

G-22- ligustrum vulgare whole solution
DNA Labs, 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.

G-22

NDC 58264-0385-2

INDICATIONS

Heavy menses.

INGREDIENTS

ACTIVE

Ligustrum vulgare (Privet) 1DH

INACTIVE

36.6% alcohol (V/V) Glycerin Macerate

SUGGESTED DOSAGE

One dropper full twice daily.

SHAKE WELL

Warnings

- Use only if cap seal is unbroken.
- If pregnant or breastfeeding, ask a healthcare professional before use.
- Keep this and all medication out of the reach of children.

To be used according to standard homeopathic indications.

PRINCIPAL DISPLAY PANEL - 2 FL. OZ. Bottle Label

DYNAMIC
NUTRITIONAL
ASSOCIATES, INC.

G-22

LIGUSTRUM

VULGARE

GEMMOTHERAPY

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Rev. 8/18



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MANUFACTURED FOR:
DNA LABORATORIES, INC.
Coeur d'Alene, ID 83814
800-426-7112

G-22

ligustrum vulgare whole solution

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:58264-0385
Route of Administration	SUBLINGUAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
LIGUSTRUM VULGARE WHOLE (UNII: FC604DJ98E) (LIGUSTRUM VULGARE WHOLE - UNII:FC604DJ98E)	LIGUSTRUM VULGARE WHOLE	1 [hp_X] in 1 mL

Inactive Ingredients

Ingredient Name	Strength
ALCOHOL (UNII: 3K9958V90M)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:58264-0385-2	59.14 mL in 1 BOTTLE, GLASS; Type 0: Not a Combination Product	01/01/1990	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		01/01/1990	

Labeler - DNA Labs, Inc. (031784339)

Revised: 6/2022

DNA Labs, Inc.