## Fabomotizole

HY-14895		
173352-21-2	L	
C <sub>15</sub> H <sub>21</sub> N <sub>3</sub> O <sub>2</sub> S	i	
307.41		
Others		
Others		
Pure form	-20°C	3 years
In solvent	-80°C	6 months
	-20°C	1 month
	173352-21-2 C <sub>15</sub> H <sub>21</sub> N <sub>3</sub> O <sub>2</sub> S 307.41 Others Others Pure form	173352-21-1 C <sub>15</sub> H <sub>21</sub> N <sub>3</sub> O <sub>2</sub> S 307.41 Others Others Pure form -20°C In solvent -80°C

## SOLVENT & SOLUBILITY

	Solvent Concentration Preparing 1 mM Stock Solutions 5 mM 10 mM		1 mg	5 mg	10 mg	
		1 mM	3.2530 mL	16.2649 mL	32.5298 mL	
		5 mM	0.6506 mL	3.2530 mL	6.5060 mL	
		10 mM	0.3253 mL	1.6265 mL	3.2530 mL	
	Please refer to the solubility information to select the appropriate solvent.					
In Vivo	1. Add each solvent one by one: 10% DMSO >> 40% PEG300 >> 5% Tween-80 >> 45% saline Solubility: ≥ 2.5 mg/mL (8.13 mM); Clear solution					
	2. Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline) Solubility: ≥ 2.5 mg/mL (8.13 mM); Clear solution					
	<ol> <li>Add each solvent one by one: 10% DMSO &gt;&gt; 90% corn oil Solubility: ≥ 2.5 mg/mL (8.13 mM); Clear solution</li> </ol>					

BIOLOGICAL ACTIVITY						
Description	Fabomotizole (CM346) is an anxiolytic agent. Fabomotizole produces anxiolytic and neuroprotective effects without any muscle relaxant actions.					
In Vitro	Afobazole's mechanism of action remains poorly defined however, with GABAergic, NGF and BDNF release promoting, MT1 receptor antagonism, MT3 receptor antagonism, and sigma agonism suggested as potential mechanisms. Afobazole was shown to inhibit MAO-A reversibly and there might be also some involvement with serotonin receptors. Afobazole has found little clinical use outside of Russia and has not been evaluated by the FDA. MCE has not independently confirmed the accuracy of these methods. They are for reference only.					

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## REFERENCES

[1]. Afobazole, From Wikipedia

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

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