of β-catenin and transcription dependent on β- catenin/T cell factor in mammalian cells <sup>[2]</sup> . MCE has not independently confirmed the accuracy of these methods. They are for reference only. Western Blot Analysis <sup>[2]</sup>			
	Cell Line:	CHO cells		
	Concentration:	10 μΜ		
	Incubation Time:	2 and 4 h		
	Result:	Reduced the nuclear b-catenin amount.		

## REFERENCES

[1]. E C Lerner, et al. Disruption of oncogenic K-Ras4B processing and signaling by a potent geranylgeranyltransferase I inhibitor. J Biol Chem. 1995 Nov 10;270(45):26770-3.

[2]. Naoyuki Nishiya, et al. A zebrafish chemical suppressor screening identifies small molecule inhibitors of the Wnt/β-catenin pathway. Chem Biol. 2014 Apr 24;21(4):530-540.

## 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