T-9- clematis vitalba 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-9

NDC 58264-0254-1

INDICATIONS

Indifference, dreaminess, inattention, unconsciousness.

INGREDIENTS

ACTIVE

Clematis vitalba 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.

CLEMATIS FLOWER ESSENCES 1 FL. OZ.

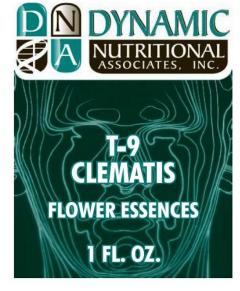
SUGGESTED DOSAGE:

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.



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

Т-9									
clematis vitalba whole	e solutior	า							
Product Informa	tion								
Product Type		HUMAN OTC DRUG	Item Code	m Code (Source)			NDC:58264-0254		
Route of Administra	ation	SUBLINGUAL							
Active Ingredient	/Active	Moiety							
						Basis d	of		
Ingredient Name						Streng	Strength		
CLEMATIS VITALBA WHOLE (UNII: 7PH07Z124Q) (CLEMATIS VITALBA WHOLE - UNII: 7PH07Z124Q)								30 [hp_X] in 1 mL	
Inactive Ingredie	nts								
Ingredient Name						Strength			
ALCOHOL (UNII: 3K9958	8V90M)								
WATER (UNII: 059QF0KC	00R)								
Packaging					_				
# Item Code	Pa	ackage Description		Mark	Date) Start	Mar	keting End Date	

Rev. 4/22

	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					
nomeoputine							

Labeler - DNA Labs, Inc. (031784339)

Revised: 5/2022

DNA Labs, Inc.