DG ANTI-ITCH- diphenhydramine hydrochloride and zinc acetate cream
Dollar General

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.

DG™ Anti-Itch
Anti-Itch Cream

Drug Facts

Active ingredients	***Purposes***
Diphenhydramine hydrochloride 2%	Topical analgesic
Zinc acetate 0.1%	Skin protectant

Uses

Temporarily relieves pain and itching associated with:
- insect bites
- minor burns
- sunburn
- scrapes
- minor skin irritations
- minor cuts

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

Warnings

For external use only.

Avoid contact with eyes.

Do Not Use
- more than directed
- on large areas of the body
- with any product containing diphenhydramine even one taken by mouth

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

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

If pregnant or breast-feeding, ask a health professional before use.

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away: 800-222-1222.

Directions

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

Other information

- store at 15 to 30°C (59 to 86°F)
- contents filled by weight, not volume
- **Tamper Evident:** DO NOT USE IF SEAL ON TUBE IS PUNCTURED OR MISSING.

Inactive ingredients

cetearyl alcohol (and) polysorbate 60, DMDM hydantoin, glyceryl stearate, isopropyl myristate, mineral oil, phenoxy ethenol, purified water, sodium benzoate

Questions or comments?

1-888-309-9030

DISTRIBUTED BY OLD
EAST MAIN CO.,
100 MISSION RIDGE
GOODLETTSVILLE, TN
37072

PRINCIPAL DISPLAY PANEL - 28 g Tube Carton

DG™|health

NDC 55910-344-24

Compare to
active
ingredients
of Benadryl®
Extra
Strength
Cream*

Extra Strength
Anti-Itch
Cream
Topical Analgesic

and Skin Protectant

Relieves Itches
From Insect Bites
& Skin Irritations

NET WT 1 OZ (28 g)

DG health

NDC 55910-344-24

Compare to active ingredients of Benadryl® Extra Strength Cream*

Extra Strength

Anti-Itch Cream

Topical Analgesic and Skin Protectant

Relieves Itches From Insect Bites & Skin Irritations

NET WT 1 OZ (28 g)

PL306638 V04

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Extra Strength
Anti-Itch Cream
Topical Analgesic and Skin Protectant
Relieves Itches From Insect Bites & Skin Irritations
NET WT 1 OZ (28 g)

DG health

NDC 55910-344-24
Compare to active ingredients of Benadryl® Extra Strength Cream*

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8 38891 00313 4

Drug Facts (continued)

hydantoin, glyceryl stearate, isopropyl myristate, mineral oil, phenoxy ethenol, purified water, sodium benzoate

Questions or comments?
1-888-309-9030

*This product is not manufactured by or distributed by Johnson & Johnson Consumer Products Co., the distributor of Benadryl®.

100% Satisfaction Guaranteed!
(888) 309-9030

DISTRIBUTED BY OLD EAST MAIN CO.,
100 MISSION RIDGE
GOODLETTSVILLE, TN
37072

MADE IN U.S.A.

DG ANTI-ITCH

diphenhydramine hydrochloride and zinc acetate cream

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:55910-344
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 g
Zinc acetate (UNII: FM5526K07A) (zinc cation - UNII:13S1S8SF37)	Zinc acetate	0.1 mg in 1 g

Inactive Ingredients

Ingredient Name	Strength
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)	
polysorbate 60 (UNII: CAL22UVI4M)	
DMDM HYDANTOIN (UNII: BYR0546TOW)	
GLYCERYL MONOSTEARATE (UNII: 230OU9XXE4)	
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
mineral oil (UNII: T5L8T28FGP)	
Sodium Benzoate (UNII: OJ245FE5EU)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:55910-344-24	1 in 1 CARTON	10/19/2017	
1		28 g in 1 TUBE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part348	05/17/2011	

Labeler - Dollar General (068331990)

Establishment

Name	Address	ID/FEI	Business Operations
Natureplex LLC		062808196	MANUFACTURE(55910-344)