

T-22- quercus robur 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-22

NDC 58264-0267-1

INDICATIONS

Despondency, despair, but never ceasing effort.

INGREDIENTS

ACTIVE

Quercus robur 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-22

OAK

FLOWER ESSENCES

1 FL. OZ.

SUGGESTED DOSAGE:

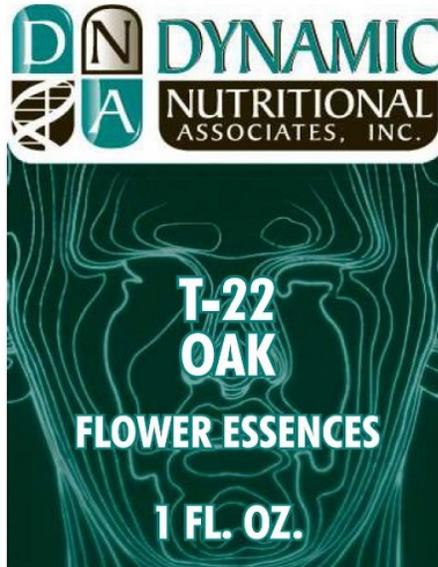
One dropper under tongue two times daily. Acute symptoms ½ dropper under tongue every 30 minutes for two hours.

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.

Rev. 4/22



NDC 58264-0267-1

INDICATIONS:

Despondency, despair, but never ceasing effort.

INGREDIENTS:

ACTIVE: Quercus robur 6/8/30x

INACTIVE: 20% alcohol in purified water.

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

T-22

quercus robur whole solution

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:58264-0267
Route of Administration	SUBLINGUAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
QUERCUS ROBUR WHOLE (UNII: R7QMG0BT2W) (QUERCUS ROBUR WHOLE - UNII:R7QMG0BT2W)	QUERCUS ROBUR 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:58264-0267-1	29.57 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.