

**SULFUR 8 MEDICATED DANDRUFF WITH SALICYLIC ACID- salicylic acid liquid
J. Strickland and Co.**

Sulfur 8 Medicated Dandruff Shampoo with Salicylic Acid

Drug Facts

Active Ingredient

Salicylic Acid, 2%

Purpose

Antidandruff, Seborrheic dermatitis, Psoriasis.

Uses

control the symptoms of

- dandruff
- seborrheic dermatitis
- 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 of this product as directed.

Keep out of reach of children.

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

Directions

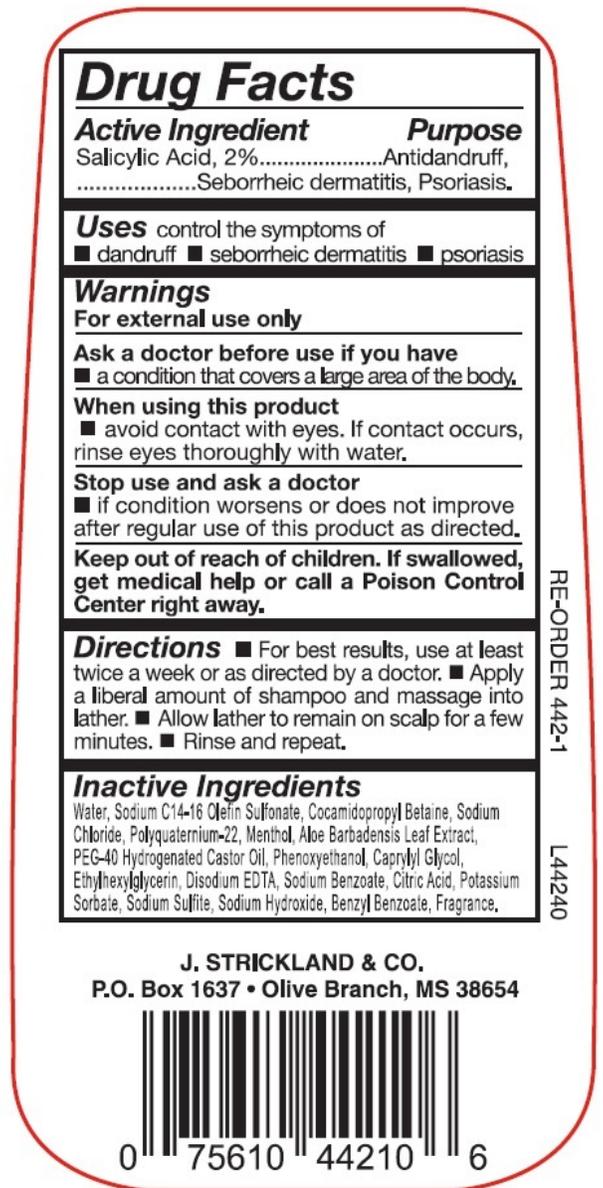
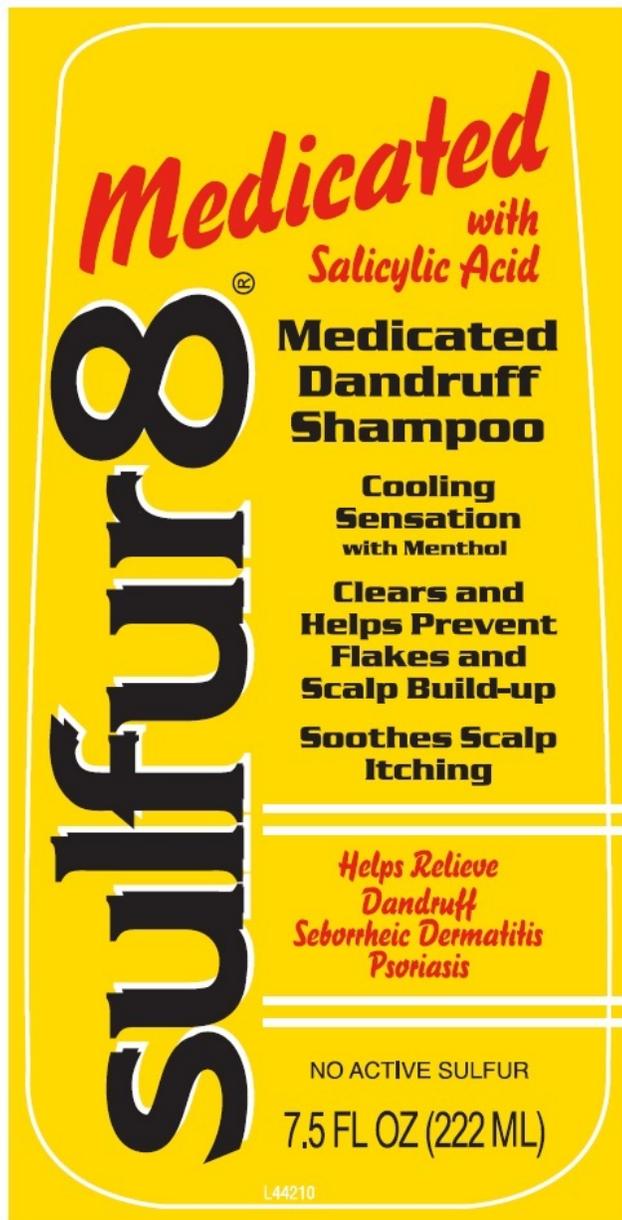
- For best results, use at least twice a week or as directed by a doctor.
- Apply a liberal amount of shampoo and massage into lather.
- Allow lather to remain on scalp for a few minutes.

- Rinse and repeat.

Inactive Ingrdients

Water, Sodium C14-16 Olefin Sulfonate, Cocamidopropyl Betaine, Sodium Chloride, Polyquaternium-22, Menthol, Aloe Barbadensis Leaf Extract, PEG-40 Hydrogenated Castor Oil, Phenoxyethanol, Caprylyl Glycol, Ethylhexylglycerin, Disodium EDTA, Sodium Benzoate, Citric Acid, Potassium Sorbate, Sodium Sulfite, Sodium Hydroxide, Benzyl Benzoate, Fragrance.

Package Labeling:



SULFUR 8 MEDICATED DANDRUFF WITH SALICYLIC ACID

salicylic acid liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:12022-039
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
SALICYLIC ACID (UNII: O414PZ4LPZ) (SALICYLIC ACID - UNII:O414PZ4LPZ)	SALICYLIC ACID	20 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
SODIUM C14-16 OLEFIN SULFONATE (UNII: O9W3D3YF5U)	
COCAMIDOPROPYL BETAINE (UNII: 5OCF3O11KX)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
MENTHOL (UNII: L7T10EIP3A)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
POLYOXYL 40 HYDROGENATED CASTOR OIL (UNII: 7YC686GQ8F)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
CAPRYLYL GLYCOL (UNII: 00YIU5438U)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
EDETATE DISODIUM ANHYDROUS (UNII: 8NLQ36F6MM)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)	
SODIUM SULFITE (UNII: VTK01UQK3G)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
BENZYL BENZOATE (UNII: N863NB338G)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:12022-039-00	222 mL in 1 BOTTLE; Type 0: Not a Combination Product	10/01/2023	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M032	10/01/2023	

Labeler - J. Strickland and Co. (007023112)