

**POISON IVY WASH- pramoxine hcl lotion**  
**Chain Drug Market Association**

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**QC Poison Ivy Wash**

Pramoxine HCl 1%

External Analgesic

For temporary relief of pain and itching associated with poison ivy, poison oak, and poison sumac.

For external use only.

When using this product:

Avoid Contact with eyes.

Do not leave on skin longer than 3 minutes

Rinse thoroughly after application

When using this product avoid contact with the eyes. Do not leave on skin longer than 3 minutes. Rinse thoroughly after application.

Stop use and ask a doctor if condition worsens, or if symptoms persist for more than 7 days or clear up and occur again within a few days.

Adults and children 2 years of age and older: wet the affected area, apply product to affected skin and surrounding area. work foam into lather and rub for up to 3 minutes if needed. do not leave on skin for longer than 3 minutes. Thoroughly rinse product from all areas. Apply to affected are not more than 3 to 4 times daily.

Water

ammonium lauryl sulfare

distearyl phtalic acid amide

glycol distereate

cocamide MIPA

Propylene glycol

Diazolidyn Urea

Methylparaben

Propylparaben

Glycerin

Jojoba esters

disodium edta

sodium hydroxide

nonoxynol-9

Keep out of reach of children. In case of accidental ingestion, seek professional assistance or contact a poison control center immediately.

| Drug Facts   |                                      |
|--|--------------------------------------|
| <b>Active ingredient</b><br>Pramoxine HCl 1% .....   | <b>Purpose</b><br>External analgesic |
| <b>Uses</b> For temporary relief of pain and itching associated with • poison ivy • poison oak • poison sumac  |                                      |
| <b>Warnings</b> For external use only  |                                      |
| When using this product: • Avoid contact with eyes • Do not leave on skin longer than 3 minutes • Rinse thoroughly after application   |                                      |
| Stop use and ask a doctor if condition worsens, or if symptoms persist for more than 7 days, or clear up and occur again within a few days.  |                                      |
| Keep out of reach of children. In case of accidental ingestion, seek professional assistance or contact a Poison Control Center immediately.   |                                      |
| <b>Directions</b>  |                                      |
| Adults and children 2 years of age and older: Wet the affected area • Apply product to affected skin and surrounding area • Work foam into a lather and rub for up to 3 minutes, if needed |                                      |
| Continued ▶  |                                      |



NDC 63868-877-06

# Poison Ivy Wash

## Poison Ivy Cleanser

Removes Urushiol from the Skin  
Unique 2-in-1 Formula  
Dual Action with Jojoba  
Helps Relieve Pain and Itch Caused by Poison Ivy, Poison Oak, Insect Bites and Minor Skin Irritations



**6 fl oz (177mL)**

| Drug Facts (continued)   |  |
|--|--|
| Do not leave on skin for longer than 3 minutes • Thoroughly rinse product from all areas • Apply to affected area not more than 3 to 4 times daily   |  |
| Children under 2 years of age: Consult a doctor  |  |
| <b>Other information</b> For best results, use near a shower or sink where it is easy to thoroughly rinse off the product.   |  |
| <b>Inactive ingredients</b> • Water, Ammonium Lauryl Sulfate, Distearyl Phthalic Acid Amide, Glycol Distearate, Cocamide MIPA, Propylene Glycol (and) Diazolidinyl Urea (and) Methylparaben (and) Propylparaben, Glycerin, Jojoba Esters, Disodium EDTA, Sodium Hydroxide, Nonoxynol-9 |  |
| Questions or Comments? 1-800-662-3435  |  |

Removes urushiol (Poison Ivy Oil) from the skin. For best results use as soon as possible after contact with poison ivy is suspected.



Distributed by C.D.M.A., Inc.®  
4357 W 9 Mile Rd  
Novi, MI 48375  
www.qualitychoice.com  
Questions: 248-443-9300  
**MADE IN USA**



## POISON IVY WASH

pramoxine hcl lotion

### Product Information

|                                |                |                           |               |
|--------------------------------|----------------|---------------------------|---------------|
| <b>Product Type</b>            | HUMAN OTC DRUG | <b>Item Code (Source)</b> | NDC:63868-877 |
| <b>Route of Administration</b> | TOPICAL        |                           |               |

### Active Ingredient/Active Moiety

| Ingredient Name   | Basis of Strength          | Strength         |
|---|----------------------------|------------------|
| <b>PRAMOXINE HYDROCHLORIDE</b> (UNII: 88AYB867L5) (PRAMOXINE - UNII:068X84E056) | PRAMOXINE<br>HYDROCHLORIDE | 10 mg<br>in 1 mL |

### Inactive Ingredients

| Ingredient Name  | Strength |
|--|----------|
| <b>GLYCOL DISTEARATE</b> (UNII: 13W7MDN21W)                    |          |
| <b>PROPYLENE GLYCOL</b> (UNII: 6DC9Q167V3)                     |          |
| <b>DIAZOLIDINYL UREA</b> (UNII: H5RIZ3MPW4)                    |          |
| <b>METHYLPARABEN</b> (UNII: A2I8C7HI9T)                        |          |
| <b>WATER</b> (UNII: 059QF0KO0R)                                |          |
| <b>AMMONIUM LAURYL SULFATE</b> (UNII: Q7AO2R1M0B)              |          |
| <b>HYDROLYZED JOJOBA ESTERS (ACID FORM)</b> (UNII: UDR641JW8W) |          |
| <b>EDETATE DISODIUM</b> (UNII: 7FLD91C86K)                     |          |
| <b>NONOXYNOL-9</b> (UNII: 48Q180SH9T)                          |          |
| <b>PROPYLPARABEN</b> (UNII: Z8IX2SC1OH)                        |          |

**GLYCERIN** (UNII: PDC6A3C0OX)

**SODIUM HYDROXIDE** (UNII: 55X04QC32I)

### Packaging

| # | Item Code        | Package Description                                      | Marketing Start Date | Marketing End Date |
|---|------------------|--|----------------------|--------------------|
| 1 | NDC:63868-877-06 | 177 mL in 1 CONTAINER; Type 0: Not a Combination Product | 04/24/2019           |                    |

### Marketing Information

| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|--------------------|--|----------------------|--------------------|
| OTC Monograph Drug | M017                                     | 04/24/2019           |                    |

**Labeler** - Chain Drug Market Association (011920774)

**Registrant** - Pharma Nobis, LLC (118564114)

### Establishment

| Name              | Address | ID/FEI    | Business Operations   |
|-------------------|---------|-----------|---|
| Pharma Nobis, LLC |         | 118564114 | manufacture(63868-877) , pack(63868-877) , analysis(63868-877) , label(63868-877) |

Revised: 12/2023

Chain Drug Market Association