DERMAREST ECZEMA MEDICATED- hydrocortisone lotion
Insight Pharmaceuticals LLC

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.

Dermarest Eczema Medicated Lotion

Dermarest™ Eczema
Medicated Lotion

Keep carton for future reference.

Drug Facts

Active ingredient

Hydrocortisone 1%

Purpose

Anti-itch

Uses

temporary relief of itching associated with
- minor skin irritations
- inflammation
- rashes due to eczema

Other uses of this product should be only under the advice and supervision of a doctor.

Warnings

For external use only

Do not use

for the treatment of diaper rash

When using this product

do not get into the eyes.

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

Do not begin use of any other hydrocortisone product unless you have consulted a doctor.

Keep out of reach of children.

If swallowed, get medical help or contact a Poison Control Center (1-800-222-1222) right away.

Directions

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	do not use, consult a doctor

Other Information

- shake well
- store at room temperature 15° - 25°C (59°- 77°F)

Inactive ingredients

Purified water, propylene glycol, glycerin, cetyl alcohol, glyceryl stearate, dimethicone, aleurites moluccana seed oil, zinc PCA, panthenol, carthamus tinctorius (safflower) flower extract, camellia sinensis leaf extract, rheum palmatum root/stalk extract, cyclohexasiloxane, cyclopentasiloxane, xanthan gum, disodium edetate, sodium hydroxide, diazolidinyl urea, methylparaben, propylparaben

Questions?

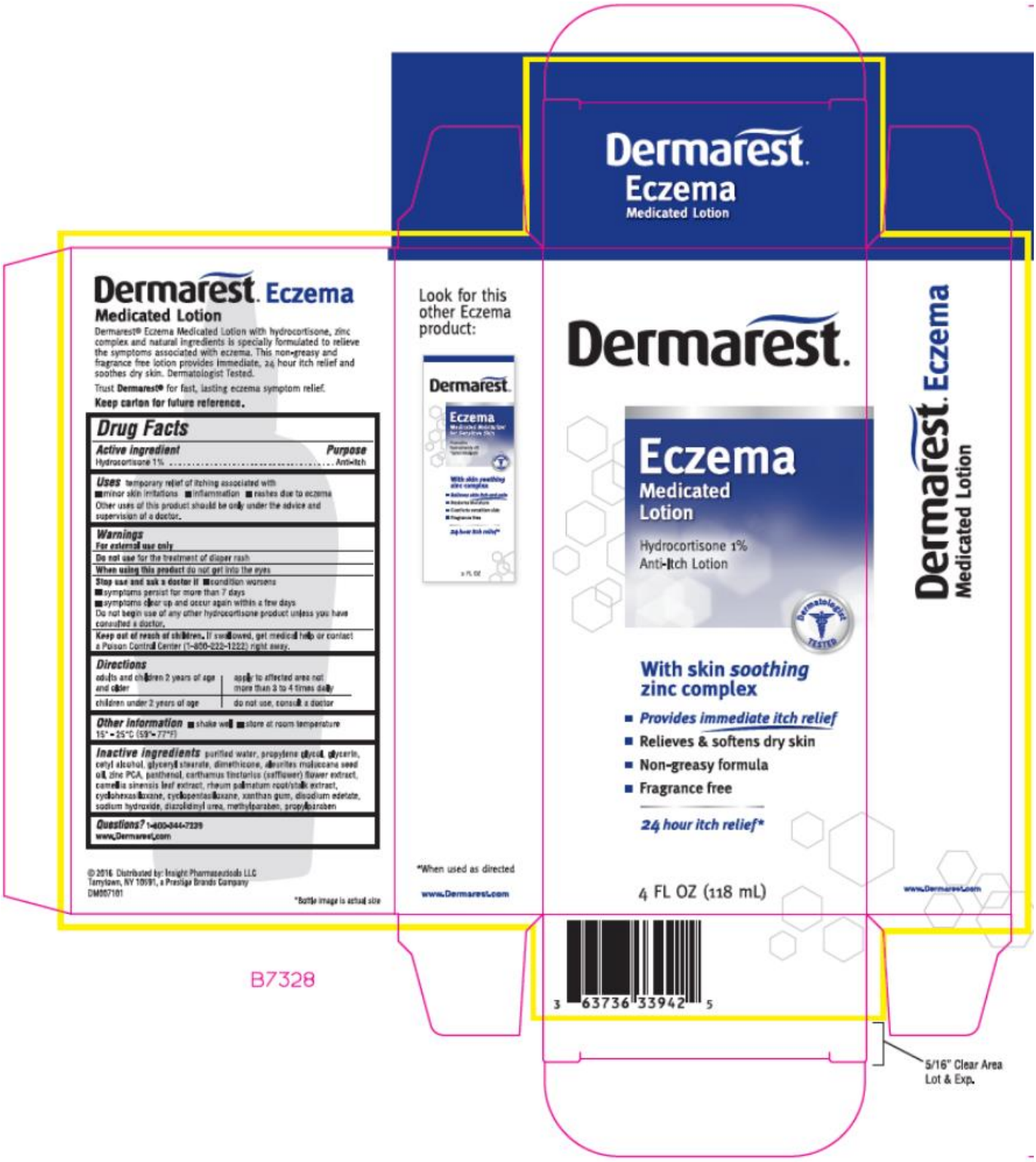
1-800-344-7239 www.Dermarest.com

PRINCIPAL DISPLAY PANEL

Dermarest®

Eczema
Medicated
Lotion

Hydrocortisone 1%
Anti-Itch Lotion
4 FL OZ (118 mL)



DERMAREST ECZEMA MEDICATED

hydrocortisone lotion

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:63736-339
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
HYDROCORTISONE (UNII: WI4X0X7BPJ) (HYDROCORTISONE - UNII:WI4X0X7BPJ)	HYDROCORTISONE	10 mg in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
GLYCERIN (UNII: PDC6A3C0OX)	
CETYL ALCOHOL (UNII: 936JST6JCN)	
GLYCERYL MONOSTEARATE (UNII: 230OU9XXE4)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
KUKUI NUT OIL (UNII: TP11QR7B8R)	
ZINC PIDOLATE (UNII: C32PQ86DH4)	
PANTHENOL (UNII: WV9CM0O67Z)	
SAFFLOWER (UNII: 4VBL71TY4Y)	
GREEN TEA LEAF (UNII: W2ZU1RY8B0)	
RHEUM PALMATUM ROOT (UNII: G025DAL7CE)	
CYCLOMETHICONE 6 (UNII: XHK3U310BA)	
CYCLOMETHICONE 5 (UNII: 0THT5PCI0R)	
XANTHAN GUM (UNII: TTV12P4NEE)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
DIAZOLIDINYL UREA (UNII: H5RIZ3MPW4)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	

Product Characteristics			
Color	WHITE	**Score**	
Shape		**Size**	
Flavor		**Imprint Code**	
Contains			

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:63736-339-24	24 in 1 CASE	09/20/2010	
1		1 in 1 CARTON		
1		118 mL in 1 BOTTLE; 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	09/20/2010	

Labeler - Insight Pharmaceuticals LLC (055665422)

Revised: 1/2020 Insight Pharmaceuticals LLC