GELSEMIUM SEMPERVIRENS- gelsemium semp tablet
STANDARD HOMEOPATHIC COMPANY

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.

GELSEMIUM SEMP. 30X

Made according to the
Homeopathic Pharmacopoeia
of the United States since 1903.

Warnings

Do not use if cap band is missing or broken.

If you are pregnant or nursing, consult a licensed health care professional before using this product.

If symptoms persist for 7 days or worsen, contact a licensed practitioner.

To be used according to label indications and/or standard homeopathic indications

Keep this and all medicines out of the reach of children.

Standard Homeopathic Company
Los Angeles, CA 90061
Questions? 800-624-9659

Directions

Adults: Dissolve 4 tablets under tongue 4 times a day.
Children: Consult a healthcare professional

In a base of Acacia Gum and Lactose N.F.

GELSEMIUM SEMP. 30X

DULL HEADACHE OR FLU

Principal Display Panel -- 250 Tablet Bottle Label

Standard

Homeopathic

Made in the USA Since 1903

NDC # 0360-0182-01

Homeopathic Medicine

GELSEMIUM SEMP.

30X

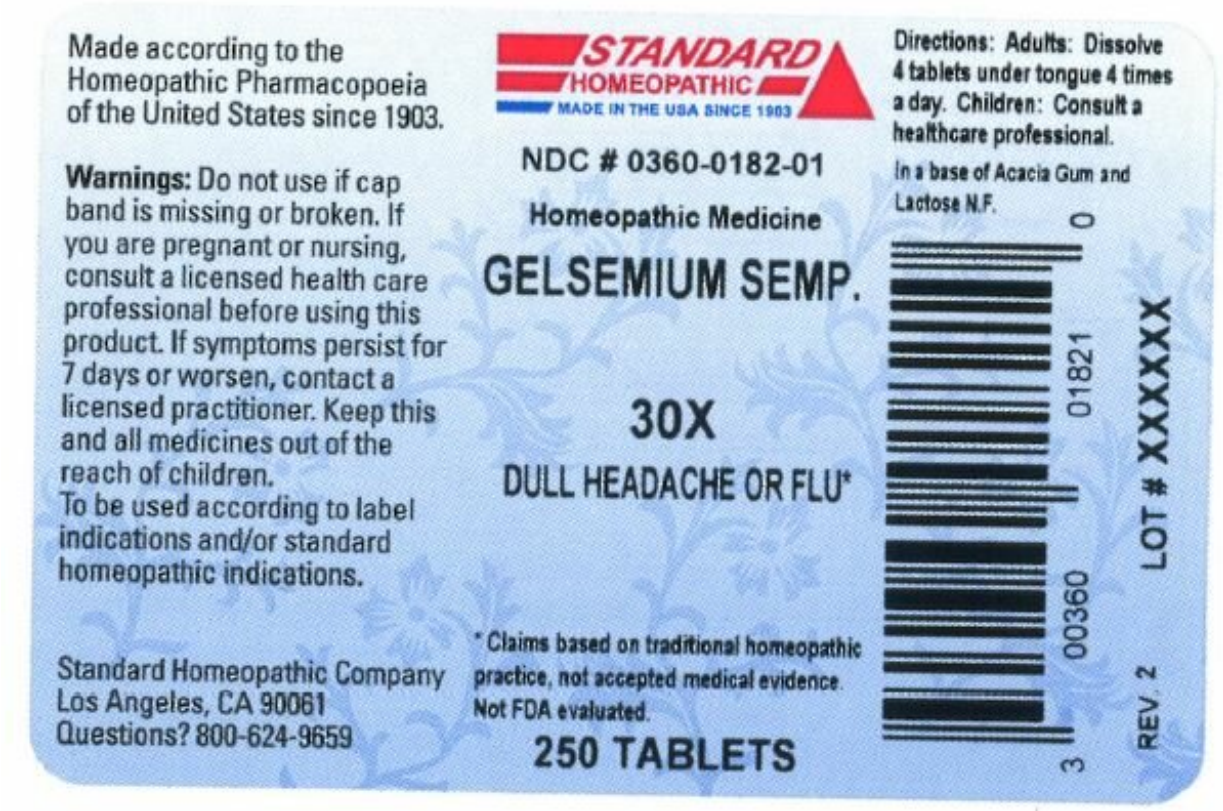
DULL HEADACHE OR FLU*

*Claims based on traditional homeopathic

practice, not accepted medical evidence.

Not FDA evaluated.

250 TABLETS



GELSEMIUM SEMPERVIRENS

gelsemium semp tablet

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:0360-0182
Route of Administration	SUBLINGUAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
GELSEMIUM SEMPERVIRENS ROOT (UNII: 639KR60Q1Q) (GELSEMIUM SEMPERVIRENS ROOT - UNII:639KR60Q1Q)	GELSEMIUM SEMPERVIRENS ROOT	30 [hp_X] in 1 g

Inactive Ingredients	
Ingredient Name	**Strength**
ACACIA (UNII: 5C5403N26O)	
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)	

Product Characteristics			
Color	white	**Score**	score with uneven pieces
Shape	ROUND	**Size**	5mm
Flavor		**Imprint Code**	

Contains	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0360-0182-01	1 g in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	02/17/1997	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		02/17/1997	

Labeler - STANDARD HOMEOPATHIC COMPANY (008316655)

Establishment

Name	Address	ID/FEI	Business Operations
STANDARD HOMEOPATHIC COMPANY		008316655	manufacture(0360-0182) , pack(0360-0182) , label(0360-0182)

Revised: 12/2018 STANDARD HOMEOPATHIC COMPANY