

EMVITA 20- strychnos ignatii seed, apis mellifera, phosphorus, pulsatilla vulgaris, and lytta vesicatoria 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 20

Homeopathic preparation for general well-being: False pride

Ingredients

Belladonna 800 C
Ignatia amara 16 LM
Apis mellifica 18 LM
Magnesia carb. 21 x
Phosphorus 21 x
Pulsatilla 21 x
Cantharis 21 x
Hypophysis 21 x
Alcohol 43 % v/v

TAMPER-EVIDENT

Do not use this product if seal at base of cap is missing or broken.

Directions

10 drops per day or as instructed by your health care provider.

Warnings

As with any drug, if you are pregnant or nursing a baby, seek the advice of a health professional before using the product.

Keep out of Reach of Children!

PRINCIPAL DISPLAY PANEL - 50 ml Bottle Label

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Homeopathic preparation
for general well-being:
False pride

RUBIMED®

1.7 Fl.oz. (50 ml)

NDC 57227-0102-1

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Made in Switzerland for Rubimed AG
 Grossmatt 3
 CH 6052 Hergiswil
 Switzerland

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 Lot No

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Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:66343-032
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
STRYCHNOS IGNATII SEED (UNII: 1NM3M2487K) (STRYCHNOS IGNATII SEED - UNII:1NM3M2487K)	STRYCHNOS IGNATII SEED	16 [hp_M] in 1 mL
APIS MELLIFERA (UNII: 7S82P3R43Z) (APIS MELLIFERA - UNII:7S82P3R43Z)	APIS MELLIFERA	18 [hp_M] in 1 mL
PHOSPHORUS (UNII: 27YLU75U4W) (PHOSPHORUS - UNII:27YLU75U4W)	PHOSPHORUS	21 [hp_X] in 1 mL
PULSATILLA VULGARIS (UNII: I76KB35JEV) (PULSATILLA VULGARIS - UNII:I76KB35JEV)	PULSATILLA VULGARIS	21 [hp_X] in 1 mL
LYTTA VESICATORIA (UNII: 3Q034RO3BT) (LYTTA VESICATORIA - UNII:3Q034RO3BT)	LYTTA VESICATORIA	21 [hp_X] in 1 mL

Inactive Ingredients

Ingredient Name	Strength
Alcohol (UNII: 3K9958V90M)	0.43 mL in 1 mL

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:66343-032-50	50 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
UNAPPROVED HOMEOPATHIC		05/07/2015	

Labeler - RUBIMED AG (480582035)

Establishment

Name	Address	ID/FEI	Business Operations
RUBIMED AG		480582035	MANUFACTURE(66343-032)

Establishment

Name	Address	ID/FEI	Business Operations
Omidia AG		483268348	MANUFACTURE(66343-032)

Revised: 5/2015

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