BELLADONNA- belladonna gel Rxhomeo Private Limited d.b.a. Rxhomeo, Inc

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.

ACTIVE INGREDIENT

BELLADONNA GEL TINC HPUS

USES

Boils, abscesses and dryness of skin, swelling from insect stings

INDICATIONS

Condition listed above or as directed by the physician

DOSAGE & ADMINISTRATION SECTION

Apply a thin layer of Gel to the affected area, repeat 3 times a day or as needed.

WARNINGS

This product is to be used for self-limiting conditions

If symptoms do not improve in 4 days, or worsen, discontinue use and seek assistance of health professional

As with any drug, if you are preganant, or nursing a baby, seek professional advice before taking this product

Keep this and all medication out of reach of children

Do not use if capseal is broken or missing. Close the cap tightly after use.

INACTIVE INGREDIENTS

ALOE VERA GEL

STORAGE

Store in a cool dark place

QUESTIONS OR COMMENTS

www.Rxhomeo.com | 1.888.2796642 | info@rxhomeo.com

Rxhomeo, Inc 9415 Burnet Road, Suite 312, Austin, TX 78758









BELLADONNA

belladonna gel

Product Information

Product TypeHUMAN OTC DRUGItem Code (Source)NDC:15631-2307

Route of Administration TOPICAL

Active Ingredient/Active Moiety

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Ingredient Name	Basis of Strength	Strength			
ATROPA BELLADONNA (UNII: WQZ3G9PF0H) (ATROPA BELLADONNA - UNII:WQZ3G9PF0H)	ATROPA BELLADONNA	1 [hp_X] in 1 g			

Inactive Ingredients

Ingredient Name Strength

ALOE VERA LEAF (UNII: ZY81Z83H0X)

Packaging# Item CodePackage DescriptionMarketing Start DateMarketing End Date1 NDC:15631-2307-01 in 1 CONTAINER03/30/20162 NDC:15631-2307-11 in 1 CONTAINER; Type 0: Not a Combination Product03/30/20162 NDC:15631-2307-21 00 g in 1 CONTAINER; Type 0: Not a Combination Product03/30/20163 NDC:15631-2307-21 in 1 CONTAINER03/30/20163 NDC:15631-2307-3200 g in 1 CONTAINER; Type 0: Not a Combination Product03/30/2016

Marketing Information						
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date			
unapproved homeopathic		03/30/2016				

500 g in 1 CONTAINER; Type 0: Not a Combination Product

Labeler - Rxhomeo Private Limited d.b.a. Rxhomeo, Inc (650833994)

Establishment							
Name	Address	ID/FEI	Business Operations				
Rxhomeo Private Limited d.b.a. Rxhomeo, Inc		650833994	manufacture(15631-2307), label(15631-2307)				

Revised: 3/2016 Rxhomeo Private Limited d.b.a. Rxhomeo, Inc