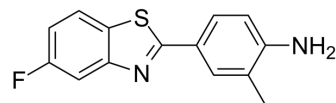


## 5F-203

<b>Cat. No.:</b>	HY-124421	
<b>CAS No.:</b>	260443-89-8	
<b>Molecular Formula:</b>	C <sub>14</sub> H <sub>11</sub> FN <sub>2</sub> S	
<b>Molecular Weight:</b>	258.31	
<b>Target:</b>	Aryl Hydrocarbon Receptor	
<b>Pathway:</b>	Immunology/Inflammation	
<b>Storage:</b>	Powder	-20°C 3 years 4°C 2 years
	In solvent	-80°C 6 months -20°C 1 month



### SOLVENT & SOLUBILITY

<b>In Vitro</b>	DMSO : 100 mg/mL (387.13 mM; Need ultrasonic)				
		Solvent Concentration	Mass 1 mg	5 mg	10 mg
	<b>Preparing Stock Solutions</b>	1 mM	3.8713 mL	19.3566 mL	38.7132 mL
		5 mM	0.7743 mL	3.8713 mL	7.7426 mL
10 mM		0.3871 mL	1.9357 mL	3.8713 mL	
Please refer to the solubility information to select the appropriate solvent.					
<b>In Vivo</b>	1. Add each solvent one by one: 10% DMSO >> 40% PEG300 >> 5% Tween-80 >> 45% saline Solubility: ≥ 2.5 mg/mL (9.68 mM); Clear solution				

### BIOLOGICAL ACTIVITY

<b>Description</b>	5F-203 (NSC-703786) is a cytotoxic molecule that forms DNA adducts and cell cycle arrest. 5F-203 induces aryl hydrocarbon receptor (AhR) signaling and elevates expression of CYP1A1. 5F-203 also increases the levels of reactive oxygen species as well as activates JNK, ERK, and p38 <sup>[1][2][3]</sup> .
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### REFERENCES

[1]. Hutchinson I, et al. Antitumor benzothiazoles. 16. Synthesis and pharmaceutical properties of antitumor 2-(4-aminophenyl)benzothiazole amino acid prodrugs. J Med Chem. 2002 Jan 31;45(3):744-7.

[2]. Hose CD, et al. Induction of CYP1A1 in tumor cells by the antitumor agent 2-[4-amino-3-methylphenyl]-5-fluoro-benzothiazole: a potential surrogate marker for patient sensitivity. Mol Cancer Ther. 2003 Dec;2(12):1265-72.

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[3]. Callero MA, et al. Biomarkers of sensitivity to potent and selective antitumor 2-(4-amino-3-methylphenyl)-5-fluorobenzothiazole (5F203) in ovarian cancer. J Cell Biochem. 2013 Oct;114(10):2392-404.

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

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