# BELLADONNA 200C- atropa belladonna liquid Natural Creations, 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.

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#### **BELLADONNA 200C**

## **ACTIVE INGREDIENTS (HPUS\*):**

Belladonna 200C

**USES:** Temporarily relieves high fever.\*\*

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**DIRECTIONS:** Adults & children over 12 years: 10 drops orally 3 times daily, or as

directed by a health care professional.

**KEEP OUT OF THE REACH OF CHILDREN.** In case of overdose (or accidental ingestion) get medical help or contact a Poison Control Center right away.

### **WARNINGS:**

- Consult a physician for use in children under 12 years of age.
- IF PREGNANT OR BREAST-FEEDING, ask a health care professional before use.
- **KEEP OUT OF THE REACH OF CHILDREN.** In case of overdose (or accidental ingestion) get medical help or contact a Poison Control Center right away.
- Do not use if **TAMPER EVIDENT** seal is broken or missing.

INACTIVE INGREDIENTS: Ethyl Alcohol USP, Purified Water.

## **QUESTIONS & COMMENTS?**

Natural Creations, Inc. / Woodbine, IA 51579 / 712-647-1600

\*The letters "HPUS" indicate the component in this product is officially monographed in the Homeopathic Pharmacopeia of the United States.

\*\*These statements have not been reviewed by the FDA. They are based on traditional homeopathic practice.

NDC: 43406-0324-1

**BELLADONNA 200C** 

**HOMEOPATHIC** 

1 fl oz (30 mL) / 20% Alcohol



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INACTIVE INGREDIENTS: Purified Water, Ethyl Alcohol USP.

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LOT: HL1703033

LOT: NC170332

## **BELLADONNA 200C**

atropa belladonna liquid

### **Product Information**

HUMAN OTC DRUG NDC:43406-0324 **Product Type Item Code (Source)** 

**Route of Administration** ORAL

## **Active Ingredient/Active Moiety**

**Basis of Ingredient Name** Strength Strength ATROPA BELLADONNA (UNII: WQZ3G9PF0H) (ATROPA BELLADONNA -ATROPA 200 [hp C] **BELLADONNA** UNII:WQZ3G9PF0H) in 1 mL

## **Inactive Ingredients**

**Ingredient Name** Strength ALCOHOL (UNII: 3K9958V90M) WATER (UNII: 059QF0KO0R)

## **Packaging**

# Item Code	Package Description	Marketing Start Date	Marketing End Date
<b>1</b> NDC:43406-0324-1	30 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product	05/29/2007	

## **Marketing Information**

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		05/29/2007	

## Labeler - Natural Creations, Inc. (018022074)

Establishment								
Name	Address	ID/FEI	Business Operations					
OHM Pharma, Inc.		030572478	manufacture(43406-0324)					

Revised: 12/2022 Natural Creations, Inc.