

**EQUATE ITCH RELIEF GEL- itch relief gel gel**  
**Walmart, Inc**

**Equate Itch Relief Gel**

Do not use more often 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.

For external use only.

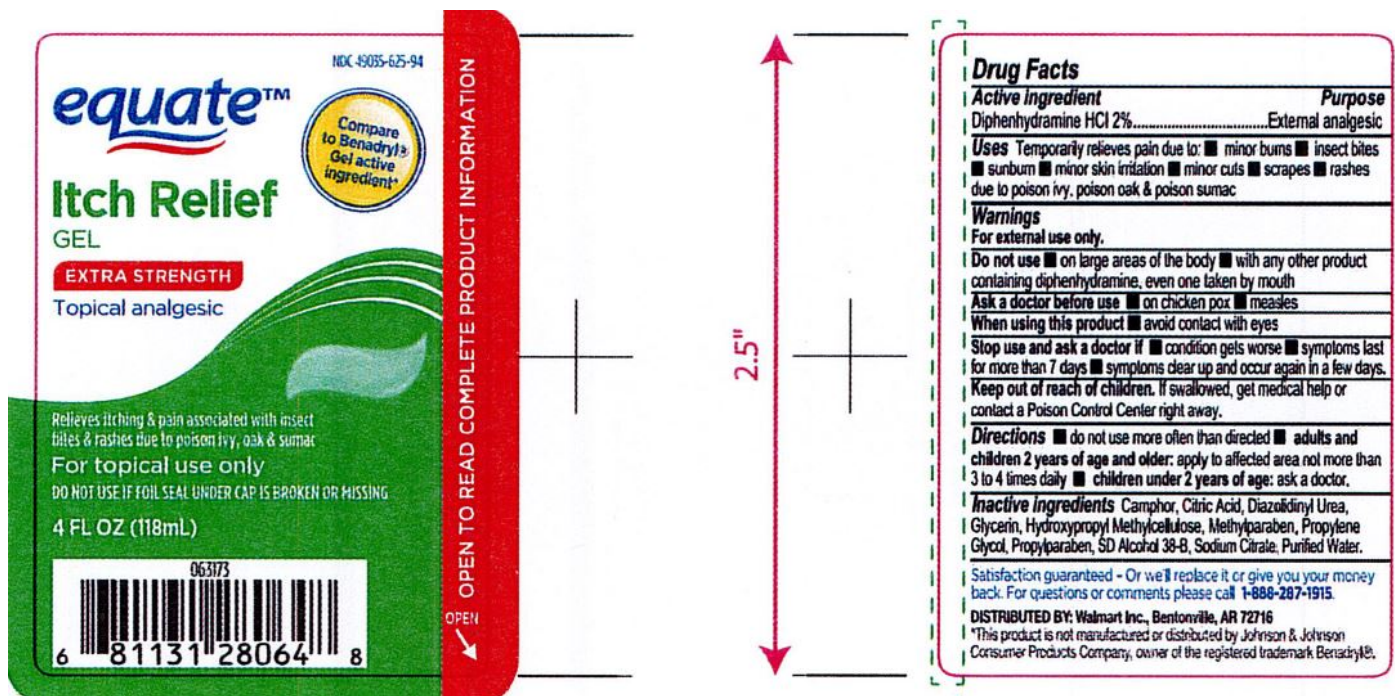
Camphor, Citric Acid, Diazolidnyl Urea, Glycerin, Hydroxypropyl Methylcellulose, Methylparaben, Propylene Glycol, Propylparaben, SD Alcohol 38-B, Sodium Citrate, Purified Water.

Temporarily relieves pain due to: Minor burns, insect bites, sunburn, minor skin irritation, 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.

External Analgesic

Diphenhydramine HCl 2%



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<b>EQUATE ITCH RELIEF GEL</b>
itch relief gel gel
<b>Product Information</b>

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:49035-625	
<b>Route of Administration</b>	TOPICAL			
<b>Active Ingredient/Active Moiety</b>				
<b>Ingredient Name</b>		<b>Basis of Strength</b>	<b>Strength</b>	
DIPHENHYDRAMINE HYDROCHLORIDE (UNII: TC2D6JAD40) (DIPHENHYDRAMINE - UNII:8GTS82S83M)		DIPHENHYDRAMINE HYDROCHLORIDE	2 mg in 100 mL	
<b>Inactive Ingredients</b>				
<b>Ingredient Name</b>			<b>Strength</b>	
PROPYLPARABEN (UNII: Z8IX2SC1OH)				
METHYLPARABEN (UNII: A2I8C7HI9T)				
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)				
ALCOHOL (UNII: 3K9958V90M)				
DIAZOLIDINYL UREA (UNII: H5RIZ3MPW4)				
SODIUM CITRATE (UNII: 1Q73Q2JULR)				
WATER (UNII: 059QF0KO0R)				
GLYCERIN (UNII: PDC6A3C0OX)				
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)				
CAMPHOR (SYNTHETIC) (UNII: 5TJD82A1ET)				
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)				
<b>Packaging</b>				
<b>#</b>	<b>Item Code</b>	<b>Package Description</b>	<b>Marketing Start Date</b>	<b>Marketing End Date</b>
1	NDC:49035-625-94	118 mL in 1 CONTAINER; Type 0: Not a Combination Product	12/20/2018	
<b>Marketing Information</b>				
<b>Marketing Category</b>	<b>Application Number or Monograph Citation</b>		<b>Marketing Start Date</b>	<b>Marketing End Date</b>
OTC Monograph Drug	M017		12/20/2018	

**Labeler** - Walmart, Inc (051957769)

**Registrant** - Pharma Nobis, LLC (118564114)

### Establishment

Name	Address	ID/FEI	Business Operations
Pharma Nobis, LLC		118564114	manufacture(49035-625) , label(49035-625) , pack(49035-625) , analysis(49035-625)

