Phone: 888-558-5227

651-644-8424

Fax: 888-558-7329 Email: getinfo@lktlabs.com

Web: lktlabs.com

Product Information

Product ID N3448

CAS No. 66085-59-4

Chemical Name 1,4-Dihydro-2,6-dimethyl-4-(3-nitrophenyl)-3,5- pyridinedicarboxylic

acid 2-methoxyethyl 1-methyl- ethyl ester

Synonym Periplum, Nimotop

Formula $C_{21}H_{26}N_2O_7$ Formula Wt. 418.44 Melting Point 125°C Purity ≥98%

Solubility Soluble in methanol or DMSO. Insoluble in water.

OCH₃ CH₃ O

Pricing and Availability

Bulk quanitites available upon request

Product ID	Size	List Price
N3448	100 mg	\$49.70
N3448	500 mg	\$126.80
N3448	1 g	\$220.50
N3448	5 g	\$529.20

Store Temp ambient Ship Temp Ambient

Description Nimodipine is an antagonist at L-type Ca2+ channels that exhibits antihypertensive, neuroprotective, neuromodulatory, cognition enhancing, and anti-inflammatory activities. In animal models of subarachnoid hemorrhage, nimodipine prevents cerebral ischemia and vasospasm. Nimodipine also inhibits methylmercury-induced behavioral neurotoxicity in vivo. In animal models of Alzheimer's disease, nimodipine inhibits amyloid-8 (AB)-stimulated release of IL-1B and resulting neurotoxicity. This compound also inhibits neurological symptoms of drug-induced withdrawal, suppressing anxiety, depression, and memory impairment.

References Bia A GY, Polak P, Michalak A, et al. Influence of calcium channel antagonists on nonsomatic signs of nicotine and damphetamine withdrawal in mice. Pharmacol Rep. 2014 Apr;66(2):212-22. PMID: 24911072.

> Bailey JM, Hutsell BA, Newland MC. Dietary nimodipine delays the onset of methylmercury neurotoxicity in mice. Neurotoxicology. 2013 Jul; 37:108-17. PMID: 23583802.

Sanz JM, Chiozzi P, Colaianna M, et al. Nimodipine inhibits IL-1B release stimulated by amyloid B from microglia. Br J Pharmacol. 2012 Dec;167(8):1702-11. PMID: 22831460.

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