# QUALITY CHOICE CALAMINE- calamine 8% and zinc oxide 8% lotion Chain Drug Market Association

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### **Quality Choice Calamine Lotion, USP**

**Drug Facts** 

### **Active Ingredients**

Calamine 8% and Zinc Oxide 8%

### **Purpose**

Skin protectant

#### Uses

dries the oozing and weeping o poison ivy, poison oak, and poison sumac.

#### Warnings

For external use only. Use only as directed.

Avoid contact with eyes and mucous membranes.

Ask a doctor before using on chilren 6 months of age.

### When using this product

Discontinue use if condition worsens or does not improve within 7 days and consult a doctor.

### Keep out of reach of children

In case of accidental ingestion, seek professional assistance or contact a Poison Control center immediately.

#### **Directions**

Adults and chidren 2 years of age and older: shake well before using. Cleanse the skin with soap and water and let it dry befroe each use. Apply lotion to the affected area using a cotton or soft cloth, as often as needed for comfort.

Children under 6 months of age: Consult a doctor before use.

#### Other information

Store at room temperature 13-30C (50-86F)

### **Inactive ingredients**

Bentonite magma, calcium hydroxide, glycerin, purified water.

### **Package Principal Display Panel**



### **OUALITY CHOICE CALAMINE**

calamine 8% and zinc oxide 8% lotion				
Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source) NDC:63868-317		NDC:63868-317
Route of Administration	TOPICAL			
Active Ingredient/Active Moiety				
Ingredient Name			Basis of Strengt	h Strength
ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC CATION - UNII:13S1S8SF37)			ZINC CATION	160 mg in 1 mL

Inactive Ingredients	
Ingredient Name	Strength

CALCIUM HYDROXIDE (UNII: PF5DZW74VN)			
GLYCERIN (UNII: PDC6A3C0OX)			
WATER (UNII: 059QF0KO0R)			

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:63868- 317-06	177 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	12/12/2017	

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC Monograph Drug	M016	03/25/1998		

## Labeler - Chain Drug Market Association (011920774)

# Registrant - Pharma Nobis, LLC (118564114)

Establishment			
Name	Address		Business Operations
Pharma Nobis, LLC		118564114	label(63868-317), manufacture(63868-317), analysis(63868-317), pack(63868-317)

Revised: 12/2023 Chain Drug Market Association