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**KAMPOYAKI NATURAL
PRODUCTS BIO-CHEMISTRY**

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OXYMATRINE

Datasheet

Kampoyaki Novo-Drug Screening Libraries 4th Edition (Revised in July, 2016)

PRODUCT INFORMATION

Name: Oxymatrine

Catalog No.: KRN99805

Cas No.: 16837-52-8

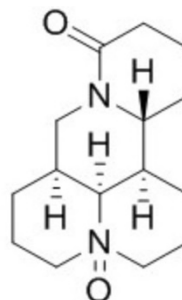
Purity: >=98%

M.F: C₁₅H₂₄N₂O₂

M.W: 264.4

Physical Description: Powder

Synonyms: Ammothamnine; Matrine N-oxide



POTENTIAL USES

1. Reference standards; **2.** Pharmacological research; **3.** Food and cosmetic research;
4. Synthetic precursor compounds; **5.** Active Pharmaceutical Intermediates (API) & Fine Chemicals; **6.** Ingredient in supplements, beverages; **7.** Agricultural research; **8.** Botanical Bio- Allelopathy, **9.** Natural Botanical Molecules as Botanical Bio-Herbicides **10.** As Botanical Bio- Anti-Blight Fungicides

SOURCE

The root of *Sophora flavescens* Ait

BIOLOGICAL ACTIVITY OR INHIBITORS

Oxymatrine, one of the major components of *Sophora flavescens* ait, has exhibited anti-hepatitis virus infection, anti-hepatic fibrosis, anti-inflammation, anti-anaphylaxis and other immune-regulation, it induces human pancreatic cancer PANC-1 cells apoptosis via regulating expression of Bcl-2 and IAP families, and releasing of cytochrome C.[1] Oxymatrine is proven to protect ischemic and reperfusion injury in liver, intestine and heart, this effect is via anti-inflammation and anti-apoptosis, it has protective effect applies to ischemic injury in brain to reduce infarct volume induced by pMCAO, this effect may be through the decreasing of NF-kappaB expression. [2]

Oxymatrine has anti-inflammatory activity, it can reduce the serum levels of TNF- α , IL-6, and the expression of NF- κ B and ICAM-1 in colonic mucosa in dextran sulfate sodium (DSS)--induced colitis of rats, indicates that oxymatrine may ameliorate the colonic inflammation and thus alleviate diarrhea and bloody stool.[3]

Oxymatrine has a beneficial effect on acute lung injury induced by oleic acid in mice and may inhibit the production of proinflammatory cytokine, TNF- α , by means of the inhibition of p38 MAPK.[4]

Oxymatrine has inhibition of hepatitis B virus in vivo, it (200 mg/kg/d,20day)can reduce the contents of HBsAg and HBeAg in transgenic miceliver, and longer treatment time and larger dosage do not yield better effects.[5]

Oxymatrine can attenuate diabetes-associated cognitive deficits in rats, which is associated with oxidative stress, inflammation and apoptotic cascades.[6]

Oxymatrine can induce cell cycle arrest and apoptosis, which makes it a potentially useful agent for targeting cancer cells; it causes a dose-dependent reduction in the proliferation of MCF-7 cells and a decrease in SP cells, the growth inhibitory effects of oxymatrine treatment on MCF-7 cells may be due to the inhibition of SP and Wnt/0205-catenin signaling pathway. [7]

Oxymatrine and astragalus polysaccharide can synergistically improve the immune efficacy of Newcastle disease vaccine in chicken, because of astragalus polysaccharide and oxymatrine possesses synergistical immunoenhancement.[8]

SOLVENT

Pyridine, Methanol, Ethanol, Hot water, etc.

HPLC METHOD (9)

Mobile phase: Methanol-3% Phosphoric acid H₂O=80:20;

Flow rate: 1.0 ml/min;

Column temperature: 30 °C;

The wave length of determination: 220 nm.

