T-10- malus pumila 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-10

NDC 58264-0255-1

INDICATIONS

The cleansing remedy, despondency, despair.

INGREDIENTS

ACTIVE

Malus pumila 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-10 CRAB APPLE 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.

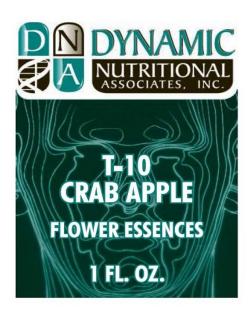
SHAKE WELL

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 58264-0255-1

INDICATIONS:

The cleansing remedy, despondency, despoir.

INGREDIENTS:

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

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

T-10

malus pumila whole solution

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:58264-0255

Route of Administration SUBLINGUAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
MILA WHOLE (UNII: 04W636S1V3) (MALUS PUMILA WHOLE -	MALUS PUMILA	30 [hp X]

MALUS PUMILA WHOLE (UNII: 04W636S1V3) (MALUS PUMILA WHOLE - MALUS PUMILA WHOLE - UNII:04W636S1V3) WHOLE 30 [hp_X] in 1 mL

Inactive Ingredients

Ingredient Name	Strength
ALCOHOL (UNII: 3K9958V90M)	

WATER (UNII: 059QF0KO0R)

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" Date Date	# Item Code			Marketing End Date
	NDC:58264-	29.57 mL in 1 BOTTLE, GLASS; Type 0: Not a	01/01/1000	

0255-1	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)

Establishment			
Name	Address	ID/FEI	Business Operations
RNA Bio/Pharma, Inc		883081523	manufacture(58264-0255)

Revised: 5/2022 DNA Labs, Inc.