

产品名称: **Pamidronic Acid See P172500**

产品别名: **帕米膦酸; Pamidronic acid**

生物活性:					
Description	Pamidronic acid is a drug used to treat a broad spectrum of bone absorption diseases.				
IC₅₀ & Target	Wnt, β -catenin[1]				
In Vitro	Osteosarcoma cell viability decreases significantly in a concentration- and time-dependent manner at pamidronate concentrations ranging from 100 to 1000 μ M, most consistently after 48 and 72 hours' exposure. In treated osteosarcoma cells, the lowest percentage cell viability is 34% (detected after 72 hours' exposure to 1000 μ M pamidronate)[1]. Pamidronate disodium inhibits Wnt and β -catenin signaling, which controls osteogenic differentiation in BMSCs. Wnt3a, a Wnt and β -catenin signaling activator, reverses the negative effects caused by pamidronate disodium to salvage the osteogenic defect in BMSCs[2].				
In Vivo	Pamidronic acid can significantly inhibit and even reverse early osteoarthritic subchondral bone loss, thus alleviating the process of cartilaginous degeneration. The mechanisms involved may be associated with the upregulation of OPG expression, and downregulation of RANKL, MMP-9 and TLR-4 expression[3].				
Solvent&Solubility	In Vitro: H ₂ O : 6 mg/mL (25.52 mM; Need ultrasonic and warming)				
		Solvent \ Mass Concentration	1 mg	5 mg	10 mg
	Preparing	1 mM	4.2541 mL	21.2703 mL	42.5405 mL
	Stock Solutions	5 mM	0.8508 mL	4.2541 mL	8.5081 mL
		10 mM	0.4254 mL	2.1270 mL	4.2541 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。					
References	<p>[1]. Ashton JA, et al. Investigation of the effect of pamidronate disodium on the in vitro viability of osteosarcoma cells from dogs. Am J Vet Res. 2005 May;66(5):885-91.</p> <p>[2]. Xu Y, et al. Pamidronate Disodium Leads to Bone Necrosis via Suppression of Wnt/β-Catenin Signaling in Human Bone Marrow Mesenchymal Stem Cells In Vitro. J Oral Maxillofac Surg. 2017 Mar 22.</p> <p>[3]. Lv Y, et al. Effects of pamidronate disodium on the loss of osteoarthritic subchondral bone and the expression of cartilaginous and subchondral osteoprotegerin and RANKL in rabbits. BMC Musculoskelet Disord. 2014 Nov 6;15:370.</p>				
实验参考:					
Cell Assay	Cell counts and cell viability assays are performed in cultures of osteosarcoma cells (POS, HMPOS, and COS31 cell lines) and fibroblasts after 24, 48, and 72 hours of incubation with pamidronate at concentrations of 0.001 to 1000 microM or with no drug (control treatment). Percentage viability is determined in cell samples for each concentration of pamidronate and each incubation time. A DNA fragmentation analysis is performed to assess bisphosphonate-induced apoptosis[1].				
	Rabbits: The rabbits are randomly divided into four groups. Sham-operated with vehicle treatment, OA induced by ACLT with vehicle treatment, OA-induced ACLT treated with short-term pamidronic				

Animal Administration	acid treatment after ACLT, and ACLT treated with long-term PAM treatment. PAM is injected at the 4th week after ACLT in PAM-S and PAM-L groups, and followed by once monthly ear vein injections at a dosage of 3 mg/kg body weight. In the other groups, only saline infusions of equal volumes are administered. 10 animals are humanely sacrificed at both 2 and 10 weeks after pamidronic acid treatment. In the ACLT and Sham groups, five animals are sacrificed at 2, 4, 6, and 14 weeks after model establishment[3].
References	<p>[1]. <u>Ashton JA, et al. Investigation of the effect of pamidronate disodium on the in vitro viability of osteosarcoma cells from dogs. Am J Vet Res. 2005 May;66(5):885-91.</u></p> <p>[2]. <u>Xu Y, et al. Pamidronate Disodium Leads to Bone Necrosis via Suppression of Wnt/β-Catenin Signaling in Human Bone Marrow Mesenchymal Stem Cells In Vitro. J Oral Maxillofac Surg. 2017 Mar 22.</u></p> <p>[3]. <u>Lv Y, et al. Effects of pamidronate disodium on the loss of osteoarthritic subchondral bone and the expression of cartilaginous and subchondral osteoprotegerin and RANKL in rabbits. BMC Musculoskelet Disord. 2014 Nov 6;15:370.</u></p>



源叶生物