STORAGE

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

REFERENCES

- [1] Qi L, Xiao X, Wei X, et al. J. Exp. Clin. Cancer Res., 2011, 30(1):66-66.
[2] Zhang X J, Yang C H, Fan H G. Brain Res., 2009, 1268:174-80.
[3] Zheng P, Niu F L, Liu W Z, et al. World J. Gastroenterol., 2005, 11(31):4912-5.
[4] Lao L, Rao S Y, Gong Z N, et al. J. Ethnopharmacol., 2005, 98(1-2):177-83.
[5] Xiao, Song, Chen, et al. World J. Gastroenterol., 2001, 7(1):49-52.
[6] Suo-bin, WANG, Jian-ping. Acta Pharmacol. Sin., 2014, 35(3):331-8.
[7] Zhang Y, Piao B, Zhang Y, et al. Med. Oncol., 2011, 28(1):99-107.
[8] Chen Y, Wang D, Hu Y, et al. Int. J. Biol. Macromol., 2010, 46(4):425-8.
[9] Liu J, Yuan L M, Li-Qin H E, et al. Chemical World, 2009, 50(7):410-1.



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CERTIFICATE OF ANALYSIS

Name: Oxymatrine
Catalog No.: KRN99805
Cas No.: 16837-52-8
Purity: >= 99.4%
M.F: C₁₅H₂₄N₂O₂

Physical Description: Yellow powder

Solvent: Pyridine, Methanol, Ethanol, etc.

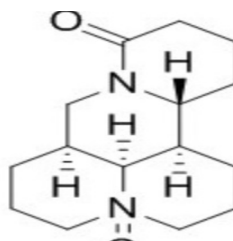
Weight 20mg

Lot No. KRS201801

Storage Protected from air and light, refrigerate or freeze (2-8 °C)

Intended Use For laboratory use only

Shelf Life 2 years



CHARACTERIZATION DATA SUMMARY

Analytical Test

Identification by ¹H-NMR , HPLC
Purity tested

Results

Consistent with the above structure
>= 99.4%



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GHS SAFETY DATA SHEET

Version 4.2

Revision Date 01/01/2018

Print Date 01/08/2019

1. PRODUCT AND COMPANY IDENTIFICATION

GHS Product Name: Oxymatrine

Product code: KRN99805

Company: KAMPOYAKI HERS PTE LTD

Address: 16 New Industrial Road, #05-05 Hudson Techno Centre Singapore 536204

Tel: +65-63833202

Fax: +65-63833632

Website: www.kampoyaki-research.com

E-mail: thiru-sam@kampoyaki-research.com | kampoyak@singnet.com.sg

2. HAZARDS IDENTIFICATION

2.1 GHS classification

Physical Hazards: Not classified

Health Hazards: Not classified

Environmental Hazards: Not classified

2.2 GHS label elements, including precautionary statements

Pictograms or hazard symbols: None

Signal word: No signal word

Hazard statements: None

Precautionary statements: None

3. COMPOSITION/INFORMATION ON INGREDIENTS

Chemical Name: Oxymatrine

CAS#: 16837-52-8

Purity: $\geq 98\%$

Formula: $C_{15}H_{24}N_2O_2$

Molecular Weight: 264.4

Hazard Symbols: ---

Risk Phrases: ---

4. FIRST AID MEASURES

4.1 Description of first aid measures

Eyes: Immediately flush eyes with plenty of water for at least 15 minutes, occasionally lifting the upper and lower eyelids. Consult a doctor.

Skin: Flush skin with plenty of water for at least 15 minutes while removing contaminated clothing and shoes. Consult a doctor.

Ingestion: Do NOT induce vomiting. If conscious and alert, rinse mouth and drink 2-4 cupfuls of milk or water. Consult a doctor.

Inhalation: Remove from exposure and move to fresh air immediately. Consult a doctor.

4.2 Indication of immediate medical attention and special treatment needed

Show this safety data sheet to the doctor in attendance. Immediate medical attention is required.

5. FIRE FIGHTING MEASURES

5.1 Suitable extinguishing

Media: Dry chemical, foam, water spray, carbon dioxide.

Precautions for firefighters: Fire-extinguishing work is done from the windward and the suitable fire-extinguishing method according to the surrounding situation is used. Uninvolved persons should evacuate to a safe place. In case of fire in the surroundings: Remove movable containers if safe to do so.

5.2 Special protective

Equipment for firefighters: When extinguishing fire, be sure to wear personal protective equipment.

6. ACCIDENTAL RELEASE MEASURES

6.1 Personal precautions, protective equipment and emergency procedures

Avoid dust formation. Avoid breathing vapors, mist or gas.

6.2 Environmental precautions

Do not let product enter drains.

6.3 General Information

Use proper personal protective equipment as indicated in Section 8.

