

CHAVITA 4- aur met, crataegus, cetraria, eucalyptus globulus, ferrum met, thymus gland, zinc val 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.

Chavita 4

Drug Facts Active Ingredients: (HPUS*) 14.3% of each

Aur met 21X Crataegus 21X
Cetraria 21X Eucalyptus globulus 21X
Ferrum met 50C Thymus gland 21X
Zinc val 200C

*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 hay fever and allergies.

Warnings:

Stop use if symptoms persist or worsen.

If you are pregnant or breastfeeding, consult a healthcare professional prior to use.

Keep out of reach of children.

Directions: (adults & children 6 years & older)

Take 12 drops 2 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 the 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.
9169 W State St #196
Garden City, ID 83714
www.privianaturals.com
Info@privianaturals.com Product of USA.

NDC 66343-085-60

RUBIMED

Chavita 4

Homeopathic Medicine For
Hay Fever and Allergies

2.0 fl oz. 59mL 20% Ethanol

Drug Facts

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UPC-A: 658877924747
Mag: 83%
Height: 0.5 in.

Chavita™ 4



NDC 66343-085-60

RUBIMED

Chavita™ 4

Homeopathic Medicine For Hayfever and Allergies



TM

2.0 fl oz. 59mL
20% Ethanol by Volume

Drug Facts (continued)

Inactive ingredients:
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Chavita™ 4

Homeopathic Medicine For Hayfever and Allergies



TM

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PR10005



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Hay Fever And Allergies

2.0 fl oz. 59 mL 20% Ethanol



PRIVIA
NATURALS



Drug Facts (continued)

Inactive ingredients: Ethanol 20% USP, Purified Water.

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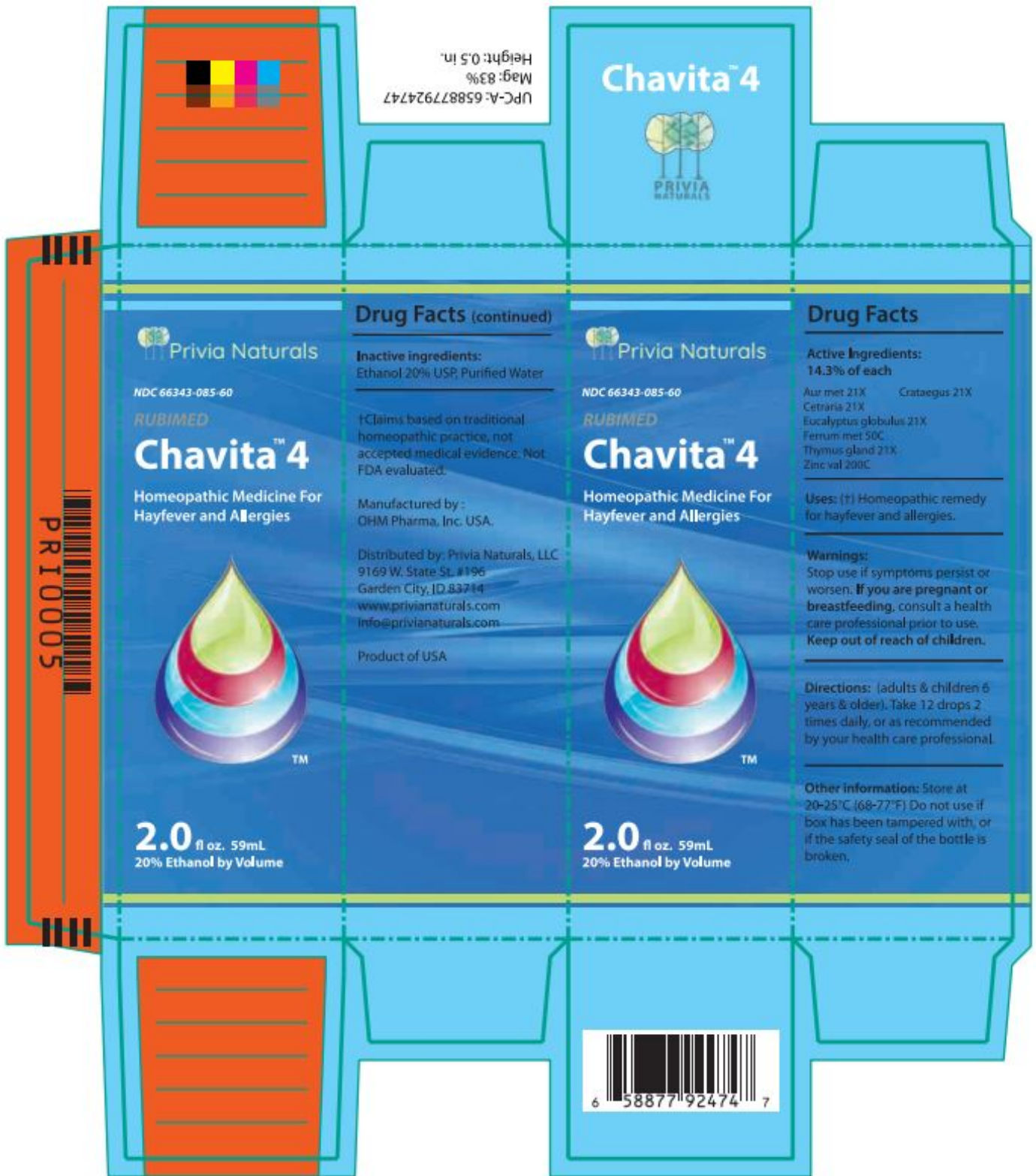
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Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:66343-085
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
GOLD (UNII: 79Y1949PYO) (GOLD - UNII:79Y1949PYO)	GOLD	21 [hp_X] in 60 mL
HAWTHORN LEAF WITH FLOWER (UNII: 6OM09RPY36) (HAWTHORN LEAF WITH FLOWER - UNII:6OM09RPY36)	HAWTHORN LEAF WITH FLOWER	21 [hp_X] in 60 mL
CETRARIA ISLANDICA WHOLE (UNII: BJ7YPN79A1) (CETRARIA ISLANDICA WHOLE - UNII:BJ7YPN79A1)	CETRARIA ISLANDICA WHOLE	21 [hp_X] in 60 mL
EUCALYPTUS GLOBULUS LEAF (UNII: S546YLW6E6) (EUCALYPTUS GLOBULUS LEAF - UNII:S546YLW6E6)	EUCALYPTUS GLOBULUS LEAF	21 [hp_X] in 60 mL
IRON (UNII: E1UOL152H7) (IRON - UNII:E1UOL152H7)	IRON	50 [hp_C] in 60 mL
BOS TAURUS THYMUS (UNII: 8XEJ88V2T8) (BOS TAURUS THYMUS - UNII:8XEJ88V2T8)	BOS TAURUS THYMUS	21 [hp_X] in 60 mL
ZINC VALERATE DIHYDRATE (UNII: MN0RX54EQA) (VALERIC ACID - UNII:GZK92PJM7B)	ZINC VALERATE DIHYDRATE	200 [hp_C] in 60 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:66343-085-60	60 mL in 1 BOTTLE; Type 0: Not a Combination Product	12/19/2022	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		12/01/2022	

Labeler - RUBIMED AG (480582035)