

Phone: 888-558-5227

651-644-8424

Fax: 888-558-7329

Email: getinfo@lktlabs.com Web: lktlabs.com

## **Product Information**

Product ID V0146

CAS No. 137862-53-4

Chemical Name N-(1-Oxopentyl)-N-[[2'-1H-tetrazol-5-yl)[1,1'-biphenyl]-4-yl]methyl]-

L-valine

Svnonvm Diovan, Tareg

Formula  $C_{24}H_{29}N_5O_3$ Formula Wt. 435.52 Melting Point 116-117°C Purity ≥98%

Solubility Solubility in DMSO - 20 mg/mL.

Sparingly soluble in aqueous medium.

N=N

## **Pricing and Availability**

Bulk quanitites available upon request

Product ID	Size	List Price
V0146	50 mg	\$54.60
V0146	1 g	\$63.00
V0146	5 g	\$113.30
V0146	25 a	\$415.10

Store Temp Ambient Ship Temp Ambient

**Description** Valsartan is an angiotensin II (ATII) type 1 receptor (AT1) inhibitor that is clinically used to lower blood pressure and coronary resistance, and to decrease cardiac hypertrophy. Valsartan exhibits antihypertensive, cardioprotective, neuroprotective, antiangiogenic, and anti-inflammatory activities. Inhibition of the ATII type 1 receptor results in inhibition of NF-κB and AP-1 activation. Valsartan induces autophagy through alteration of Akt/mTOR signaling in the rat heart, decreasing infarct size in an ischemia-reperfusion model. In neurons, valsartan promotes spinogenesis, increasing the number of AMPA receptors on the cell surface and altering levels of CaMKIIa and phospho-CDK5. In vivo, valsartan prevents induction of cardiotrophin-1, inhibiting increases in creatine kinase, atrial natriuretic peptide, and the heart weight/body weight ratio during heart failure. Additionally, inhibition of the angiotensin II type 1 (AT1) receptor prevents phosphorylation of Akt and decreases expression of VEGF in bone marrow stromal cells. Valsartan also inhibits release of pro-inflammatory cytokines such as IL-6, TNF-α, and IL-18 independent of its activity on ATII type 1 receptors.

References Wu X, He L, Cai Y, et al. Induction of autophagy contributes to the myocardial protection of valsartan against ischemia reperfusion injury. Mol Med Rep. 2013 Dec;8(6):1824-30. PMID: 24084854.

> Sohn YI, Lee NJ, Chung A, et al. Antihypertensive drug Valsartan promotes dendritic spine density by altering AMPA receptor trafficking, Biochem Biophys Res Commun. 2013 Oct 4;439(4):464-70. PMID: 24012668.

Cheng CI, Hsiao CC, Wu SC, et al. Valsartan impairs angiogenesis of mesenchymal stem cells through Akt pathway. Int J Cardiol. 2013 Sep 10;167(6):2765-74. PMID: 22805546.

Al-Mazroua HA, Al-Rasheed NM, Korashy HM, Downregulation of the cardiotrophin-1 gene expression by valsartan and spironolactone in hypertrophied heart rats in vivo and rat cardiomyocyte H9c2 cell line in vitro: a novel mechanism of cardioprotection. J Cardiovasc Pharmacol. 2013 Apr;61(4):337-44. PMID: 23288202.

Iwashita M, Sakoda H, Kushiyama A, et al. Valsartan, independently of AT1 receptor or PPARy, suppresses LPS-induced macrophage activation and improves insulin resistance in cocultured adipocytes. Am J Physiol Endocrinol Metab. 2012 Feb 1;302 (3):E286-96. PMID: 22045314.

Müller DN, Mervaala EM, Dechend R, et al. Angiotensin II (AT(1)) receptor blockade reduces vascular tissue factor in angiotensin II-induced cardiac vasculopathy. Am J Pathol. 2000 Jul;157(1):111-22. PMID: 10880382.

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