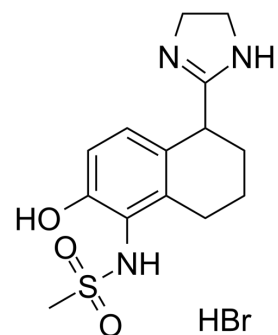


## A-61603

<b>Cat. No.:</b>	HY-101366
<b>CAS No.:</b>	107756-30-9
<b>Molecular Formula:</b>	C <sub>14</sub> H <sub>20</sub> BrN <sub>3</sub> O <sub>3</sub> S
<b>Molecular Weight:</b>	390.3
<b>Target:</b>	Adrenergic Receptor
<b>Pathway:</b>	GPCR/G Protein; Neuronal Signaling
<b>Storage:</b>	-20°C, sealed storage, away from moisture * In solvent : -80°C, 6 months; -20°C, 1 month (sealed storage, away from moisture)



### SOLVENT & SOLUBILITY

#### In Vitro

DMSO : 20 mg/mL (51.24 mM; Need ultrasonic and warming)

Concentration	Mass			
	1 mg	5 mg	10 mg	
1 mM	2.5621 mL	12.8107 mL	25.6213 mL	
5 mM	0.5124 mL	2.5621 mL	5.1243 mL	
10 mM	0.2562 mL	1.2811 mL	2.5621 mL	

Please refer to the solubility information to select the appropriate solvent.

### BIOLOGICAL ACTIVITY

#### Description

A-61603 is a selective  $\alpha_{1A}$ -adrenergic receptor agonist<sup>[1]</sup>. A-61603 increases the frequency of spontaneous Ca<sup>2+</sup> transients in rat ventricular myocytes in vitro<sup>[2]</sup>.

#### IC<sub>50</sub> & Target

$\alpha_{1A}$ -adrenergic receptor

### REFERENCES

[1]. Meyer MD, et al. Synthesis and in vitro characterization of N-[5-(4,5-dihydro-1H-imidazol-2-yl)-2-hydroxy-5,6,7,8-tetrahydronaphthalen-1-yl]methanesulfonamide and its enantiomers: A novel selective  $\alpha_{1A}$  receptor agonist. J. Med. Chem. 39(20), 4116-4119 (1

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

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