

T-3- fagus sylvatica 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.

T-3

NDC 58264-0248-1

INDICATIONS

Intolerance, criticism, passing judgments.

INGREDIENTS

ACTIVE

Fagus sylvatica 6/8/30x

INACTIVE

20% alcohol in purified water.

SUGGESTED DOSAGE

One dropper under tongue two times daily. Acute symptoms ½ dropper under tongue every 30 minutes for two hours.

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 - 1 FL. OZ. Bottle Label

DYNAMIC

NUTRITIONAL

ASSOCIATES, INC.

T-3

BEECH

FLOWER ESSENCES

1 FL. OZ

SUGGESTED DOSAGE:

One dropper under tongue two times daily. Acute symptoms 1/2 dropper under tongue every 30 minutes for two hours.

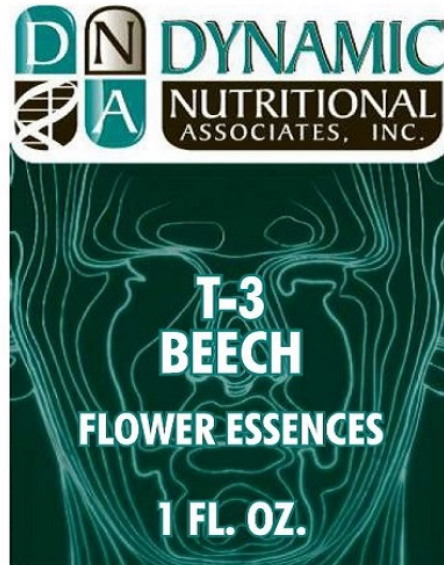
SHAKE WELL

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.

Rev. 4/22



NDC 58264-0248-1

INDICATIONS:

Intolerance, criticism, passing judgments.

INGREDIENTS:

ACTIVE: Fagus sylvatica 6/8/30x

INACTIVE: 20% alcohol in purified water.

Mfg for: **DNA LABORATORIES, INC.**
Chelan, WA 98816

T-3

fagus sylvatica whole solution

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:58264-0248
Route of Administration	SUBLINGUAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
FAGUS SYLVATICA WHOLE (UNII: SW8UOC4GS) (FAGUS SYLVATICA WHOLE - UNII:SW8UOC4GS)	FAGUS SYLVATICA WHOLE	30 [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-	29.57 mL in 1 BOTTLE, GLASS; Type 0: Not a	01/01/2000	

0248-1	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.