GLOVERS CLINICAL CARE SEBORRHEIC DERMATITIS ANTIDANDRUFF- pyrithione zinc shampoo
J. Strickland and Co.

Glover's Clinical Care - Seborrheic Dermatitis, Antidandruff - Shampoo

Drug Facts

Active Ingredient

Pyrithione Zinc, 1%

Purpose

Antidandruff, Seborrheic Dermatitis

Use:

controls scalp itching and flaking due to

• dandruff • seborrheic dermatitis

Warnings:

For external use only

Ask a doctor before use if you have

• serborrheic 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 at least twice a week or as directed by a doctor • wet hair, apply shampoo, lather and rinse.

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, Mthylchloriothiazolinone, Triethylene Glycol, Magnesium Nitrate, Magnesium Chloride, Citric Acid, Fragrance, Blue 1.

Package Labeling:

GLOVERS CLINICAL CARE SEBORRHEIC DERMATITIS ANTIDANDRUFF

pyrithione zinc shampoo

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:12022-038
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)	
TRIETHYLENE GLYCOL (UNII: 3P5SU53360)	
MAGNESIUM NITRATE (UNII: 77CBG3UN78)	
MAGNESIUM CHLORIDE (UNII: 02F3473H9O)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:12022-038-00	118 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/01/2023	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M032	04/01/2023	

Labeler - J. Strickland and Co. (007023112)

Revised: 10/2023 J. Strickland and Co.