SULFUR 8 SCALP THERAPY MEDICATED DANDRUFF CONTROL- pyrithione zinc shampoo, suspension
J. Strickland and Co.

Sulfur 8 Scalp Therapy Medicated Dandruff Control Shampoo

Drug Facts

Active Ingredient

Pyrithione Zinc, 1%

Purpose

Antidandruff

Use

Controls scalp itching an flaking due to dandruff

Warnings

For external use only

Ask a doctor before use if you have

seborrheic dermatitis 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 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 right away.

Directions

- shake well.
- for best results, use twice a week or as directed by a doctor
- wet hair, apply shampoo. lather, rinse & repeat

Inactive Ingredients

Water, Ammonium Lauryl Sulfate, Propylene Glycol, Ammonium Laureth Sulfate,

Cocamidopropyl Betaine, PEG-3 Glyceryl Cocoate, Hydroxypropyl Methylcellulose, Sodium Cocoamphopropionate, Glycol Distearate, Glycol Stearate, Glycol, Magnesium Aluminum Silicate, Benzyl Alcohol, Methylchloroisothiazolinone, Methylisothiazolinone, Triethylene Glycol, Magnesium Nitrate, Copper Sulfate, Citric Acid, Fragrance, Blue 1

Package Labeling:

SULFUR 8 SCALP THERAPY MEDICATED DANDRUFF CONTROL

pyrithione zinc shampoo, suspension

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:12022-036
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
PYRITHIONE ZINC (UNII: R953O2RHZ5) (PYRITHIONE ZINC - UNII:R953O2RHZ5)	PYRITHIONE ZINC	10 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
AMMONIUM LAURYL SULFATE (UNII: Q7AO2R1M0B)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
COCAMIDOPROPYL BETAINE (UNII: 5OCF3O11KX)	
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
GLYCOL DISTEARATE (UNII: 13W7MDN21W)	
GLYCOL STEARATE (UNII: 0324G66D0E)	
ETHYLENE GLYCOL (UNII: FC72KVT52F)	
MAGNESIUM ALUMINUM SILICATE (UNII: 6M3P64V0NC)	
BENZYL ALCOHOL (UNII: LKG8494WBH)	
METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)	
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	
TRIETHYLENE GLYCOL (UNII: 3P5SU53360)	
MAGNESIUM NITRATE (UNII: 77CBG3UN78)	
CUPRIC SULFATE (UNII: LRX7AJ16DT)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:12022-036-00	280 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: 11/2025 J. Strickland and Co.