

SULFUR 8 SCALP THERAPY MEDICATED DANDRUFF CONTROL SCALP BUTTER CREAM- sulfur, salicylic acid ointment
J. Strickland and Co.

Sulfur 8 Scalp Therapy Medicated Dandruff Control Scalp Butter Cream

Drug Facts

Active Ingredient

Sulfur, 5%

Salicylic Acid, 3%

Purpose

Antidandruff

Use

- relieves the itching and scaling associated with dandruff

Warnings

For external use only

When using this product

avoid contact with eyes. If contact occurs rinse eyes thoroughly with water.

Stop use and consult a doctor if

- condition worsens or does not improve after regular use as directed.
- HAIR IS FLAMMABLE. Flammability is increased by product build-up. Keep hair away from sparks, flame, extreme heat or lit tobacco.

Keep out of reach of children.

If swallowed, get medical help or call a Poison Control Center right away.

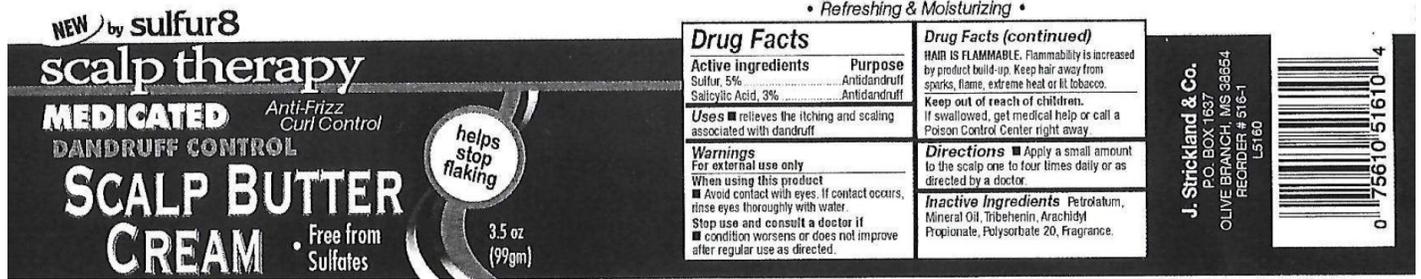
Directions

- Apply a small amount to the scalp one to four times daily or as directed by a doctor.

Inactive Ingredients

Petrolatum, Mineral Oil, Tribehenin, Arachidyl Propionate, Polysorbate 20, Fragrance

Package Labeling:



SULFUR 8 SCALP THERAPY MEDICATED DANDRUFF CONTROL SCALP BUTTER CREAM

sulfur, salicylic acid ointment

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:12022-035
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
SULFUR (UNII: 70FD1KFU70) (SULFUR - UNII:70FD1KFU70)	SULFUR	50 mg	in 1 g
SALICYLIC ACID (UNII: O414PZ4LPZ) (SALICYLIC ACID - UNII:O414PZ4LPZ)	SALICYLIC ACID	30 mg	in 1 g

Inactive Ingredients		
Ingredient Name	Strength	
PETROLATUM (UNII: 4T6H12BN9U)		
MINERAL OIL (UNII: T5L8T28FGP)		
TRIBEHENIN (UNII: 8OC9U7TQZ0)		
ARACHIDYL PROPIONATE (UNII: QV5DAH3MSB)		
POLYSORBATE 20 (UNII: 7T1F30V5YH)		

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:12022-035-00	99 g in 1 JAR; Type 0: Not a Combination Product	01/01/2020	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M032	01/01/2020	

Revised: 10/2023

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