# T-24- pinus sylvestris 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.

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### T-24

NDC 58264-0269-1

### INDICATIONS

Self-reproach, guilt feelings, despondency.

### INGREDIENTS

### ACTIVE

Pinus sylvestris 6/8/30x

### INACTIVE

20% alcohol in purified water.

### SUGGESTED DOSAGE

One dropper under tongue two times daily. Acute symptoms  $\frac{1}{2}$  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.

# PRINCIPAL DISPLAY PANEL - 1 FL. OZ. Bottle Label

## DYNAMIC

### NUTRITIONAL

**ASSOCIATES, INC.** 

### T-24

# PINE 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.



#### NDC 58264-0269-1

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INGREDIENTS: ACTIVE: Pinus sylvestris 6/8/30x INACTIVE: 20% alcohol in purified water.

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

Rev. 4/22

## **T-24**

pinus sylvestris whole solution

Product Info	rmation									
Product Type		HUMAN OTC DRUG	Item Code (Source)			NDC:58264-0269				
Route of Admin	istration	SUBLINGUAL								
Active Ingredient/Active Moiety										
Ingredient Name					Basis of Strength		Strength			
PINUS SYLVESTRIS WHOLE (UNII: VP00HZL9IC) (PINUS SYLVESTRIS WHOLE - UNII: VP00HZL9IC)				PINUS SYLVESTRIS WHOLE		30 [hp_X] in 1 mL				
Inactive Ingr	edients									
Ingredient Name					Strength					
ALCOHOL (UNII: 3	K9958V90M)									
WATER (UNII: 0590	QF0KO0R)									
Packaging										
# Item Code	Pa	ackage Description	M	Marketing Start Ma Date		Mar	rketing End Date			
		• •		Dat	e		Date			

1 NDC:58264- 0269-1	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.