T-13- ulex europaeus 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-13

NDC 58264-0258-1

INDICATIONS

Hopelessness, despair.

INGREDIENTS

ACTIVE

Ulex europaeus 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-13

GORSE

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 58264-0258-1

INDICATIONS:

Hopelessness, despair.

INGREDIENTS:

ACTIVE: Ulex europaeus 6/8/30x INACTIVE: 20% alcohol in purified water.

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

T-13

ulex europaeus whole solution

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:58264-0258

Route of Administration SUBLINGUAL

Active Ingredient/Active Moiety		
Ingredient Name	Basis of Strength	Strength
ULEX EUROPAEUS WHOLE (UNII: 3349V12Y8N) (ULEX EUROPAEUS WHOLE - UNII: 3349V12Y8N)	ULEX EUROPAEUS WHOLE	30 [hp_X] in 1 mL

Inactive Ingredients	
Ingredient Name	Strength
ALCOHOL (UNII: 3K9958V90M)	
WATER (UNII: 059QF0KO0R)	

ı	Packaging			
	# Item Code	Package Description	Marketing Start Date	Marketing End Date

1 NDC:58264- 0258-1	29.57 mL in 1 BOTTLE, GLASS; Type 0: Not a Combination Product	01/01/1990	
Marketing Information			
	 		
Marketing Category	Application Number or Monograph	n Marketing Start Date	Marketing End Date
Marketing	Application Number or Monograph	_	_

Labeler - DNA Labs, Inc. (031784339)

Revised: 5/2022 DNA Labs, Inc.