

**EMVITA 4- apis mellifica, oophorinum, platinum met, orchitinum, petroleum,
stramonium 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 4

Drug Facts

Active Ingredients: (HPUS*) 16.7% of each

Apis mellifica 21X Orchitinum 21X

Oophorinum 21X Petroleum 16LM

Platinum met 800C Stramonium 18LM

*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 extreme self-control.

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 - 25°C (68 - 77°F). 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 66343-053-50

RUBIMED

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Emvita 4

Homeopathic Medicine For Extreme Self-Control

1.7 fl oz. 50 mL 20% Ethanol

PRIVIA NATURALS

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**Homeopathic Medicine For
Extreme Self-Control**

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Extreme Self-Control**

EMVITA 4

apis mellifica, oophorinum, platinum met, orchitinum, petroleum, stramonium liquid

Product Information			
Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:66343-053
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
APIS MELLIFERA (UNII: 7S82P3R43Z) (APIS MELLIFERA - UNII:7S82P3R43Z)	APIS MELLIFERA	21 [hp_X] in 50 mL
SUS SCROFA OVARY (UNII: S7YTV04R8O) (SUS SCROFA OVARY - UNII:S7YTV04R8O)	SUS SCROFA OVARY	21 [hp_X] in 50 mL
PLATINUM (UNII: 49DFR088MY) (PLATINUM - UNII:49DFR088MY)	PLATINUM	800 [hp_C] in 50 mL
SUS SCROFA TESTICLE (UNII: KM02613O28) (SUS SCROFA TESTICLE - UNII:KM02613O28)	SUS SCROFA TESTICLE	21 [hp_X] in 50 mL
KEROSENE (UNII: 1C89KKC04E) (KEROSENE - UNII:1C89KKC04E)	KEROSENE	16 [hp_M] in 50 mL
DATURA STRAMONIUM (UNII: G6W4F0V8Z3) (DATURA STRAMONIUM - UNII:G6W4F0V8Z3)	DATURA STRAMONIUM	18 [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:66343-053-50	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)**Registrant** - OHM PHARMA INC. (030572478)**Establishment**

Name	Address	ID/FEI	Business Operations
OHM PHARMA INC.		030572478	manufacture(66343-053)

Revised: 6/2019

RUBIMED AG