SULFUR 8 SCALP THERAPY MEDICATED DANDRUFF CONTROL- salicylic acid spray
J. Strickland and Co.

Sulfur 8 Scalp Therapy Medicated Dandruff Control Scalp Spray

Drug Facts

Active Ingredient

Salicylic Acid, 2%

Purpose

Antidandruff

Use:

Controls scalp itching and flaking due to dandruff

Warnings

For external use only

When using this product

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

Stop use and consult 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 at once.

Directions

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

Inactive Ingredients

Water, Glycerin, Sodium Hydroxide, PEG-40 Hydrogenate Castor Oil, PEG-12 Dimethicone, Propylene Glycol, Hydrolyzed Collagen, Diazolidinyl Urea, Disodium EDT, Methylparaben, Propylparaben, Ethylhexylglycerin, PEG-12 Alyl Ether, PEG-12, Butylene Glycol, Phenoxyethanol, Fragrance

Package Labeling:

SULFUR 8 SCALP THERAPY MEDICATED DANDRUFF CONTROL

salicylic acid spray

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:12022-032
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)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
PEG-12 DIMETHICONE (300 CST) (UNII: ZEL54N6W95)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
DIAZOLIDINYL UREA (UNII: H5RIZ3MPW4)	
EDETATE DISODIUM ANHYDROUS (UNII: 8NLQ36F6MM)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
POLYETHYLENE GLYCOL 600 (UNII: NL4J9F21N9)	
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:12022-032-00	355 mL in 1 BOTTLE; 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	

Labeler - J. Strickland and Co. (007023112)

Revised: 10/2023 J. Strickland and Co.