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产品名称: **T807**  
 产品别名: **AV-1451**

生物活性:					
<b>Description</b>	T807 a novel tau positron emission tomography (PET) tracer.				
<b>In Vitro</b>	Aggregated tau protein is a major neuropathological substrate central to the pathophysiology of neurodegenerative diseases such as Alzheimer's disease (AD). <i>In vitro</i> autoradiography results show that [ <sup>18</sup> F]T807 exhibits strong binding to PHF-tau positive human brain sections ( $K_d=14.6$ nM). A comparison of autoradiography and double immunohistochemical staining of PHF-tau and Ab on adjacent sections demonstrates that [ <sup>18</sup> F]T807 binding colocalizes with immunoreactive PHF-tau pathology, but does not highlight Ab plaques <sup>[1]</sup> . [ <sup>18</sup> F]T807 strongly binds to tau lesions primarily made of paired helical filaments in Alzheimer's brains e.g. intra and extraneuronal tangles and dystrophic neurites. [ <sup>18</sup> F]T807 off-target binding to neuromelanin- and melanin-containing cells and, to a lesser extent, to brain hemorrhagic lesions is identified <sup>[2]</sup> .				
<b>In Vivo</b>	[ <sup>18</sup> F]T807 is able to cross the blood-brain barrier and ished out quickly in mice model. [ <sup>18</sup> F]T807 clears rapidly from the brain, with activity values decreasing from 4.43% ID/g at 5 minutes to 0.62% ID/g at 30 minutes. Kidney elimination is a significant clearance pathway, resulting in a maximum tracer concentration of 14.99% ID/g in the kidneys at 5 minutes, which decreases to 5.52% ID/g at 30 minutes. The accumulation of activity in muscle and bone remain relatively low throughout the PET scan <sup>[1]</sup> .				
<b>Solvent&amp;Solubility</b>	<b><i>In Vitro:</i></b> DMSO : $\geq 16.6$ mg/mL (63.05 mM) * ">" means soluble, but saturation unknown.				
		Solvent / Mass / Concentration	1 mg	5 mg	10 mg
	<b>Preparing</b>	1 mM	3.7984 mL	18.9919 mL	37.9838 mL
	<b>Stock Solutions</b>	5 mM	0.7597 mL	3.7984 mL	7.5968 mL
		10 mM	0.3798 mL	1.8992 mL	3.7984 mL
	*请根据产品在不同溶剂中的溶解度选择合适的溶剂配制储备液, 一旦配成溶液, 请分装保存, 避免反复冻融造成的产品失效。 储备液的保存方式和期限: -80°C, 6 months; -20°C, 1 month。-80°C 储存时, 请在 6 个月内使用, -20°C 储存时, 请在 1 个月内使用。				
<b>References</b>	[1]. Xia CF, et al. [(18)F]T807, a novel tau positron emission tomography imaging agent for Alzheimer's disease. <i>Alzheimers Dement.</i> 2013 Nov;9(6):666-76. [2]. Marquié M, et al. Validating novel tau positron emission tomography tracer [F-18]-AV-1451 (T807) on postmortem brain tissue. <i>Ann Neurol.</i> 2015 Nov;78(5):787-800.				
实验参考:					
<b>Animal Administration</b>	Mice: Six male mice at each time point are administered 250 mCi [ <sup>18</sup> F]T807 (in 200 mL saline) via tail vein injection. At 5, 15, and 30 minutes after administration, the mice are anesthetized and 500-mL whole blood samples and centrifuged. After euthanasia, the liver, kidneys, skeletal muscle (right quadriceps), brain, and bone (femur) are harvested and weighed. Each of the tissue samples				



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	are transferred to gamma counter tubes and counted <sup>[1]</sup> .
<b>Kinase Assay</b>	10 mg/mL frozen brain homogenate aliquots are thawed and diluted 10-fold in binding buffer to 1 mg/mL. 500 $\mu$ L of appropriate concentrations of non-radioactive T807 to be tested are combined with 400 $\mu$ L of [ <sup>3</sup> H] T807 (29.7 Ci/mmol) in a volume of 900 $\mu$ L of binding buffer. The assay begins by addition of 100 $\mu$ L of the 1 mg/mL brain homogenate to achieve a final concentration of 0.10 mg tissue/mL for radioligands. The final concentration of [ <sup>3</sup> H] T807 is typically 1-2 nM. After incubation at room temperature for 60 minutes, the binding mixture is filtered and rapidly washed 5 times with 3 mL binding buffer. The filters are counted <sup>[2]</sup> .
<b>References</b>	[1]. Xia CF, et al. [(18)F]T807, a novel tau positron emission tomography imaging agent for Alzheimer's disease. <i>Alzheimers Dement.</i> 2013 Nov;9(6):666-76. [2]. Marquié M, et al. Validating novel tau positron emission tomography tracer [F-18]-AV-1451 (T807) on postmortem brain tissue. <i>Ann Neurol.</i> 2015 Nov;78(5):787-800.



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