



LKT Laboratories, Inc.

Propofol

Phone: 888-558-5227
651-644-8424
Fax: 888-558-7329
Email: getinfo@lktlabs.com
Web: lktlabs.com

Product Information

Product ID P6870

CAS No. 2078-54-8

Chemical Name

Synonym 2,6-Bis(isopropyl)phenol, 2,6-Diisopropylphenol

Formula $C_{12}H_{18}O$

Formula Wt. 178.27

Melting Point

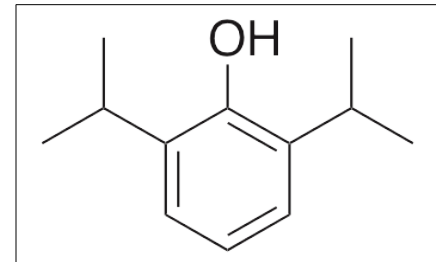
Purity $\geq 98\%$

Solubility 100mM in DMSO and ethanol

Store Temp Ambient

Ship Temp Ambient

Description Propofol is an agonist at GABA-A receptors and antagonist at voltage-gated Na⁺ channels and NMDA receptors; it also modulates Ca²⁺ signaling. Propofol displays anesthetic, sedative/hypnotic, neuroprotective, and nephroprotective activities. Propofol is a short-acting intravenous anesthetic that depresses the central nervous system, decreases cerebral blood flow, and decreases intracranial pressure. In a middle cerebral artery occlusion (MCAO) model of cerebral ischemia/reperfusion, propofol decreases expression of aquaporins 1 and 4 (AQ-1 and AQ-4), matrix metalloproteinases 2 and 9 (MMP2/9), and HIF-1 α , decreasing edema and disruption of the blood-brain barrier. Propofol also increases expression of μ -opioid receptors in neuroblastoma cells. Additionally, this compound normalizes osmolality, serum creatine, and levels of AQP-2, ICAM-1, TNF- α , Bcl-2, and Bax in kidneys, improving renal function in animal models of endotoxemia-induced kidney injury.



Pricing and Availability

Bulk quantities available upon request

Product ID	Size	List Price
P6870	5 g	\$33.20
P6870	25 g	\$63.70
P6870	100 g	\$104.50

References Lee JH, Cui HS, Shin SK, et al. Effect of propofol post-treatment on blood-brain barrier integrity and cerebral edema after transient cerebral ischemia in rats. *Neurochem Res.* 2013 Nov;38(11):2276-86. PMID: 23990224.

Li Z, Pei Q, Cao L, et al. Propofol increases μ -opioid receptor expression in SH-SY5Y human neuroblastoma cells. *Mol Med Rep.* 2012 Dec;6(6):1333-6. PMID: 22965315.

Cui WY, Tian AY, Bai T. Protective effects of propofol on endotoxemia-induced acute kidney injury in rats. *Clin Exp Pharmacol Physiol.* 2011 Nov;38(11):747-54. PMID: 21824173.

Kotani Y, Shimazawa M, Yoshimura S, et al. The experimental and clinical pharmacology of propofol, an anesthetic agent with neuroprotective properties. *CNS Neurosci Ther.* 2008 Summer;14(2):95-106. PMID: 18482023.

Salmi E, Kaisti KK, Metsähonkala L, et al. Sevoflurane and propofol increase 11C-flumazenil binding to gamma-aminobutyric acid receptors in humans. *Anesth Analg.* 2004 Nov;99(5):1420-6. PMID: 15502041.

Yamanoue T, Brum JM, Estafanous FG. Vasodilation and mechanism of action of propofol in porcine coronary artery. *Anesthesiology.* 1994 Aug;81(2):443-51. PMID: 8053594.

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