

T-19- larix decidua 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-19

NDC 58264-0264-1

INDICATIONS

Lack of confidence, anticipation of failure, despondency.

INGREDIENTS

Larix decidua 6/8/30x, 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.
- As with drugs if you are pregnant or nursing a baby seek professional advice before using this product.
- 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-19

LARCH

FLOWER ESSENCES

1 FL. OZ.

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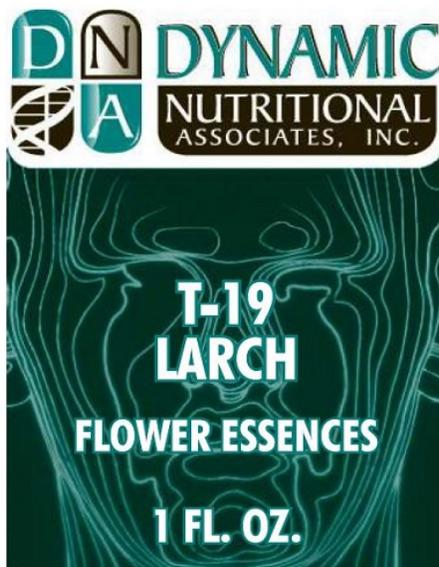
SHAKE WELL

Warnings:

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- If pregnant or breastfeeding, ask a healthcare professional before use.
- Keep this and all medication out of the reach of children.

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Rev. 4/22



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INDICATIONS:

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INGREDIENTS:

ACTIVE: Larix decidua 6/8/30x

INACTIVE: 20% alcohol in purified water.

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

T-19

larix decidua whole solution

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:58264-0264
Route of Administration	SUBLINGUAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
LARIX DECIDUA WHOLE (UNII: O8554H2U9W) (LARIX DECIDUA WHOLE - UNII:O8554H2U9W)	LARIX DECIDUA WHOLE	30 [hp_X] in 1 mL

Inactive Ingredients

Ingredient Name	Strength
ALCOHOL (UNII: 3K9958V90M)	
WATER (UNII: 059QF0K00R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:58264-0264-1	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.