

# **SEBORRHEIC DERMATITIS AND PSORIASIS CREAM- roycederm seborrheic dermatitis and psoriasis cream cream**

**Ehy Holdings LLC**

*Disclaimer: This drug has not been found by FDA to be safe and effective, and this labeling has not been approved by FDA. For further information about unapproved drugs, click here.*

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**81799-006**

**Seborrheic Dermatitis and Psoriasis Cream**

## **Active Ingredient(s)**

ARTEMISIA EXTRACT □FHFII□3%

## **Purpose**

Seborrheic

## **Use**

Relieves and helps prevent recurrence of scalp: itching, flaking, scaling, redness, and irritation associated with psoriasis and seborrheic dermatitis.

## **Warnings**

For external use only

## **Do not use**

Do not use more than 3 times daily. Children under 12 years: ask a doctor

When using this product avoid contact with eyes.

condition worsens or does not improve after regular use as directed

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

## **Directions**

Adult and children over 12 years or old. Wash hands before and after applying the cream. Apply to affected area at the first sign of blisters, sores, or tingling sensation. Rub in gently and thoroughly. Do not use more than 3 times daily. Children under 12 years: ask a doctor

## **Other information**

Store at room temperature and out of direct sunlight.

## Inactive ingredients

Purified water, Artemisia extracts Sophora Flavescens, Cnidium Monnieri (L) Cuss., Kochia scoparia, Dictamni Cortex, Smilax Glabra Roxb, Solidago Decurrens, Phellodendron amurense Ruprecht., Gleditsia Sinensis

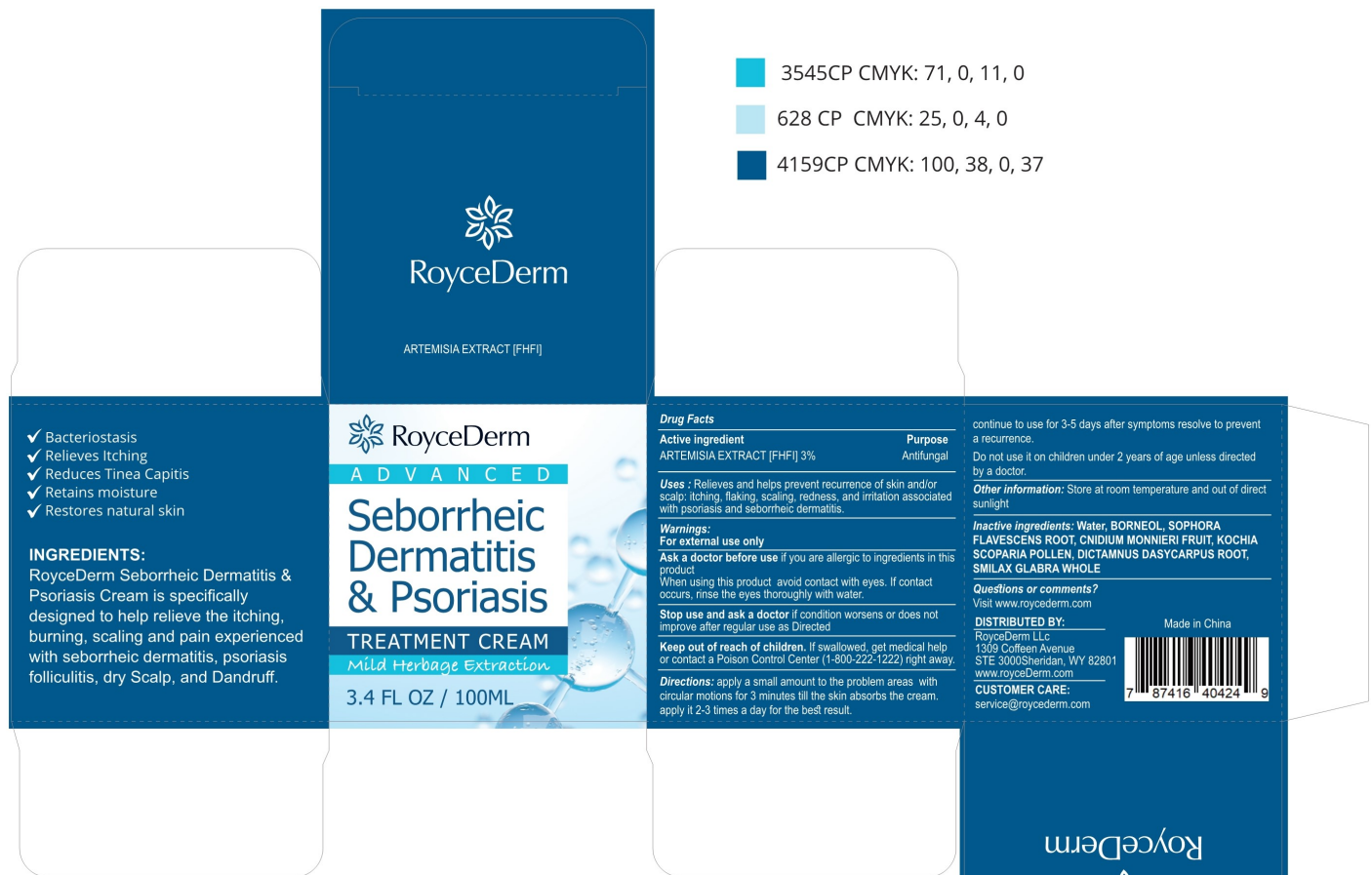
Apply 2-3 times daily. Children under 12 years: ask a doctor

Visit [www.roycederm.com](http://www.roycederm.com)

## Package Label - Principal Display Panel

81799-006-1

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- 3545CP CMYK: 71, 0, 11, 0
- 628 CP CMYK: 25, 0, 4, 0
- 4159CP CMYK: 100, 38, 0, 37

Sticker size: 35 mm x 150 mm



81799-006-1



## SEBORRHEIC DERMATITIS AND PSORIASIS CREAM

roycederm seborrheic dermatitis and psoriasis cream cream

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:81799-006
<b>Route of Administration</b>	TOPICAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>WORMWOOD</b> (UNII: F84709P2XV) (WORMWOOD - UNII:F84709P2XV)	WORMWOOD	3 mg in 100 mL

### Inactive Ingredients

Ingredient Name	Strength
<b>DICTAMNUS DASYPARPUS ROOT</b> (UNII: 6153LEN214)	
<b>WATER</b> (UNII: 059QF0KO0R)	
<b>BORNEOL</b> (UNII: M89NIB437X)	
<b>CNIDIUM MONNIERI FRUIT</b> (UNII: V1IA3S3CUS)	
<b>BASSIA SCOPARIA POLLEN</b> (UNII: 07A108ZKW5)	
<b>SOPHORA FLAVESCENS ROOT</b> (UNII: IYR6K8KQ5K)	

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:81799-006-01	100 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/11/2023	

## Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		01/11/2023	

**Labeler** - Ehy Holdings LLC (117322715)

## Establishment

Name	Address	ID/FEI	Business Operations
Ehy Holdings LLC		117322715	manufacture(81799-006)

Revised: 1/2023

Ehy Holdings LLC