POISON IVY WASH- pramoxine hcl lotion
Chain Drug Market Association

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 worses, 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 form 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

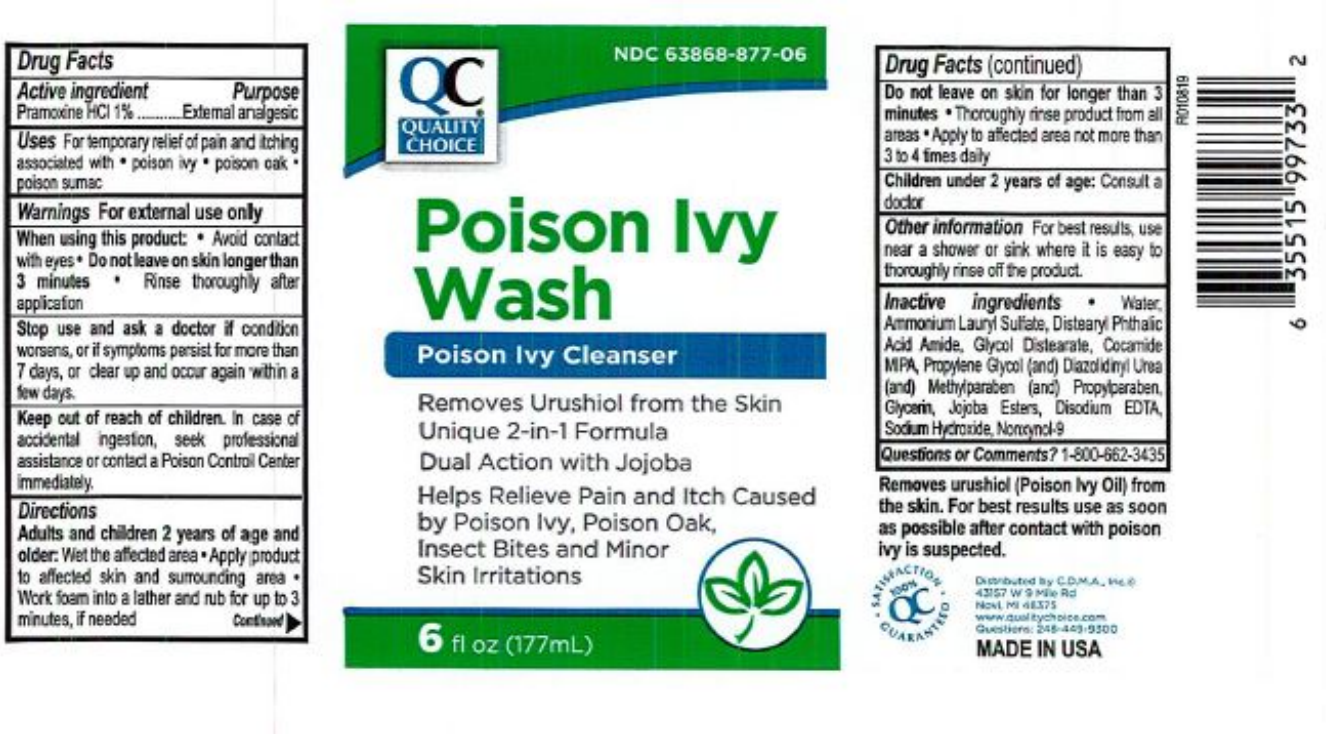
Jojoba esters

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.



POISON IVY WASH

pramoxine hcl lotion

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:63868-877
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
PRAMOXINE HYDROCHLORIDE (UNII: 88AYB867L5) (PRAMOXINE - UNII:068X84E056)	PRAMOXINE HYDROCHLORIDE	10 mg in 1 mL

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