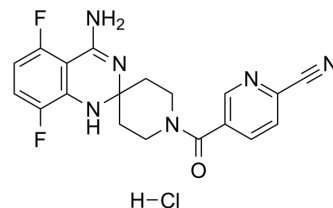


AR-C102222 hydrochloride

Cat. No.:	HY-12122A
CAS No.:	1781934-50-6
Molecular Formula:	C ₁₉ H ₁₇ ClF ₂ N ₆ O
Molecular Weight:	418.83
Target:	NO Synthase
Pathway:	Immunology/Inflammation
Storage:	4°C, sealed storage, away from moisture * In solvent : -80°C, 6 months; -20°C, 1 month (sealed storage, away from moisture)



BIOLOGICAL ACTIVITY

Description	AR-C102222 hydrochloride is a potent, competitive, orally active and highly selective inducible nitric oxide synthase (iNOS) inhibitor, with an IC ₅₀ of 37 nM ^[1] . AR-C102222 hydrochloride has antinociception and anti-inflammatory activities ^[2] .								
In Vivo	<p>AR-C102222 (3, 10, 30, 100 mg/kg, P.O.) attenuates arachidonic acid-induced ear inflammation and possesses anti-inflammatory activity^[2].</p> <p>AR-C102222 shows good efficacy in a rat adjuvant-induced arthritis model^[3].</p> <p>MCE has not independently confirmed the accuracy of these methods. They are for reference only.</p> <table border="1"> <tr> <td>Animal Model:</td> <td>Male Balb/c mice (20-25 g)^[2].</td> </tr> <tr> <td>Dosage:</td> <td>3, 10, 30, 100 mg/kg.</td> </tr> <tr> <td>Administration:</td> <td>P.O. 55 min before the administration of 0.6% acetic acid (i.p., 1 mL/100 g volume).</td> </tr> <tr> <td>Result:</td> <td>Significantly inhibited arachidonic acid-induced ear inflammation following a dose of 100 mg/kg of has a maximal inhibition of approximately 79%.</td> </tr> </table>	Animal Model:	Male Balb/c mice (20-25 g) ^[2] .	Dosage:	3, 10, 30, 100 mg/kg.	Administration:	P.O. 55 min before the administration of 0.6% acetic acid (i.p., 1 mL/100 g volume).	Result:	Significantly inhibited arachidonic acid-induced ear inflammation following a dose of 100 mg/kg of has a maximal inhibition of approximately 79%.
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REFERENCES

[1]. Tinker AC, et al. 1,2-Dihydro-4-quinazolinamines: potent, highly selective inhibitors of inducible nitric oxide synthase which show antiinflammatory activity in vivo. *J Med Chem.* 2003 Mar 13;46(6):913-6.

[2]. LaBuda CJ, et al. Antinociceptive activity of the selective iNOS inhibitor AR-C102222 in rodent models of inflammatory, neuropathic and post-operative pain. *Eur J Pain.* 2006 Aug;10(6):505-12. Epub 2005 Aug 24.

[3]. Yoon J, et al. Syntheses of 1,2,3-triazolyl salicylamides with inhibitory activity on lipopolysaccharide-induced nitric oxide production. *Bioorg Med Chem Lett.* 2011 Apr 1;21(7):1953-7.

Caution: Product has not been fully validated for medical applications. For research use only.

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