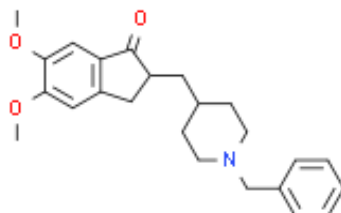


Product Data Sheet

Cas No.:	120011-70-3	Cat. No:	PL09449
Product Name:	Donepezil Hydrochloride		
Product synonym:	2,3-二氢-5,6-二甲氧基-2-[[[(1-苯甲基)-4-哌啶基]甲基]-1H-茚-1-酮]盐酸盐;盐酸多奈哌齐;多奈哌齐盐酸盐;多奈哌齐盐酸盐;2-[[[(1-苄基-4-哌啶基)甲基]-5,6-二甲氧基-2,3-二氢-1-茚酮]盐酸盐;1-苄基-4-[[[5,6-二甲氧基-1-茚满酮-2-基]甲基]哌啶盐酸盐;3-氟-2-氰基苯乙酰胺;Donepezil HCL 盐酸多奈哌齐;Donepezil Hydrochloride 盐酸多奈哌齐;多奈哌齐;多奈哌齐-D5盐酸;多奈哌齐杂质;多奈哌齐杂质及标准品;盐酸多奈哌齐 标准品;盐酸多奈哌齐（I）;盐酸多奈哌齐（III）;盐酸多奈哌齐III型;盐酸多奈哌齐I型;盐酸多奈哌齐标准品(JP);盐酸多奈哌齐中间体;盐酸多奈哌齐;2-(1-苄基-4-哌啶基甲基)-5,6-二甲氧基-1-二氢茚酮盐酸盐;多奈派齐;盐酸多奈哌;盐酸多奈哌齐(I)型晶;盐酸多奈哌齐(III);多奈哌齐盐酸盐 10MG		
Chemical name:	Donepezil Hydrochloride		
MF:	C24H30CLNO3	FW:	415.9529
Purity:	≥99%	Batch No.:	-
Storage:			
Structural formula:	 H Cl		
λmax:	-	Formulation:	-
Solubility :			
SMILES :	Cl[H].O=C1C2=C([H])C(=C(C([H])=C2C([H])([H])C1([H])C([H])([H])C1([H])C([H])([H])C([H])([H])N(C([H])([H])C2C([H])=C([H])C([H])=C([H])C=2[H])C([H])([H])C1([H])([H])OC([H])([H])([H])OC([H])([H])([H])[H]		
InChI Code:	-		
InChI Key:			
WARNING This product is not for human or veterinary use.			

Product Description

Donepezil Hydrochloride (E2020) 是一种可逆的选择性 AChE 抑制剂，对 AChE 的 IC₅₀ 为 6.7 nM。对 AChE 的选择性高于 BuChE。Donepezil Hydrochloride 对 Aβ₄₂ 神经毒性具有神经保护作用。

生物活性	Donepezil Hydrochloride (E2020) is a reversible, selective AChE inhibitor with an IC ₅₀ of 6.7 nM for AChE activity. Donepezil shows high selectivity for AChE over BuChE. Donepezil exhibits neuroprotective effect on Aβ ₄₂ neurotoxicity.
体外研究(In Vitro)	Donepezil's neuroprotective mechanism is related to the enhanced phosphorylation of Akt and GSK-3β and reduced phosphorylation of tau and glycogen synthase. has not independently confirmed the accuracy of these methods. They are for reference only.Cell Viability Assay

体内研究(In Vivo)	Donepezil treatment (3 mg/kg) significantly prevents the progression of scopolamine-induced memory impairment in mice. A pharmacokinetic study of Donepezil shows a mean peak plasma concentration of donepezil after oral treatment (3 and 10 mg/kg) of approximately 1.2±0.4 h and 1.4±0.5 h, respectively; absolute bioavailability is calculated as 3.6%. has not independently confirmed the accuracy of these methods. They are for reference only.
包装储存	4°C, sealed storage, away from moisture In solvent : -80°C, 6 months; -20°C, 1 month (sealed storage, away from moisture)
溶解度数据	In Vitro: H ₂ O : 25 mg/mL (60.10 mM; Need ultrasonic)DMSO : 6.2 mg/mL (14.91 mM; Need warming)配制储备液