SHINGLES PAIN RELIEF CREAM- roycederm shingles pain relief 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.

81799-009
Shingles Pain Relief Cream

Active Ingredient(s)

Artemisia extract [fhfi] 3%

Purpose

for shingles skin care

Use

For the treatment of symptomsof shingles, hives, and rashes.

Warnings

FOR EXTERNAL USE ONLY

Do not use

it on pregnant womenand children under 12 years ofage unless directed by a doctor

avoid contact with eyesIf contact occurs, rinse the eyesthoroughly with water.

condition worsens or does notimprove after regular use asdirected

If swallowed, getmedical help or contact aPoison Control Center(1-800-222-1222) right away.

Directions

Apply a small amount to theproblem areas with circularmotions for 3 minutes till theskin absorbs the cream.
Apply it 2-3 times a day for thebest result.

Other information

Store at room temperature andout of direct sunlight

Inactive ingredients

Water, Sophora Flavescens Root,Cnidium Monnieri Fruit, KochiaScoparia Pollen, Dictamnus Dasyca-rpus Root, Smilax Glabra Whole

if you are allergic to ingredientsin this product.

Visit www.roycederm.com

Package Label - Principal Display Panel

81799-009-01

SHINGLES PAIN RELIEF CREAM

roycederm shingles pain relief cream cream

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:81799-009
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
WORMWOOD (UNII: F84709P2XV) (WORMWOOD - UNII:F84709P2XV)	WORMWOOD	3 g in 100 g

Inactive Ingredients	
Ingredient Name	**Strength**

DICTAMNUS DASYCARPUS ROOT (UNII: 6153LEN214)	
WATER (UNII: 059QF0KO0R)	
SOPHORA FLAVESCENS ROOT (UNII: IYR6K8KQ5K)	
SMILAX GLABRA WHOLE (UNII: H51N91QNEB)	
CNIDIUM MONNIERI FRUIT (UNII: V1IA3S3CUS)	
BASSIA SCOPARIA POLLEN (UNII: 07A108ZKW5)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:81799-009-01	60 g in 1 BOTTLE; Type 0: Not a Combination Product	02/15/2023	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		02/15/2023	

Labeler - Ehy Holdings LLC (117322715)

Establishment

Name	Address	ID/FEI	Business Operations
Ehy Holdings LLC		117322715	manufacture(81799-009)

Revised: 2/2023 Ehy Holdings LLC