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Product Information

Product ID F0275

CAS No. 105628-07-7

Chemical Name 1-(5-isoquinolinesulfonyl)homopiperazine hydrochloride

Synonym HA-1077; Fasudil HCl; Eril; Fasudil (Hydrochloride); AT 877

hydrochloride

Formula C₁₄H₁₇N₃O₂S • HCl

Formula Wt. 327.83

Melting Point

Purity ≥98% Solubility

HN HCI

Bulk quanitites available upon request

Product ID	Size
F0275	100 mg
F0275	250 mg
F0275	1 g

Store Temp Ambient Ship Temp Ambient

Description Fasudil is a rho-associated kinase (ROCK) inhibitor that exhibits cardioprotective, vasodilatory, neuroprotective, antiinflammatory, and anti-angiogenic activities. In animal models of myocardial ischemia/reperfusion, fasudil increases expression of Bcl-2 and p-Akt and decreases expression of Bax and caspase 3, decreasing myocardial infarction size. In other heart failure models, fasudil decreases activation of JNK, translocation of ERK, and expression of c-fos and c-jun. Fasudil also decreases activity of matrix metalloproteinase 9 (MMP9). This compound decreases aneurysm size in models of abdominal aortic aneurysm and also inhibits progression of existing aneurysms in vivo. In animal models of amyotrophic lateral sclerosis (ALS), fasudil decreases motor neuron loss, slowing disease progression and increasing survival time. In animal models of experimental autoimmune encephalitis (EAE), this compound decreases production of toll-like receptor 4 (TLR4), NF-κB, IL-1β, IL-6, and TNFα and increases production of IL-10 and cannabinoid receptor 2 (CBR2). Additionally, fasudil decreases expression of fibrotic mediators and upregulates expression of prolyl hydroxylase 2, downregulating expression of HIF-1 α in diabetic mice.

References Peng C, Gu P, Zhou J, et al. Inhibition of rho-kinase by fasudil suppresses formation and progression of experimental abdominal aortic aneurysms. PLoS One. 2013 Nov 14;8(11):e80145. PMID: 24244631.

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Caution: This product is intended for laboratory and research use only. It is not for human or drug use.