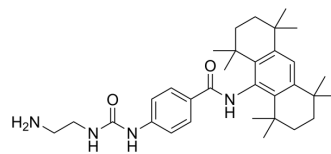


## OAB-14

<b>Cat. No.:</b>	HY-139973		
<b>CAS No.:</b>	2140911-49-3		
<b>Molecular Formula:</b>	C <sub>32</sub> H <sub>46</sub> N <sub>4</sub> O <sub>2</sub>		
<b>Molecular Weight:</b>	518.73		
<b>Target:</b>	Amyloid- $\beta$		
<b>Pathway:</b>	Neuronal Signaling		
<b>Storage:</b>	Powder	-20°C	3 years
		4°C	2 years
	In solvent	-80°C	6 months
		-20°C	1 month



### SOLVENT & SOLUBILITY

#### In Vitro

DMSO : 20 mg/mL (38.56 mM; ultrasonic and warming and adjust pH to 3 with HCl and heat to 60°C)

Concentration	Mass		
	1 mg	5 mg	10 mg
1 mM	1.9278 mL	9.6389 mL	19.2779 mL
5 mM	0.3856 mL	1.9278 mL	3.8556 mL
10 mM	0.1928 mL	0.9639 mL	1.9278 mL

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

### BIOLOGICAL ACTIVITY

#### Description

OAB-14, is a Bexarotene (HY-14171) derivative, improves Alzheimer's disease-related pathologies and cognitive impairments by increasing  $\beta$ -amyloid clearance in APP/PS1 mice. OAB-14 effectively ameliorates the dysfunction of the endosomal-autophagic-lysosomal pathway in APP/PS1 transgenic mice<sup>[1][2]</sup>.

#### In Vivo

OAB-14 significantly alleviates cognitive impairments in amyloid precursor protein (APP)/presenilin 1 (PS1) transgenic mice after administration for 15 days or 3 months. OAB-14 rapidly cleared 71% of A $\beta$  by promoting microglia phagocytosis and increasing IDE and NEP expression. OAB-14 also attenuates the downstream pathological events of A $\beta$  accumulation, such as synaptic degeneration, neuronal loss, tau hyperphosphorylation and neuroinflammation in APP/PS1 mice. OAB-14 has no significant effect on body weight or liver toxicity after acute and chronic treatment<sup>[1]</sup>.

OAB-14 facilitates receptor-mediated endocytosis and restores autophagy flux via the AMPK/mTOR pathway. OAB-14 enhances the lysosomal activity, and reduced A $\beta$  accumulation in lysosomes is observed in OAB-14-treated AD mice<sup>[2]</sup>.

MCE has not independently confirmed the accuracy of these methods. They are for reference only.

### REFERENCES

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[1]. Guo X, et al. OAB-14 Effectively Ameliorates the Dysfunction of the Endosomal-Autophagic-Lysosomal Pathway in APP/PS1 Transgenic Mice. ACS Chem Neurosci. 2021;12(21):3985-3993.

[2]. Yuan C, et al. OAB-14, a bexarotene derivative, improves Alzheimer's disease-related pathologies and cognitive impairments by increasing  $\beta$ -amyloid clearance in APP/PS1 mice. Biochim Biophys Acta Mol Basis Dis. 2019;1865(1):161-180.

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

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