

**DERMACORT ANTI-ITCH- hydrocortisone cream**  
**Melaleuca, Inc.**

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**DermaCort**

**Active ingredient**

Hydrocortisone 1%

**Purpose**

Anti-itch

**Uses**

- For the temporary relief of itching associated with minor skin irritations, inflammation and rashes due to eczema, psoriasis, , seborrheic dermatitis, poison ivy, poison oak, poison sumac, insect bites, soaps, detergents, and jewelry
- Other use of this product should be only under the advice and supervision of a doctor.

**Warnings**

**For external use only.**

**When using this product** avoid contact with the eyes.

**Stop use and ask a doctor if** conditions worsens, or if symptoms persist for more than 7 days or clear up and occur again within a few days, stop use of this product and do not begin use of any other hydrocortisone product unless you have consulted a doctor.

**Do not use** for the treatment of diaper rash. Consult a doctor.

**Keep out of reach of children.**

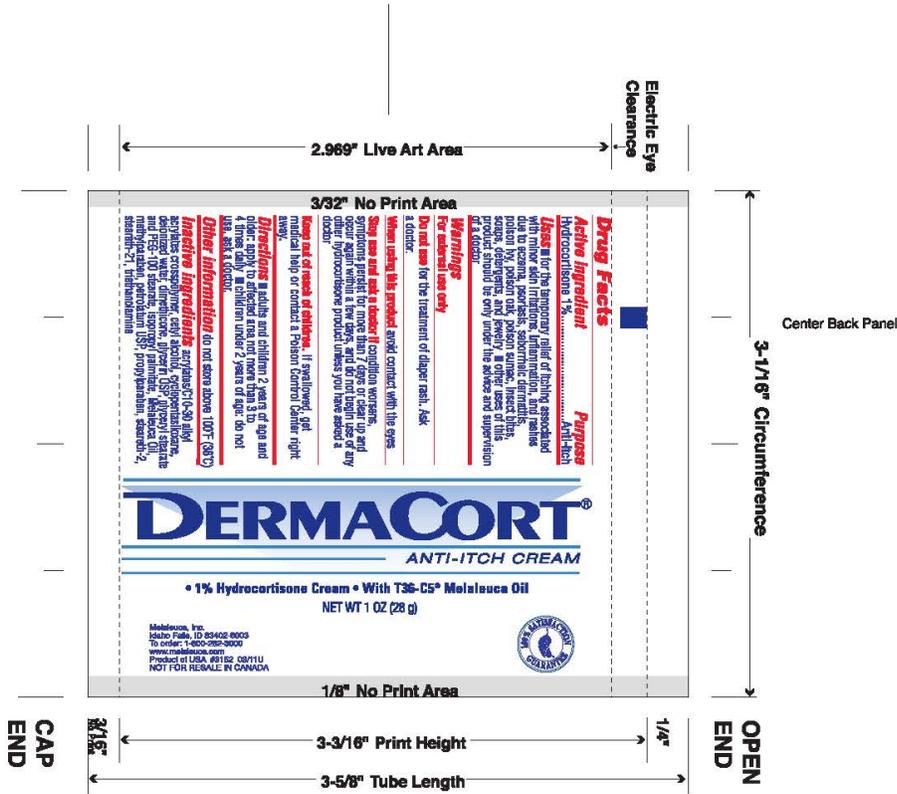
In case of accidental ingestion, seek professional assistance or contact a Poison Control Center immediately.

**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** Do not store above 100°F.

**Inactive ingredients** Acrylates/C10-30 Alkyl Acrylates Crosspolymer, Cetyl Alcohol, Cyclopentasiloxane, Deionized Water, Dimethicone, Glycerin USP, Glyceryl Stearate and PEG-100 Stearate, Isopropyl Palmitate, Melaleuca Oil, Methylparaben, Petrolatum USP, Propylparaben, Serareth-2, Steareth-21, Triethanolamine.

# Compax 1" x 3-5/8"



505003152

LEGEND	
	Pantone 279 C
	PANTONE 2746 C
	PANTONE 485 CVC

Customer \_\_\_\_\_

Date \_\_\_\_\_

NOTE: CHANGES ARE COSTLY AND WILL CAUSE ADDITIONAL EXPENSE AND DELAY

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hydrocortisone cream

## Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:54473-117
<b>Route of Administration</b>	TOPICAL		

## Active Ingredient/Active Moiety

<b>Ingredient Name</b>	<b>Basis of Strength</b>	<b>Strength</b>
<b>HYDROCORTISONE</b> (UNII: W4X0X7BPJ) (HYDROCORTISONE - UNII:W4X0X7BPJ)	HYDROCORTISONE	10 mg in 1 g

## Inactive Ingredients

<b>Ingredient Name</b>	<b>Strength</b>
<b>POLYOXYL 100 STEARATE</b> (UNII: YD01N1999R)	
<b>PROPYLPARABEN</b> (UNII: Z8IX2SC1OH)	
<b>STEARETH-2</b> (UNII: V56DFE46J5)	
<b>STEARETH-21</b> (UNII: 53J3F32P58)	
<b>TEA TREE OIL</b> (UNII: VIF565UC2G)	
<b>TROLAMINE</b> (UNII: 9O3K93S3TK)	
<b>WATER</b> (UNII: 059QF0KO0R)	
<b>CARBOMER COPOLYMER TYPE B (ALLYL PENTAERYTHRITOL CROSSLINKED)</b> (UNII: 809Y72KV36)	
<b>CETYL ALCOHOL</b> (UNII: 936JST6JCN)	
<b>CYCLOMETHICONE 5</b> (UNII: 0THT5PCI0R)	
<b>DIMETHICONE</b> (UNII: 92RU3N3Y1O)	
<b>GLYCERIN</b> (UNII: PDC6A3C0OX)	
<b>GLYCERYL MONOSTEARATE</b> (UNII: 230OU9XXE4)	
<b>ISOPROPYL PALMITATE</b> (UNII: 8CRQ2TH63M)	
<b>METHYLPARABEN</b> (UNII: A2I8C7HI9T)	
<b>PETROLATUM</b> (UNII: 4T6H12BN9U)	

## Packaging

<b>#</b>	<b>Item Code</b>	<b>Package Description</b>	<b>Marketing Start Date</b>	<b>Marketing End Date</b>
1	NDC:54473-117-28	28 g in 1 TUBE; Type 0: Not a Combination Product	01/01/2018	

## Marketing Information

<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	08/02/2000	

**Labeler** - Melaleuca, Inc. (139760102)

**Registrant** - Melaleuca, Inc. (139760102)

**Establishment**

Name	Address	ID/FEI	Business Operations
Melaleuca, Inc.		079711683	manufacture(54473-117)

Revised: 3/2025

Melaleuca, Inc.