G-41- ulmus glabra 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-41

NDC 58264-0404-2

INDICATIONS

Support for the joints and skin.

INGREDIENTS

ACTIVE

Ulmus campestris (Elm) 1DH

INACTIVE

36.6% alcohol (V/V) Glycerin Macerate

Rx CAUTION

Federal law prohibits dispensing without a prescription.

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-41

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GEMMOTHERAPY

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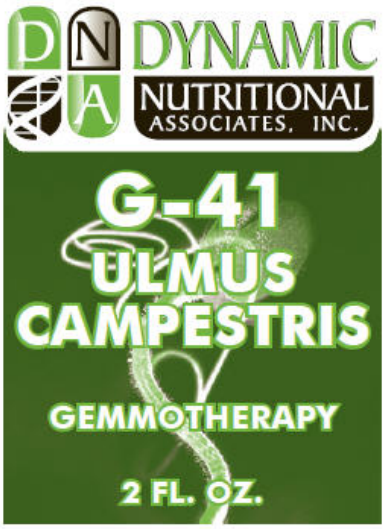
Rev. 8/18



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

G-41
ulmus glabra whole solution

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:58264-0404
Route of Administration	SUBLINGUAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ULMUS GLABRA WHOLE (UNII: I2AHJ0JD5V) (ULMUS GLABRA WHOLE - UNII:I2AHJ0JD5V)	ULMUS GLABRA 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:58264-0404-2	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.