

DHS ZINC- pyrithione zinc shampoo

Person and Covey

DHS ZINC

WARNINGS AND PRECAUTIONS

For external use only. Ask a physician before use if you have Psoriasis or Seborrheic Dermatitis that covers a large area of the body. Do not get into eyes. If contact occurs, rinse eyes thoroughly with water. Stop use and ask a physician if condition worsens or does not improve after regular use.

Dosage and Administration

Shake well before using. Wet hair thoroughly. Apply a liberal amount of shampoo and massage into a rich lather. Allow lather to remain on scalp for several minutes. Rinse hair well and repeat application. For best results use at least twice a week or as directed by a physician.

Indications and Use

Helps prevent recurrence of scalp itching and flaking associated with Seborrheic Dermatitis and Dandruff.

OTC - KEEP OUT OF REACH OF CHILDREN

Yes. If swallowed, get medical help or contact a Poison Control Center right away.

PURPOSE

Recommended for control of Dandruff of Seborrheic Dermatitis

INACTIVE INGREDIENTS

Purified Water

TEA Lauryl Sulfate

Cocamidopropyl Hydroxysultaine

Sodium Chloride

PEG 8 Distearate

Methocel E4M Premium

Veegum

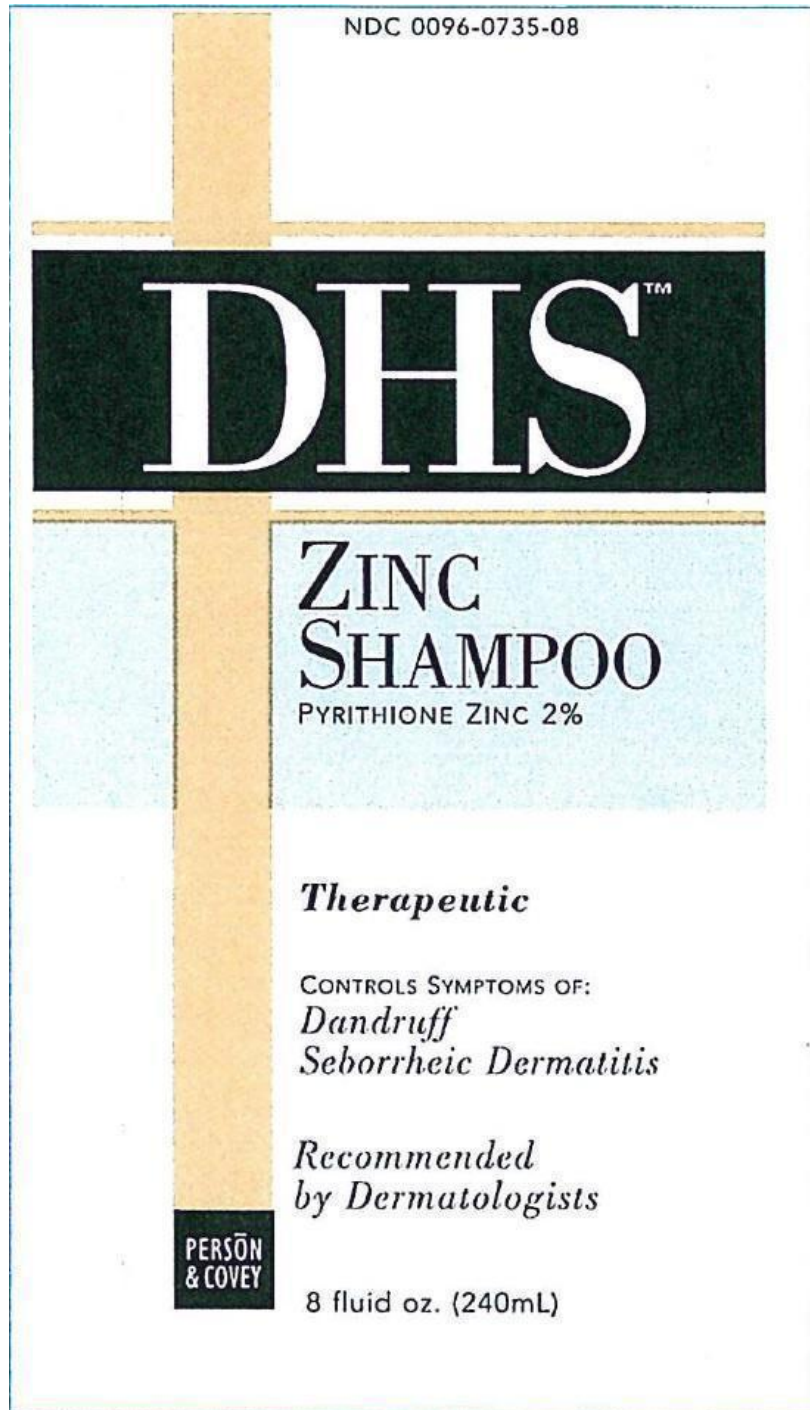
Herbal Bouquet No. 1751 Perfume

Active Ingredient

pyrithione zinc

PACKAGE LABEL.PRINCIPAL DISPLAY PANEL

DHS Zinc 8oz.jpg



Principal Display Panel



DHS ZINC

pyrithione zinc shampoo

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:0096-0735
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
PYRITHIONE ZINC (UNII: R953O2RHZ5) (PYRITHIONE ZINC - UNII:R953O2RHZ5)	PYRITHIONE ZINC	20 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
COCAMIDOPROPYL HYDROXYSULTAINE (UNII: 62V75NI93W)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
PEG-8 DISTEARATE (UNII: 7JNC8VN07M)	
HYPROMELLOSES (UNII: 3NXW29V3WO)	
MAGNESIUM ALUMINUM SILICATE (UNII: 6M3P64V0NC)	
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	
TRIETHANOLAMINE LAURYL SULFATE (UNII: E8458C1KAA)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0096-0735-08	240 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/01/1995	
2	NDC:0096-0735-16	480 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/01/1995	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M032	06/01/1995	

Labeler - Person and Covey (008482473)

Establishment

Name	Address	ID/FEI	Business Operations
Person and Covey		008482473	manufacture(0096-0735)

Revised: 7/2025

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