WALGREENS CLINICAL STRENGTH MEDICATED DANDRUFF ANTI-DANDRUFF, ANTI-SEBORRHEIC DERMATITIS- selenium sulfide shampoo Walgreen Company

Walgreens Clinical Strength Medicated Dandruff Shampoo Anti-dandruff, Anti-seborrheic dermatitis

Drug Facts

Active ingredient

Selenium Sulfide 1%

Purpose

Anti-dandruff, Anti-seborrheic dermatitis

Uses

Helps prevent recurrence of scalp itching and flaking associated with dandruff and seborrheic dermatitis.

Warnings

For external use only

If condition covers a large area of the body, consult your doctor before using this product.

When using this product

• avoid contact with the 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 contact a Poison Control Center right away.

Directions

- shake well.
- wet hair.
- massage onto scalp.
- rinse.
- repeat if desired.
- for best results use at least twice a week or as directed by a doctor.
- for maximum dandruff control, use every time you shampoo.

Inactive ingredients

water, ammonium laureth sulfate, cocamidopropyl betaine, ammonium lauryl sulfate, glycol distearate, cocamide MEA, ammonium xylenesulfonate, cetyl alcohol, stearyl alcohol, hydroxypropyl methylcellulose, dimethicone, sodium benzoate, fragrance, sodium citrate, disodium EDTA, methylchloroisothiazolinone, methylisothiazolinone, citric acid, sodium chloride, red 40.

Questions or comments?

1-800-925-4733

DISTRIBUTED BY: WALGREEN CO.
200 WILMOT RD., DEERFIELD, IL 60015

PRINCIPAL DISPLAY PANEL - 400 mL Bottle Label

Walgreens

Compare to the active ingredient in Head & Shoulders[®] Clinical Strength^{††}

• WALGREENS • PHARMACIST RECOMMENDED

DANDRUFF SHAMPOO

Clinical Strength

SELENIUM SULFIDE 1%

Medicated

 Strong medicated formula to help treat dandruff & seborrheic dermatitis

05-3271C

13.5 FL OZ (400 mL)



DANDRUFF SHAMPOO

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WALGREENS CLINICAL STRENGTH MEDICATED DANDRUFF ANTI-DANDRUFF, ANTI-SEBORRHEIC DERMATITIS

selenium sulfide shampoo

Product Information Product Type HUMAN OTC DRUG Item Code (Source) NDC:0363-9422 Route of Administration TOPICAL

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
Selenium Sulfide (UNII: 769D9E3810) (Selenium Sulfide - UNII: 769D9E3810)	Selenium Sulfide	10 mg in 1 ml	

Inactive Ingredients

Water (UNII: 059QF0KOOR) Ammonium Laureth-5 Sulfate (UNII: 43Z H89 48) Cocamidopropyl Betaine (UNII: 50CF3011KX) Ammonium Lauryl Sulfate (UNII: Q7AO2R1M0B) Glycol Distearate (UNII: 13W7MDN21W) COCO MONOETHANOLAMIDE (UNII: C80684146D) Ammonium Xylenesulfonate (UNII: 4FZY6L6XCM) Cetyl Alcohol (UNII: 936)ST6jCN) Stearyl Alcohol (UNII: 2KR89 4H1Y) Hypromellose, Unspecified (UNII: 3NXW29V3WO) Dimethicone (UNII: 92RU3N3Y1O) Sodium Benzoate (UNII: Oj245FE5EU) SODIUM CITRATE, UNSPECIFIED FORM (UNII: 1Q73Q2JULR) EDETATE DISODIUM (UNII: 7FLD91C86K) Methylchloroisothiazolinone (UNII: 298DP1CFA) Citric Acid Monohydrate (UNII: 2968PHW8QP) Sodium Chloride (UNII: 451W47IQ8X) FD&C RED NO. 40 (UNII: WZB9127XOA)	Ingredient Name	Strength
Cocamidopropyl Betaine (UNII: 5OCF3O11KX) Ammonium Lauryl Sulfate (UNII: Q7AO2R1M0B) Glycol Distearate (UNII: 13W7MDN21W) COCO MONOETHANOLAMIDE (UNII: C80684146D) Ammonium Xylenesulfonate (UNII: 4FZY6L6XCM) Cetyl Alcohol (UNII: 936JST6JCN) Stearyl Alcohol (UNII: 2KR89I4H1Y) Hypromellose, Unspecified (UNII: 3NXW29V3WO) Dimethicone (UNII: 92RU3N3Y1O) Sodium Benzoate (UNII: OJ245FE5EU) SODIUM CITRATE, UNSPECIFIED FORM (UNII: 1Q73Q2JULR) EDETATE DISODIUM (UNII: 7FLD91C86K) Methylchloroisothiazolinone (UNII: DEL7T5QRPN) Methylisothiazolinone (UNII: 2990DE1QFA) Citric Acid Monohydrate (UNII: 2968PHW8QP) Sodium Chloride (UNII: 451W47IQ8X)	Water (UNII: 059QF0KO0R)	
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Sodium Chloride (UNII: 451W47IQ8X)	Methylisothiazolinone (UNII: 229D0E1QFA)	
	Citric Acid Monohydrate (UNII: 2968PHW8QP)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	Sodium Chloride (UNII: 451W47IQ8X)	
	FD&C RED NO. 40 (UNII: WZB9127XOA)	

Product Characteristics			
Color	ORANGE	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

Packaging				
	# Item Code	Package Description	Marketing Start Date	Marketing End Date
	1 NDC:0363-9422-01	400 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	12/14/2016	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M032	12/14/2016	

Labeler - Walgreen Company (008965063)

Registrant - Garcoa, Inc. (036464697)

Establishment			
Na me	Address	ID/FEI	Business Operations

Revised: 5/2024 Walgreen Company