

## alpha-Viniferin Datasheet

4<sup>th</sup> Edition (Revised in July, 2016)

### [ Product Information ]

**Name:** alpha-Viniferin

**Catalog No.:** CFN97068

**Cas No.:** 62218-13-7

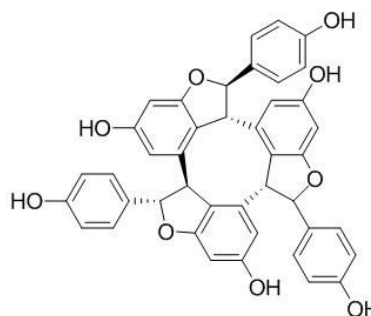
**Purity:** > 98%

**M.F:** C<sub>42</sub>H<sub>30</sub>O<sub>9</sub>

**M.W:** 678.7

**Physical Description:** Powder

**Synonyms:** (2R,2aR,7R,7aR,12S,12aS)-2,7,12-tris(4-hydroxyphenyl)-2,2a,7,7a,12,12a-hexahydrobis[1]benzofuro[3',4':4,5,6;3'',4'':7,8,9]cyclonona[1,2,3-cd][1]benzofuran-4,9,14-triol.



### [ Intended Use ]

1. Reference standards;
2. Pharmacological research;
3. Synthetic precursor compounds;
4. Intermediates & Fine Chemicals;
5. Others.

### [ Source ]

The herb of *Carex humilis* Leyss.

## **[ Biological Activity or Inhibitors ]**

Alpha-viniferin isolated from *Caragana chamlagu* is a trimer of resveratrol, has anti-inflammatory, anti-oxidant, anti-arthritis, and anti-tumor activities; alpha-viniferin strongly inhibits 7 of the 9 CYP isoforms (except CYP2A6 and CYP2E1); alpha-viniferin strongly inhibits CYP2C19-mediated omeprazole 5-hydroxylation and CYP3A4-catalyzed midazolam 1-hydroxylation with IC<sub>50</sub> values of 0.93 and 1.2 μM, respectively.<sup>[1]</sup>

(+)-Alpha-viniferin, a stilbene trimer from *Caragana chamlague*, inhibits acetylcholine -sterase (AChE) activity in a dose-dependent manner, the IC<sub>50</sub> value is 2.0 microM.<sup>[2]</sup>

Alpha-viniferin at doses > 30 mg/kg (p. o.) or > 3 mg/kg (i. v.) shows significant anti-inflammatory activity through inhibition of cyclooxygenase-2 and inducible nitric oxide synthase on carrageenin-induced paw edema in mice; alpha-Viniferin showed an inhibitory effect with an IC<sub>50</sub> value of 4.9 microM on COX-2 activity, but a very weak inhibitory effect at 100 microM on COX-1 activity; alpha-Viniferin inhibited synthesis of iNOS transcript with an IC<sub>50</sub> value of 4.7 microM.<sup>[3]</sup>

Alpha-viniferin is a prostaglandin H<sub>2</sub> synthase inhibitor, the inhibitory potency of (+)-alpha-viniferin is about 3- to 4-fold stronger than that of resveratrol on cyclooxygenase activity of prostaglandin H<sub>2</sub> synthase partially purified from sheep seminal vesicles.<sup>[4]</sup>

## **[ Solvent ]**

Chloroform, Dichloromethane, DMSO, Acetone, etc.

## **[ HPLC Method ]<sup>[5]</sup>**

Mobile phase: Methanol-Acetonitrile- Buffer (pH=4.50)= 16.2 : 12.8 : 71.0 ;

Flow rate: 1.0 ml/min;

Column temperature: 30 °C;

The wave length of determination: 284nm.

## **[ Storage ]**

2-8°C, Protected from air and light, refrigerate or freeze.

## **[ References ]**

- [1] Sim J, Jang H W, Min S, *et al. Food Chem. Toxicol.* , 2014, 69:276-80.
- [2] Sung S H, Kang S Y, Lee K Y, *et al. Biol. Pharm. Bull.*, 2002, 25(1):125-7.
- [3] Chung E Y, Kim B H, Lee M K, *et al. Planta Med.*, 2003, 69(8):710-4.
- [4] Lee S H, Shin N H, Kang S H, *et al. Planta Med.*, 1998, 64(3):204-7.
- [5] Shu N, Zhou H, Hu C. *Biol. Pharm. Bull.*, 2006, 29(4):608-12.

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