

**G-59- populus nigra leaf bud and cornus sanguinea 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-59

NDC 58264-0422-2

INDICATIONS

Congestion, stagnation, inflammation.

Energy Meridian

Heart Master-Triple Heater

INGREDIENTS

ACTIVE

Populus nigra (Poplar) 1DH, Cornus sanguinea (Hound's Tree) 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-59

HEART-SMALL
INTESTINE

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-59

populus nigra leaf bud and cornus sanguinea whole solution

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:58264-0422
Route of Administration	SUBLINGUAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
POPULUS NIGRA LEAF BUD (UNII: GJU5F9OXMT) (POPULUS NIGRA LEAF BUD - UNII:GJU5F9OXMT)	POPULUS NIGRA LEAF BUD	1 [hp_X] in 1 mL
CORNUS SANGUINEA WHOLE (UNII: PQF087FG15) (CORNUS SANGUINEA WHOLE - UNII:PQF087FG15)	CORNUS SANGUINEA 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:58264-0422-2	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: 1/2025

DNA Labs, Inc.