EMVITA 7- anacard occ, lachesis mutus, phosphorus, glandula sup, lycopodium liquid
RUBIMED AG

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.

Emvita 7

Drug Facts

Active Ingredients: (HPUS*) 20% of each

Anacard occ 18LM Glandula sup 21X

Lachesis mutus 800C Lycopodium 16LM

Phosphorus 21X

*The letters "HPUS" indicate that the components in this product are officially monographed in the Homeopathic Pharmacopoeia of the United States.

†Claims based on traditional homeopathic practice, not accepted medical evidence. Not FDA evaluated.

Uses: (†) Homeopathic remedy for show of strength.

Warnings:

Stop use if symptoms persist or worsen.

If you are pregnant or breastfeeding,

consult a health care professional prior to use.

Keep out of reach of children.

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Directions: (adults & children 6 years & older)

Take 5 drops 3 - 6 times daily, or as recommended by your health care professional.

Other information: Store at 20 - 25ºC (68 - 77ºF). Do not use if box has been tampered with, or if safety seal of the

bottle is broken.

Inactive ingredients: Ethanol 20% USP, Purified Water.

Manufactured by: OHM pharma, Inc., USA.
Distributed by: Privia Naturals, LLC.
197 Woodland Pkwy Suite 104 #813
San Marcos, CA 92069
www.privianaturals.com
1 (888) 526-9695 Product of USA.

NDC 66343-056-50

RUBIMED

Emvita 7

Homeopathic Medicine For

Show of Strength

1.7 fl oz. 50 mL 20% Ethanol

Homeopathic Medicine For

Show of Strength

EMVITA 7 anacard occ, lachesis mutus, phosphorus, glandula sup, lycopodium liquid			
Product Information			
Product Type	HUMAN PRESCRIPTION DRUG	**Item Code (Source)**	NDC:66343-056

Route of Administration	ORAL

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ANACARDIUM OCCIDENTALE FRUIT (UNII: 4A10JR4E7E) (ANACARDIUM OCCIDENTALE FRUIT - UNII:4A10JR4E7E)	ANACARDIUM OCCIDENTALE FRUIT	18 [hp_M] in 50 mL
LACHESIS MUTA VENOM (UNII: VSW71SS07I) (LACHESIS MUTA VENOM - UNII:VSW71SS07I)	LACHESIS MUTA VENOM	800 [hp_C] in 50 mL
PHOSPHORUS (UNII: 27YLU75U4W) (PHOSPHORUS - UNII:27YLU75U4W)	PHOSPHORUS	21 [hp_X] in 50 mL
SUS SCROFA ADRENAL GLAND (UNII: 398IYQ16YV) (SUS SCROFA ADRENAL GLAND - UNII:398IYQ16YV)	SUS SCROFA ADRENAL GLAND	21 [hp_X] in 50 mL
LYCOPODIUM CLAVATUM SPORE (UNII: C88X29Y479) (LYCOPODIUM CLAVATUM SPORE - UNII:C88X29Y479)	LYCOPODIUM CLAVATUM SPORE	16 [hp_M] in 50 mL

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

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:66343-056-50	50 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/03/2019	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		06/03/2019	

Labeler - RUBIMED AG (480582035)

Revised: 1/2021 RUBIMED AG