

PROCURE DRYDROX- aluminum hydroxide gel 2% ointment
Twin Med LLC

Active Ingredient

Aluminum Hydroxide Gel 2%

Purpose

Skin Protectant

Uses

Uses

- Dries the oozing and weeping of poison ivy, oak and sumac

Directions

Directions

- Apply as needed
- Children under 6 months of age: ask a doctor

Inactive Ingredients

Calcium Carbonate, Cholecalciferol, Citric Acid, Ethylhexylglycerin, Glyceryl Monostearate, Glyceryl Stearate/ PEG-100 Stearate, Lanolin, Magnesium Hydroxide, Petrolatum, Phenoxyethanol, Propylene Glycol, Retinyl Palmitate, Sodium Chloride, Sodium Laureth Sulfate, Stearyl Alcohol, Water, Zea Mays Corn Oil, Zinc Chloride

Warnings

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For external use only

When using this product

When using this product do not get into the eyes

Stop use and ask a doctor if

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- Condition worsens
- Symptoms last more than 7 days or clear up and occur again within a few days

Keep out of reach of children

Keep out of reach of children. In case of accidental ingestion contact a physician or Poison Control Center right away.

Drug Facts

Active Ingredient	Purpose
Aluminum Hydroxide Gel 2%.....	Skin Protectant

Uses ■ Dries the oozing and weeping of poison
■ ivy ■ oak ■ sumac

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Directions

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■ Children under 6 months of age: ask a doctor

Other Information

■ Store between 15°-30°C (59°-86°F)

Inactive Ingredients

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Questions or Comments:

Call 1-877-894-6633 8 am – 6 pm EST M-F

Patient Name:
Room #:

PROCURE[®]
HEALTHCARE PRODUCTS

DryDrox
Aluminum
Hydroxide Gel 2%



Manufactured for:

Twin Med LLC.

11333 Greenstone Ave
Santa Fe Springs, CA 90670

ProCureProducts.com

Made in India

Item No.

PCMG35

**Poison ivy, oak and
sumac drying ointment**



(01) 10840330706268

SKIN PROTECTANT

NDC: 55681-031-04

NET WT 3.75 OZ (106g)

PROCURE DRYDROX

aluminum hydroxide gel 2% ointment				
Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:55681-031	
Route of Administration	TOPICAL			
Active Ingredient/Active Moiety				
Ingredient Name		Basis of Strength	Strength	
ALUMINUM HYDROXIDE (UNII: 5QB0T2IUN0) (ALUMINUM HYDROXIDE - UNII:5QB0T2IUN0)		ALUMINUM HYDROXIDE	2 g in 100 g	
Inactive Ingredients				
Ingredient Name			Strength	
ZEA MAYS (CORN) OIL (UNII: 8470G57WFM)				
GLYCERYL MONOSTEARATE (UNII: 230OU9XXE4)				
LANOLIN (UNII: 7EV65EAW6H)				
MAGNESIUM HYDROXIDE (UNII: NBZ3QY004S)				
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)				
CALCIUM CARBONATE (UNII: H0G9379FGK)				
CHOLECALCIFEROL (UNII: 1C6V77QF41)				
CITRIC ACID (UNII: 2968PHW8QP)				
GLYCERYL STEARATE/PEG-100 STEARATE (UNII: RD25J5V947)				
PETROLATUM (UNII: 4T6H12BN9U)				
PHENOXYETHANOL (UNII: HIE492ZZ3T)				
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)				
RETINYL PALMITATE (UNII: 1D1K0N0VVC)				
SODIUM CHLORIDE (UNII: 451W47IQ8X)				
SODIUM LAURETH SULFATE (UNII: BPV390UAP0)				
STEARYL ALCOHOL (UNII: 2KR89I4H1Y)				
WATER (UNII: 059QF0KO0R)				
ZINC CHLORIDE (UNII: 86Q357L16B)				
Product Characteristics				
Color	white (White to off-white)	Score		
Shape		Size		
Flavor		Imprint Code		
Contains				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:55681-031-04	106 g in 1 TUBE; Type 0: Not a Combination Product	09/15/2025	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M016	09/15/2025	

Labeler - Twin Med LLC (009579330)

Revised: 9/2025

Twin Med LLC