

DERMAHARMONY SULFUR AND SALICYLIC ACID- salicylic acid, sulfur soap
D3 Development, Inc.

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.

Dermaharmony Sulfur and Salicylic Acid Soap

Drug Facts

Active ingredients

Salicylic acid 2%

Sulfur 5%

Purpose

Dandruff / seborrheic dermatitis / psoriasis soap

Uses

☐Helps prevent and control recurrence of itching, flaking and irritation associated with dandruff, seborrheic dermatitis, and psoriasis

Warnings

For external use only

Ask a doctor before use if you have a condition that covers a large area of the body.

When using this product, avoid contact with eyes. If contact occurs, rinse eyes thoroughly with water.

Stop use and ask a doctor if condition worsens or does not improve after regular use as directed.

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

Directions

- Use on affected areas in place of your regular soap.
- For best results use at least twice a week or as directed by a doctor.
- Work up a lather using warm water and massage into affected areas.
- Rinse.

Other information

- Store at 59° to 77°F (15° to 25°C)
- Lot number and expiration date are printed on back panel

Inactive ingredients

Sodium palmate, sodium cocoate*, sodium palm kernelate*, water, glycerin, soyethyl morpholinium ethosulfate (soy derived), sodium chloride, pentasodium pentetate

*May contain this ingredient

Questions?

1-800-827-3730

www.dermaharmony.com

Distributed by:

D3 Development, Inc., Portland, ME 04101

Made in the USA

dermaharmony

5% Sulfur & 2% Salicylic Acid SOAP

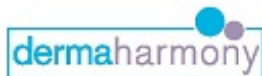
For Dandruff, Seborrheic Dermatitis & Psoriasis

NET WT 4 OZ (113 G)

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DERMAHARMONY SULFUR AND SALICYLIC ACID

salicylic acid, sulfur soap

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:71819-004
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
SULFUR (UNII: 70FD1KFU70) (SULFUR - UNII:70FD1KFU70)	SULFUR	5 g in 100 g
SALICYLIC ACID (UNII: O414PZ4LPZ) (SALICYLIC ACID - UNII:O414PZ4LPZ)	SALICYLIC ACID	2 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
WATER (UNII: 059QF0KO0R)	
SODIUM COCOATE (UNII: R1TQH25F4I)	
GLYCERIN (UNII: PDC6A3C0OX)	
SODIUM PALM KERNELATE (UNII: 6H91L1NXTW)	
SODIUM PALMATE (UNII: S0A6004K3Z)	
SOYETHYL MORPHOLINIUM ETHOSULFATE (UNII: J8C5W5HH18)	
PENTASODIUM PENTETATE (UNII: 961TOZ5L7T)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:71819-004-04	113 g in 1 CELLO PACK; Type 0: Not a Combination Product	07/02/2019	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part358H	07/02/2019	

Labeler - D3 Development, Inc. (043195877)

Revised: 7/2019

D3 Development, Inc.