Phone: 888-558-5227

651-644-8424 Email: getinfo@lktlabs.com

Fax: 888-558-7329

Web: lktlabs.com

## **Product Information**

Product ID A5313

Chemical Name  $[1R-[1\alpha[E(S^*)],4ab,5\alpha,6\alpha,8a\alpha]]-3-[2-[Decahydro-6-hydroxy-5-$ 

(hydroxymethyl)-5,8a-dimethyl-2- methylene-1-naphthalenyl]

ethylidene]dihydro-4- hydroxy-2(3H)-furanone

Synonym Andrographis

Formula C<sub>20</sub>H<sub>30</sub>O<sub>5</sub> Formula Wt. 350.45 Melting Point 218-221°C Purity ≥98%

**Solubility** Soluble in methanol,

acetone, chloroform or ether. Slightly soluble in

water.

Store Temp -20°C

Ship Temp Ambient

CAS No. 5508-58-7

## Pricing and Availability

Bulk quanitites available upon request

Product ID	Size	List Price
A5313	10 mg	\$27.80
A5313	100 mg	\$50.00
A5313	250 mg	\$94.40
A5313	1 g	\$305.10

**Description** Andrographolide is a labdane diterpene found in *Andrographis* that exhibits antioxidative, anti-inflammatory, antidiabetic, immunomodulatory, anti-metastatic, and anticancer activities. In vitro, andrographolide decreases TNF-α-induced generation of ROS and expression of ICAM-1 and increases levels of glutathione and heme oxygenase 1 (HO-1). Andrographolide also decreases influenza-induced cell mortality in vitro by inhibiting activation of RIG1-like receptors (RLRs). Additionally, andrographolide suppresses development of diabetes in vivo by decreasing expression of IL-2, IL-17, and IFN-γ and increasing expression of IL-10 and TGF-8. This compound also decreases production of NO, PGE2, iNOS, TNF-α, COX-2, and IFN-β in LPSstimulated macrophages. In leukemia cells, andrographolide inhibits HSP90 activity, decreases Bcr-Abl levels, and induces apoptosis. TEST!!!!!!

References Lu CY, Yang YC, Li CC, et al. Andrographolide inhibits TNFα-induced ICAM-1 expression via suppression of NADPH oxidase activation and induction of HO-1 and GCLM expression through the PI3K/Akt/Nrf2 and PI3K/Akt/AP-1 pathways in human endothelial cells. Biochem Pharmacol. 2014 Sep 1;91(1):40-50. PMID: 24998495.

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