GELSEMIUM SEMP- gelsemium sempervirens root tablet
Hyland's 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.

GELSEMIUM SEMP 30X

DULL HEADACHE & FLU

GELSEMIUM SEMP. 30X

DULL HEADACHE & FLU

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.

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

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

Directions

Adults: Dissolve 4 tablets under tongue 4 times a day.

Children: consult a healthcare professional.

Inactive Ingredients

Acacia Gum and Lactose

Questions?

800-624-9659

PRINCIPAL DISPLAY PANEL - 250 Tablet Bottle Label

SINCE 1903
Hyland's ®
HOMEOPATHIC

NDC 54973-2913-4

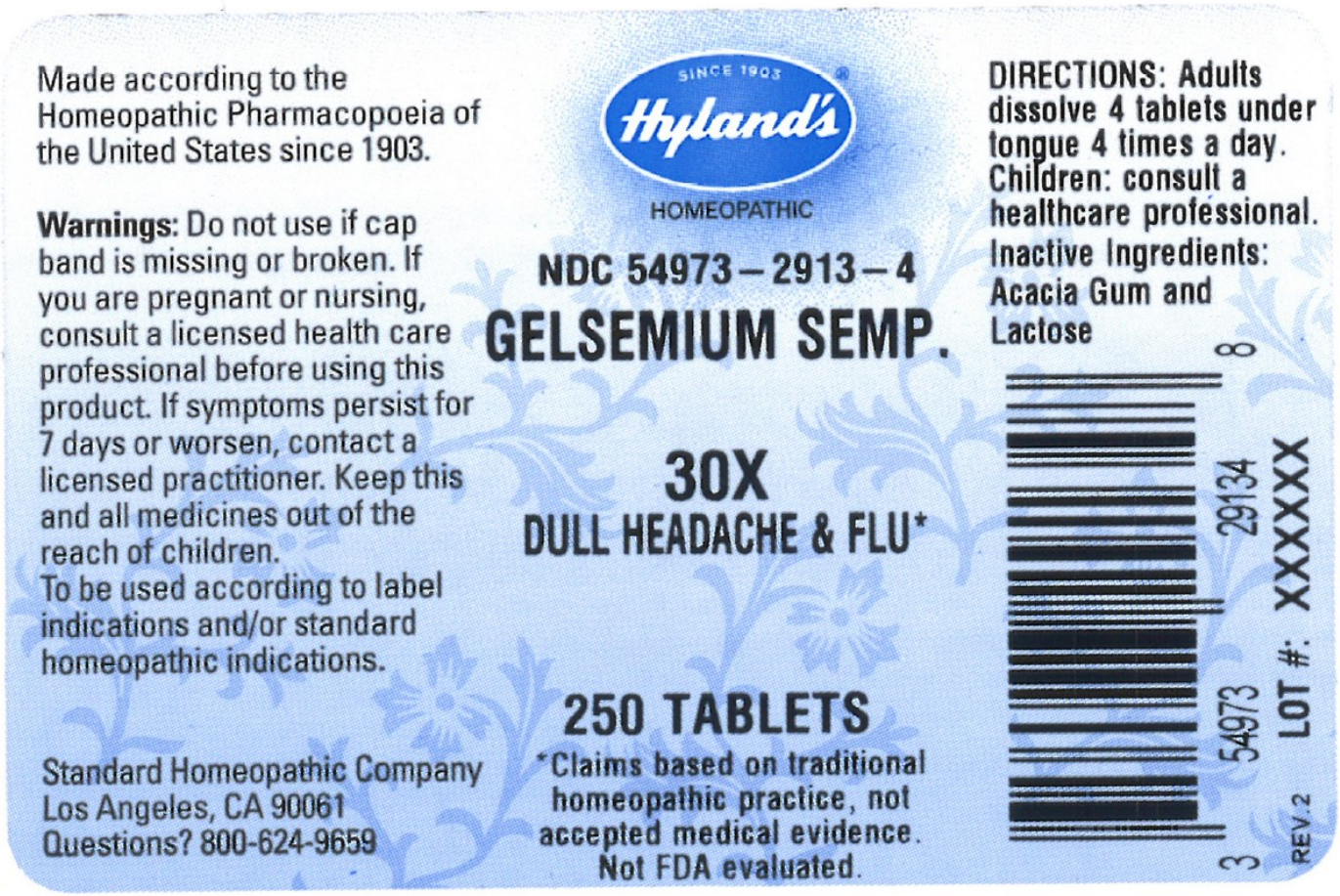
GELSEMIUM SEMP.
30X

DULL HEADACHE & FLU*

250 TABLETS

***Claims are based on traditional homeopathic practice, not accepted medical evidence.**

Not FDA evaluated.



GELSEMIUM SEMP

gelsemium sempervirens root tablet

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:54973-2913
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	30 [hp_X]

Inactive Ingredients

Ingredient Name	Strength
ACACIA (UNII: 5C5403N26O)	
LACTOSE (UNII: J2B2A4N98G)	

Product Characteristics

Color	yellow	**Score**	no score
Shape	ROUND	**Size**	9mm
Flavor		**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:54973-2913-1	250 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	01/01/1955	12/20/2017
2	NDC:54973-2913-4	250 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	01/01/1955	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		01/01/1955	

Labeler - Hyland's Inc. (008316655)

Establishment

Name	Address	ID/FEI	Business Operations
Hyland's Inc.		008316655	manufacture(54973-2913) , pack(54973-2913) , label(54973-2913)

Revised: 12/2022 Hyland's Inc.