For Research Use Only Naftifine hydrochloride



Catalog Number: CM02945

产品信息	Catalog Number: CM02945 CAS号: 65473-14-5 分子式: C ₂₁ H ₂₂ ClN	分子量: 323.86 溶解度: H2O:<1 mg/mL,Ethanol:8 mg/mL (24.7 mM),DMSO:<1 mg/mL HCI	
	主要靶点: Hedgehog/Smoothened Antibiotic 主要通路: 干细胞 G蛋白偶联受体 微生物学	Antifungal	Í
体外活性	Naftifine exhibits an interesting in vitro speconcentration (MIC) range 0.1 to 0.2 mg/mL), strains; MICs, 0.8 and 1.5 mg/mL), and yeasts. The MIC of naftifine for C. albicans \triangle 63 is 1 than 99% inhibition of sterol biosynthesis b naftifine appears to be the blocking of fungational strains of the strain	ctrum of activity against dermatophytes (38 strains; minimal inh aspergilli (6 strains; MIC range, 0.8 to 12.5 mg/mL), Sporothrix s of the genus Candida (77 strains; MIC range, 1.5 to greater than 00 mg/L in Sabouraud medium (initial pH 6.5). Naftifine (50 mg/ oth in whole cells and in cell extracts of C. albicans. The primary l squalene epoxidation. [2]	ibitory chenckii (2 100 mg/mL). [1] L) gives greater action of
体内活性	Naftifine HCl 2% cream results in clinical cu week 2 efficacy response rates in Naftifine H week 2 vehicle-treated counterparts. [3] Naf squalene in fungal organisms. Naftifine also production and a reduction in polymorphonu efficacy and safety for a variety of condition inflammatory signs and symptoms. Few adv transient burning, stinging, or itching in the a	re rate and clinical success rate of 33% and 84% after treatment ICl 2% subjects are all lower than at week 4 but are significantly tifine causes interruption of fungal ergosterol synthesis and accu has demonstrated anti-inflammatory properties such as a reduc clear leukocyte chemotaxis/endothelial adhesion. Naftifine has s and is a useful treatment that provides both antifungal action a erse events have been noted with naftifine use, the most frequer application area. [4]	for 4 weeks, and r higher than mulation of tion in superoxide shown good and relief of nt being mild and
描述	Naftifine Hydrochloride is the hydrochloride antifungal activity. Although the exact mech appears to selectively inhibit the enzyme sc a decreased amount of sterols, especially en accumulation of squalene in fungal cells. Na on the concentration and the organisms invo	salt form of naftifine, an allylamine derivate with synthetic brow nanism through which naftifine hydrochloride exerts its effect is a ualene 2, 3-epoxidase, thereby inhibiting the biosynthesis of ste gosterol which is the primary fungal membrane sterol, and a corr ftifine hydrochloride can be fungicidal as well as fungistatic to y lved.	ad-spectrum unknown, it rrol. This results in responding reasts depending
储存	Powder: -20°C for 3 years In solvent: -	80°C for 2 years	