

## Faradiol Datasheet

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

### [ Product Information ]

**Name:** Faradiol

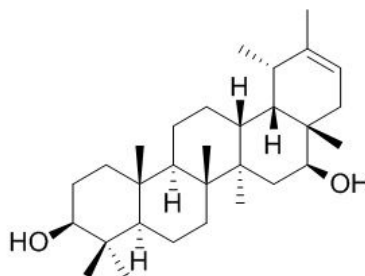
**Catalog No.:** CFN93020

**Cas No.:** 20554-95-4

**Purity:** > 95%

**M.F:** C<sub>30</sub>H<sub>50</sub>O<sub>2</sub>

**M.W:** 442.7



**Physical Description:** Powder

**Synonyms:** (18 $\alpha$ ,19 $\alpha$ )-Urs-20-ene-3 $\beta$ ,16 $\beta$ -diol.

### [ Intended Use ]

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

### [ Source ]

The herbs of *Tussilago farfara*.

### [ Biological Activity or Inhibitors ]

Faradiol shows inhibitory activity against 12-O-tetradecanoylphorbol-13-acetate (TPA)-

induced inflammation in mice; it can inhibit markedly the tumor-promoting effect of TPA (1 ug/mouse) on skin tumor formation following initiation with 7,12-dimethylbenz[ $\alpha$ ]anthracene (50 ug/mouse).<sup>[1]</sup>

Faradiol shows inhibitory effects against Epstein-Barr virus early antigen (EBV-EA) activation with potencies either comparable with or stronger than that of glycyrrhetic acid, a known natural anti-tumor-promoter.<sup>[2]</sup>

## **[ Solvent ]**

Chloroform, Dichloromethane, Ethyl Acetate, DMSO, Acetone, etc.

## **[ HPLC Method ]<sup>[3]</sup>**

Mobile phase: Methanol- Aqueous trifluoroacetic acid(pH4.0),gradient elution ;

Flow rate: 1.5 ml/min;

Column temperature: 30 °C;

The wave length of determination: 210 nm.

## **[ Storage ]**

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

## **[ References ]**

[1] Yasukawa K, Akihisa T, Oinuma H, *et al. Oncology*, 1996, 53(4):341-4.

[2] Ukiya M, Akihisa T, Tokuda H, *et al. Cancer Lett.*, 2002, 177(1):7-12.

[3] Baumann D, Adler S, Grüner S, *et al. Phytochem. Anal.*, 2004, 15(4):226-30.

## **[ Contact ]**

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