SUN HEADACHE RHEUMATISM- kalmia latifolia leaf pellet Natural Health Supply

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.

Indications

Indications: To be used for acute self-limiting conditions according to standard homeopathic indications

Active Ingredient

KALMIA LATIFOLIA

Directions

Take at onset of symptoms. Repeat every 2 hours or as needed until relieved. If condition persists or worsens discontinue use and consult a practitioner.

Adults: dissolve 5-10 pellets in 1 oz. of filtered water or take dry by mouth. Children and infants: 1-5 pellets.

Keep Out of Reach of Children

Keep these and all medications out of the reach of children.

Warning

If pregnant or nursing, consult a practitioner before using.

Inactive Ingredients

Inactive Ingredients - Lactose, Sucrose

Purpose

Purpose: Sun Headache Rheumatism

Package Label

Est. No. 1724217

Natural Health Supply

505-474-9175

Homeopathic Medicated Pellets

Do not use if clear bottle seal is broken

KALMIA LATIFOLIA 30C

Sun Headache Rheumatism

Lot#

Natural Health Supply 505-474-9175

Homeopathic Medicated Pellets

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7.5ML(1/40Z)LOT #

KALMIA LATIFOLIA

Wheadache & Rheumatis

Est. No. 1724217 Natural Health Supply 505-474-9175

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1.875 ml (1/16 oz.)

LOT #

Do not use if perforated cap seal is broken.

KALMIA LATIFOLIA

30C

Sun Headache & Rheumatism

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SUN HEADACHE RHEUMATISM

kalmia latifolia leaf pellet

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:64117-155

Route of Administration ORAL

Active Ingredient/Active Moiety

Ingredient NameBasis of StrengthStrengthKALMIA LATIFOLIA LEAF (UNII: 79N6542N18) (KALMIA LATIFOLIA LEAF -KALMIA LATIFOLIA30 [hp_C]

UNII:79 N6542N18)

KALMIA LATIFOLIA 30 [hp_C] in 1 mL

Inactive Ingredients

Ingredient Name Strength
LACTOSE (UNII: J2B2A4N98G)

SUCROSE (UNII: C151H8M554)

Packaging

Item Code Package Description Marketing Start Date Marketing End Date

NDC:64117-155- 1.875 mL in 1 BOTTLE, GLASS; Type 0: Not a Combination

01	Product	00/04/1330	
2 NDC:64117-155- 02	7.5 mL in 1 BOTTLE, GLASS; Type 0: Not a Combination Product	06/04/1998	
Marketing I	nformation		
Marketing I		Marketing Start Date	Marketing End Date
•	gory Application Number or Monograph Citation	Marketing Start Date 06/04/1998	Marketing End Date

Labeler - Natural Health Supply (018504618)

Registrant - Natural Health Supply (018504618)

Establishment						
Name	Address	ID/FEI	Business Operations			
Natural Health Supply		0 18 50 46 18	manufacture(64117-155)			

Revised: 1/2019 Natural Health Supply