T-34- hottonia palustris 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-34

NDC 58264-0279-1

INDICATIONS

Pride, aloofness.

INGREDIENTS

ACTIVE

Hottonia palustris 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-34 WATER VIOLET **FLOWER ESSENCES** 1 FL. OZ.

SUGGESTED DOSAGE:

One dropper under tongue two times daily. Acute symptoms 1/2 dropper under tongue every 30 minutes for two hours.

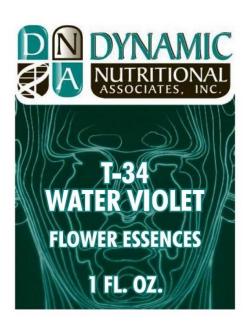
SHAKE WELL

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.

Rev. 4/22



NDC 58264-0279-1

INDICATIONS:

Pride, aloofness.

INGREDIENTS:

ACTIVE: Hottonia palustris 6/8/30x **INACTIVE**: 20% alcohol in purified water.

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

T-34

hottonia palustris whole solution

Product Information

Product Type HUMAN OTC DRUG **Item Code (Source)** NDC:58264-0279

Route of Administration SUBLINGUAL

Active Ingredient/Active Moiety

Basis of Ingredient Name Strength Strength

HOTTONIA PALUSTRIS WHOLE (UNII: GS042K765G) (HOTTONIA PALUSTRIS WHOLE - UNII:GS042K765G)

HOTTONIA PALUSTRIS 30 [hp_X] WHOLE in 1 mL

Inactive Ingredients

Ingredient Name Strength ALCOHOL (UNII: 3K9958V90M)

WATER (UNII: 0590F0KO0R)

Packaging

# Item Code	Package Description	Marketing Start Date	Marketing End Date
NDC F03C4	20 F7 and in 1 POTTLE CLASS. Trans 0. Not a		

NDC:58264- 29.57 mL in 1 BOTTLE, GLASS; Type 0: Not a

0279-1	Combination Product	01/01/1320			
Marketing Information					
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date		
Category	Citation	Date	Date		
unapproved homeopathic	Citation	01/01/1990	Dute		

Labeler - DNA Labs, Inc. (031784339)

Revised: 5/2022 DNA Labs, Inc.