SULFUR 8 ANTI-DANDRUFF BRAID- salicylic acid solution J. Strickland & Co.

Sulfur 8 Anti-Dandruff Braid Spray

Active Ingredients

Salicylic Acid, 2%

Purpose

Antidandruff

Use:

Controls scalp itching and flaking due to dandruff

Warnings:

For external use only

When using this product

• do not get into 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

Keep out of reach of children

If swallowed, get medical help or call a poison control center at once.

Directions

• Spray to apply to the affected area 1-4 times daily, or as directed by a doctor.

Inactive Ingredients

Water, Glycerin, PEG-12 Dimethicone, Propylene Glycol, Sodium Hydroxide, Hydrolyzed Collagen, Disodium EDTA, Diazolidinyl Urea, Fragrance, Methylparaben, Propylparaben.

Package Labeling



Use **Sulfur 8**° for softer more comfortable braids. Beautiful braids without the Itch. Regular use keeps your braids shiny and natural looking.

Drug Facts

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J. Strickland & Co.

P.O. Box 1637 OLIVE BRANCH, MS 38654

Reorder # 445-4

L44592

SULFUR 8 ANTI-DANDRUFF BRAID

salicylic acid solution

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:12022-021

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)			
GLYCERIN (UNII: PDC6A3C0OX)			
PEG-12 DIMETHICONE (LINII) ZEL54N6W95)			

PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
EDETATE DISODIUM ANHYDROUS (UNII: 8NLQ36F6MM)	
DIAZOLIDINYL UREA (UNII: H5RIZ 3MPW4)	
METHYLPARABEN (UNII: A218C7HI9T)	
PROPYLPARABEN (UNII: Z8IX2SC10H)	

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
	NDC:12022-021- 00	356 mL in 1 BOTTLE; Type 0: Not a Combination Product	02/06/2008	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M032	07/14/1994	

Labeler - J. Strickland & Co. (007023112)

Registrant - J. Strickland & Co. (007023112)

Establishment				
Name	Address	ID/FEI	Business Operations	
J. Strickland & Co.		007023112	manufacture(12022-021)	

Revised: 10/2023 J. Strickland & Co.