6.4 Spills/Leaks

Clean up spills immediately, observing precautions in the Protective Equipment section. Sweep up, then place into a suitable container for disposal. Decontaminate spill site with 10% caustic solution and ventilate area until after disposal is complete

7. HANDLING AND STORAGE

7.1 Precautions for safe handling:

Wash thoroughly after handling. Remove contaminated clothing and wash before reuse. Avoid contact with eyes, skin, and clothing. Avoid ingestion and inhalation. Keep away from sources of ignition. Avoid prolonged or repeated exposure.

7.2 Storage

Store in a well closed container. Protected from air and light, refrigerate or freeze.(2-8°C)

7.3 Specific end uses

Use in a laboratory fume hood where possible. Refer to employer is COSHH risk assessment.

8. EXPOSURE CONTROLS / PERSONAL PROTECTION

8.1 Engineering controls

Use adequate general or local exhaust ventilation to keep airborne concentrations below the permissible exposure limits. Use process enclosure, local exhaust ventilation, or other engineering controls to control airborne levels.

Control parameters: Not set up

8.2 Personal protective equipment

Respiratory protection: Dust respirator. Follow local and national regulations.

Hand protection: Protective gloves.

Eye protection: Wear safety glasses and chemical goggles if splashing is possible.

Skin and body protection: Wear appropriate protective gloves and clothing to prevent skin exposure.

9. PHYSICAL AND CHEMICAL PROPERTIES

- a) Appearance Yellow powder
- b) Odour no data available
- c) Odour Threshold no data available
- d) pH no data available
- e) Melting point/freezing point no data available
- f) Initial boiling point and boiling range no data available
- g) Flash point no data available
- h) Evaporation rate no data available
- i) Flammability (solid, gas) no data available
- j) Flammability or explosive limits no data available
- k) Vapour pressure no data available
- l) Vapour density
- m) Relative density no data available
- n) Water solubility no data available
- o) Partition coefficient: no data available
- p) Autoignition temperature no data available
- q) Decomposition temperature no data available
- r) Viscosity no data available
- s) Explosive properties no data available
- t) Oxidizing properties no data available

10 - STABILITY AND REACTIVITY

10.1 Reactivity

Stable under recommended transport or storage conditions.

10.2 Chemical Stability

Stable under normal temperatures and pressures.

10.3 Conditions to Avoid

Incompatible materials, strong oxidants, heat.

10.4 Incompatibilities with Other Materials

Strong oxidising/reducing agents, strong acids/alkalis.

10.5 Hazardous Decomposition Products

Nitrogen oxides, carbon monoxide, irritating and toxic fumes and gases, carbon dioxide, nitrogen.

10.6 Hazardous Polymerization

Has not been reported.

11. TOXICOLOGICAL INFORMATION

| | |
|---------------------------------------|-------------------|
| Acute Toxicity: | No data available |
| Skin corrosion/irritation: | No data available |
| Serious eye damage/irritation: | No data available |
| Germ cell mutagenicity: | No data available |
| Carcinogenicity: | --- |
| IARC: | No data available |
| NTP: | No data available |
| Reproductive toxicity: | No data available |

12. ECOLOGICAL INFORMATION

| | |
|--|--|
| Toxicity: | No data available |
| Persistence and degradability: | No data available |
| Bioaccumulative potential: | No data available |
| Mobility in soil: | No data available |
| Results of PBT and vPvB assessment: | No data available |
| Other adverse effects: | May be harmful to the aquatic environment. |

13. DISPOSAL CONSIDERATIONS

Dispose of in a manner consistent with federal, state, and local regulations.

14. TRANSPORT INFORMATION

14.1 Hazards Class:

Does not meet the criteria for classification as hazardous for transport

14.2 UN proper shipping name

ADR/RID: Not dangerous goods

IMDG: Not dangerous goods

IATA: Not dangerous goods

14.3 Transport hazard class(es)

Does not meet the criteria for classification as hazardous for transport.

15. REGULATORY INFORMATION

15.1 Safety, health and environmental regulations/legislation specific for the substance or mixture

No data available

15.2 Chemical Safety Assessment

No data available

16. ADDITIONAL INFORMATION

This GHS SDS above is believed to be accurate and represents the best information currently available to us. However, we make no warranty of merchantability or any other warranty, express or implied, with respect to such information, and we assume no liability resulting from its use. Users should make their own investigations to determine the suitability of the information for their particular purposes. In no way shall the company be liable for any claims, losses, or damages of any third party or for lost profits or any special, indirect, incidental, consequential or exemplary damages, howsoever arising, even if the company has been advised of the possibility of such damages.

End of GHS safety data sheet



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