

OPIUM- raw opium gum pellet

Remedy Makers

Disclaimer: This homeopathic product has not been evaluated by the Food and Drug Administration for safety or efficacy. FDA is not aware of scientific evidence to support homeopathy as effective.

CII

WARNING:

"The FDA has not determined that this product is safe, effective and not misbranded for its intended use".

INDICATION:

VERTIGO, DIARRHEA, PAIN OR OTHER INDICATIONS

WARNING:

Use only if cap and seal are unbroken. Keep this and all medication out of reach of children. If symptoms persist more than 3 days or worsen, discontinue (STOP) use and consult your physician. As with any drug. If you are pregnant or nursing (breast-feeding) a baby, seek the advice of a health professional before using this product. Store tightly closed in a cool area.

Directions (adult/children)

Dissolve 3 or 4 pellets in mouth under tongue 3 times a day or as directed by a physician. Children 2 years and older take 1/2 adult dose.

Inactive Ingredients:

Lactose and Sucrose. Free from yeast, wheat, corn, and soy.

Questions or comments:

(877)REM4YOU Fax (909) 594-4205 Pomona, CA. 91768 USA www.remedy-makers.com

Other Information:

Contains approx. 160 - 164 pellets.

CAUTION:

Federal law prohibits dispensing without prescription.

REMEDY MAKERS™



R_x
ONLY

OPIUM

6X_H

(Raw opium Gum)

1 gram of pellets contain approx. 0.1mg Opium

VERTIGO, DIARRHEA, PAIN,
OR OTHER INDICATIONS

2 Drams (1/4 ounce) Lot # XXX-XXX-X Exp.N/A

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Drug Facts: Active Ingredient Listed above. To be used according to standard homeopathic indications for self-limiting conditions such as those indicated above or as directed by a physician.

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NDC 10191-1082-2

HOMEOPATHIC
MEDICINE



OPIUM

raw opium gum pellet

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:10191-1082
Route of Administration	SUBLINGUAL	DEA Schedule	CII

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
OPIUM (UNII: 37M3MZ001L) (OPIUM - UNII:37M3MZ001L)	OPIUM	6 [hp_X]

Inactive Ingredients

Ingredient Name	Strength
SUCROSE (UNII: C151H8M554)	
LACTOSE (UNII: J2B2A4N98G)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:10191-1082-2	160 in 1 VIAL, GLASS; Type 0: Not a Combination Product	07/07/2015	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		07/06/2015	

Labeler - Remedy Makers (018543582)

Revised: 3/2023

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