EACH RELIEF EQUATE- diphenhydramine hci 2% gel
Walmart

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 HCI 2%.....Topical analgesic

Uses Temporarily relieves pain due to: • minor burns • insect bites • sunburn • minor skin irritations • minor cuts • scrapes • rashes due to poison ivy, poison oak & 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
When using this product • avoid contact with the eyes
S**top use and ask doctor if** • condition gets worse • symptoms last more than 7 days • symptoms clear up and occur again in 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 and older: apply to affected area not more than 3 to 4 times daily • children under 2 years of age: ask a doctor.

Inactive ingredients

Camphor, Citric Acid, Diazolidinyl
Urea, Glycerin, Hydroxypropyl Methylcellulose, Methylparaben,
Propylene Glycol, Propylparaben, SD Alcohol 38-B, Sodium
Citrate, Water.

EACH RELIEF EQUATE

diphenhydramine hci 2% gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:49035-246
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 mL

Inactive Ingredients	
Ingredient Name	**Strength**
CAMPHOR (SYNTHETIC) (UNII: 5TJD82A1ET)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
Diazolidinyl Urea (UNII: H5RIZ3MPW4)	
Glycerin (UNII: PDC6A3C0OX)	
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
Methylparaben (UNII: A2I8C7HI9T)	
Propylene Glycol (UNII: 6DC9Q167V3)	
Propylparaben (UNII: Z8IX2SC1OH)	
ALCOHOL (UNII: 3K9958V90M)	
Sodium Citrate (UNII: 1Q73Q2JULR)	
Water (UNII: 059QF0KO0R)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:49035-246-03	103 mL in 1 TUBE; Type 0: Not a Combination Product	04/03/2017	

Marketing Information			
Marketing Category	**Application Number or Monograph Citation**	**Marketing Start Date**	**Marketing End Date**
OTC monograph not final	part348	04/03/2017	

Labeler - Walmart (051957769)

Registrant - Product Quest Mfg (927768135)

Establishment			
Name	**Address**	**ID/FEI**	**Business Operations**
Product Quest Mfg		927768135	manufacture(49035-246) , label(49035-246)

Revised: 1/2017 Walmart