

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.

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Rev. 4/22



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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: 5/2022

DNA Labs, Inc.