



# Jatrorrhizine Hydrochloride Datasheet

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

## [ Product Information ]

**Name:** Jatrorrhizine Hydrochloride

**Catalog No.:** CFN98108

**Cas No.:** 6681-15-8

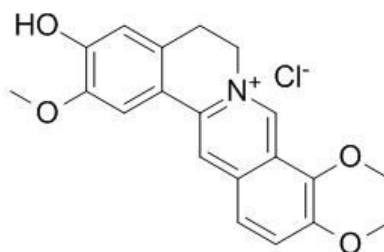
**Purity:** >=98%

**M.F:** C<sub>20</sub>H<sub>20</sub>ClNO<sub>4</sub>

**M.W:** 373.83

**Physical Description:** Powder

**Synonyms:** 3-Hydroxy-2,9,10-trimethoxy-5,6-dihydro-isochino[3,2-a]isochinolinylum, Chloride.



## [ Intended Use ]

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

## [ Source ]

The barks of *Phellodendron chinense* Schneid.

## [ Biological Activity or Inhibitors ]

Jatrorrhizine hydrochloride can inhibit the proliferation and neovascularization of C8161 metastatic melanoma cells with low toxicity, suggests that jatrorrhizine hydrochloride is a potential new antimelanoma drug candidate.<sup>[1]</sup>

Jatrorrhizine hydrochloride exhibits a potent inhibitory effect toward neuraminidase of the H7N9 (N9) avian influenza virus, it also can potentiate the neuraminidase inhibitory effect of oseltamivir towards H7N9 influenza.<sup>[2]</sup>

Jatrorrhizine hydrochloride has lipid lowering effects, it can ameliorate hyperlipidemia via the suppression of lipogenesis and the enhancement of lipid oxidation in the liver. <sup>[3]</sup>

### **[ Solvent ]**

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

### **[ HPLC Method ]<sup>[4]</sup>**

Mobile phase: Acetonitrile-H<sub>2</sub>O(0.34g potassium dihydrogen phosphate and 0.17 g sodium dodecyl sulfate per 100 mL)=45:55;

Flow rate: 1.0 ml/min;

Column temperature: 35 °C;

The wave length of determination: 345 nm.

### **[ Storage ]**

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

### **[ References ]**

[1] Liu R, Cao Z, Pan Y, *et al. Anti-cancer Drug*, 2013, 24(7):667-76.

[2] Wang Y, Yu M, Wang X, *et al. Rsc Adv.*, 2015, 5(80):64937-43.

[3] Yang W, She L, Yu K, *et al. Mol. Med. Rep.*, 2016, 14(4):3277-84.

[4] Fan X X, Wan L Y, Yuan L P, *et al. Chinese Pharmaceutical Journal*, 2006, 41(12):942-4.

## **[ Contact ]**

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