

GELSEMIUM SEMPERVIRENS - gelsemium sempervirens pellet
HOMEOLAB USA INC

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.

HOMEOPATHIC MEDICINE NDC 60512-1020-1

ACTIVE INGREDIENT HPUS

GELSEMIUM SEMPERVIRENS 3X

(Yellow jessamine)

APPREHENSION

USE

For self-limiting condition listed above or as directed by a health professional.

WARNINGS

Enter section text here

Do not use if pellet-dispenser seal is broken.

Stop use and ask a doctor if symptoms persist more than 3 days or worsen.

If pregnant or breast-feeding, ask a health professional before use.

Keep out of reach of children.

DIRECTIONS

(Adults / Children 2-18 years): Allow 3 or 4 pellets to dissolve in the mouth 3 times a day until symptoms are relieved or as directed by a health professional.

OTHER INFORMATION

Store at room temperature.

INACTIVE INGREDIENTS

Lactose, sucrose.

QUESTIONS?

1-800-404-4666

The letters HPUS indicate the ingredient is officially included in the Homeopathic Pharmacopoeia of the United States.

80 Pellets

Pellet dispenser

HOMEOLAB USA INC., 3025 De L'Assomption, Montreal, QC, H1N 2H2, CANADA

LABEL

HOMEOPATHIC MEDICINE

GELSEMIUM
SEMPERVIRENS

Yellow Jessamine

NDC 60512-1020-1

APPREHENSION

80 Pellets

Pellet dispenser

HOMEOLAB USA INC., 3025 De L'Assomption, Montreal, QC, H1N 2H2 CANADA



HOMEOLAB

usa

Product of Canada

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Break seal, turn & twist.

A-C

GELSEMIUM SEMPERVIRENS

gelsemium sempervirens pellet

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:60512-1020
Route of Administration	ORAL		

Active Ingredient/Active Moiety				
Ingredient Name			Basis of Strength	Strength
GELSEMIUM SEMPERVIRENS ROOT (UNII: 639KR60Q1Q) (GELSEMIUM SEMPERVIRENS ROOT - UNII:639KR60Q1Q)			GELSEMIUM SEMPERVIRENS ROOT	3 [hp_X]
Inactive Ingredients				
Ingredient Name			Strength	
LACTOSE (UNII: J2B2A4N98G)				
SUCROSE (UNII: C15IH8M554)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:60512-1020-1	80 in 1 TUBE		
Marketing Information				
Marketing Category	Application Number or Monograph Citation		Marketing Start Date	Marketing End Date
unapproved homeopathic			10/11/1995	

Labeler - HOMEOLAB USA INC (202032533)

Establishment			
Name	Address	ID/FEI	Business Operations
HOMEOLAB USA INC		202032533	manufacture

Revised: 11/2011

HOMEOLAB USA INC