

RITE AID MAXIMUM STRENGTH ITCH RELIEF- diphenhydramine hydrochloride and zinc acetate spray

RITE AID CORPORATION

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

DRUG FACTS

Active ingredient

Purpose

Diphenhydramine HCL 2%Topical analgesic
Zinc Acetate 0.1%.....Skin Protectant

Uses

Temporarily relieves pain and itching associated with
- minor burns - insect bites - sunburn - minor skin irritations
- minor cuts - scrapes - rashes due to poison ivy, poison oak and poison sumac

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

Uses

Temporarily relieves pain and itching associated with
- minor burns - insect bites - sunburn - minor skin irritations
- minor cuts - scrapes - rashes due to poison ivy, poison oak and poison sumac

Warnings

For external use only

Do not use - on large areas of the body
- with any other product containing diphenhydramine,
even one taken by mouth

Ask a doctor before use - on chicken pox - on measles

When using this product avoid contact with the eyes

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

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

Directions

- do not use more than directed
- **Adults and children 2 years and older:** apply to affected area not more than 3 to 4 times daily
- **Children under 2 years of age:** ask a doctor

Other information

Store at 20 degrees to 25 degrees C (68 degrees to 77 degrees F)

Inactive Ingredients

purified water, SD alcohol 40-B, glycerin, povidone, tromethamine



NON-PATENTABLE

0 11822 92812 0

DISTRIBUTED BY:
RITE AID, 30 HUNTER LANE
CAMP HILL, PA 17011

IF YOU'RE NOT SATISFIED, WE'LL HAPPILY REFUND YOUR MONEY.

* This product is not manufactured or distributed by McNeil-PPC, Inc., owner of the registered trademark Benadryl®. R248RAD20Z-1

Drug Facts **DO NOT USE IF SAFETY SEAL LABEL IS TORN OR MISSING**

Active ingredients **Purpose**
Diphenhydramine HCl 2% ... Topical analgesic
Zinc acetate 0.1% Skin protectant

Drug Facts (continued) **TEAR HERE**

Uses temporarily relieves itching and pain associated with minor irritations, burns, scrapes, cuts, insect bites and rashes due to poison ivy, poison oak, and poison sumac. Dries the oozing and weeping of poison ivy, poison oak & poison sumac.

Warnings For external use only. Flammable. Keep away from fire or flame.

Do not use ■ with any other product containing diphenhydramine, even one taken by mouth
■ on large areas of the body

Ask a doctor before use ■ on chicken pox
■ on measles

When using this product avoid contact with the eyes

Stop use and ask a doctor if ■ condition worsens or does not improve within 7 days ■ symptoms persist for more than 7 days, or clear up and occur again within a few days

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

Directions ■ Do not use more than directed
■ **Adults and children 2 years of age & older:** apply to the affected area not more than 3 to 4 times daily
■ **children under 2 years of age:** consult a doctor

Other information store at 20°–25°C (68°–77°F)

Inactive ingredients purified water, SD alcohol 40-B, glycerin, povidone, tromethamine

RITE AID MAXIMUM STRENGTH ITCH RELIEF

diphenhydramine hydrochloride and zinc acetate spray

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:11822-0371
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
DIPHENHYDRAMINE HYDROCHLORIDE (UNII: TC2D6JAD40) (DIPHENHYDRAMINE - UNII:8GTS82S83M)	DIPHENHYDRAMINE HYDROCHLORIDE	20 mg in 1 mL
ZINC ACETATE (UNII: FM5526K07A) (ZINC CATION - UNII:13S1S8SF37)	ZINC ACETATE	1 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0K00R)	
ALCOHOL (UNII: 3K9958V90M)	
GLYCERIN (UNII: PDC6A3C0OX)	
POVIDONE (UNII: FZ989GH94E)	
TROMETHAMINE (UNII: 023C2WHX2V)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11822-0371-1	59 mL in 1 BOTTLE, SPRAY		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part336	08/10/2011	

Labeler - RITE AID CORPORATION (014578892)**Registrant** - Pharma Pac, LLC (140807475)**Establishment**

Name	Address	ID/FEI	Business Operations
Pharma Pac, LLC		140807475	manufacture

Revised: 4/2012

RITE AID CORPORATION