

## Isovanillin Datasheet

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

### [ Product Information ]

**Name:** Isovanillin

**Catalog No.:** CFN90358

**Cas No.:** 621-59-0

**Purity:** >= 98%

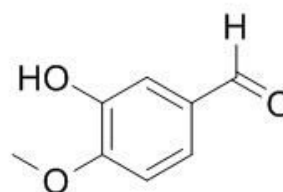
**M.F:** C<sub>8</sub>H<sub>8</sub>O<sub>3</sub>

**M.W:** 152.15

**Physical Description:** Powder

**Synonyms:** 3-Hydroxy-4-methoxybenzaldehyde; 3-Hydroxyanisaldehyde;

5-Formylguaiacol; 3-Hydroxy-p-anisaldehyde.



### [ Intended Use ]

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

### [ Source ]

The barks of *Pinus yunnanensis*.

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

Isovanillin has been proved as an inhibitor of the metabolism of heterocyclic substrates, such as phthalazine, by guinea pig liver aldehyde oxidase.<sup>[1]</sup>

Isovanillin and iso-acetovanillon are two phenolic components isolated from a number of plants including *Pycnocycla spinosa*. *P. spinosa* extract has antispasmodic and antidiarrheal activities; the antidiarrheal effect of *P. spinosa* extract is at least partially due to presence of two active compounds isovanillin and iso-acetovanillon; they are relaxant of ileum contractions induced by 5-HT and EFS and they have contribution to the relaxant effect of *P. spinosa* extract but other components are responsible for the inhibition of acetylcholine by the extract.<sup>[2]</sup>

## **[ Solvent ]**

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

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

Mobile phase: Acetonitrile -0.02% Aqueous phosphoric acid, gradient elution;

Flow rate: 1.0 ml/min;

Column temperature: Room Temperature;

The wave length of determination: 230 nm.

## **[ Storage ]**

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

## **[ References ]**

[1] Brígido C, Da F I, Parreira R, *et al. Acta Biochim. Pol.*, 2004, 51(4):943-51.

[2] Sadraei H, Ghanadian M, Asghari G, *et al. Res. Pharm. Sci.*, 2014, 9(2):187-92.

[3] Tong L, Tan X J, Lin J R, *et al. Chinese Journal of Pharmaceutical Analysis*, 2009, 29(6); 961-3.

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