FAMILY WELLNESS- diphenhydramine hydrochloride, zinc acetate cream
FRONT PHARMACEUTICAL PLC

Family Wellness Anti-Itch Cream, 28g

Active Ingredients

Diphenhydramine hydrochloride 2%

Zinc acetate 0.1%

Purpose

Topical analgesic

Skin protectant

for temprary relief from pain and itching associated **Uses**

with insect bites , sunburn and mior skin irritations. Dries the

oozing and weeping of poison: •ivy •oak •sumac

Warnings

. **For external use only**

Do not use

•avoid contact with eyes **When using this product**

Ask a doctor before use

•on chicken pox •on measles

Stop use and ask a doctor if

- over large areas of the body
- with any other product containing diphenhydramine including those taken orally
- symptoms last for more than 7 days
- the condition gets worse
- symptoms clear up and then occur again within a few days

. If swallowed, get medical help **Keep out of reach of children**

or contact a Poison Control Center right away.

Directions

•for adults and children over 2 years of age: apply to affected

area not more than 3 to 4 times daily

•children under 2 years of age: ask a doctor before use

store at room temperature 59° - 86° F **Other information**

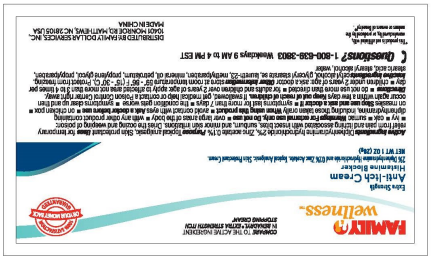
(15° - 30° C).

cetyl alcohol, glyceryl stearate se, **Inactive ingredients**

laureth-23, methylparaben, mineral oil, petrolatum, propylene glycol, propylparaben, stearic acid, stearic alcohol, water

1-800-639-3803 Weekdays 9 AM to 4 PM EST **Questions?**

label



FAMILY WELLNESS

diphenhydramine hydrochloride, zinc acetate cream

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:69571-005
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
DIPHENHYDRAMINE HYDROCHLORIDE (UNII: TC2D6JAD40) (DIPHENHYDRAMINE - UNII:8GTS82S83M)	DIPHENHYDRAMINE HYDROCHLORIDE	2 g in 100 g
ZINC ACETATE (UNII: FM5526K07A) (ZINC CATION - UNII:13S1S8SF37)	ZINC CATION	0.1 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
CETYL ALCOHOL (UNII: 936JST6JCN)	
GLYCERYL STEARATE SE (UNII: FCZ5MH785I)	
LAURETH-23 (UNII: N72LMW566G)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
MINERAL OIL (UNII: T5L8T28FGP)	
PETROLATUM (UNII: 4T6H12BN9U)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
STEARYL ALCOHOL (UNII: 2KR89I4H1Y)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69571-005-02	1 in 1 BOX	03/29/2017	
1	NDC:69571-005-01	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 Drug	M017	03/29/2017	

Labeler - FRONT PHARMACEUTICAL PLC (530897792)

Establishment

Name	Address	ID/FEI	Business Operations
FRONT PHARMACEUTICAL PLC		530897792	manufacture(69571-005)

Revised: 10/2023 FRONT PHARMACEUTICAL PLC