HEAD AND SHOULDERS CLINICAL DANDRUFF DEFENSE DRY SCALP RESCUE-pyrithione zinc lotion
The Procter & Gamble Manufacturing Company

Head and Shoulders ®

Clinical Dandruff Defense Dry Scalp Rescue

Drug Facts

Active ingredient

Pyrithione zinc .75%

Purpose

Anti-dandruff

Uses

helps prevent recurrence of flaking and itching associated with dandruff.

Warnings

For external use only.

When using this product
- avoid contact with 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 this and all drugs out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

Directions
- for best results use at least twice a week or as directed by a doctor.
- apply to wet hair after shampooing by gently massaging into hair and scalp, rinse well.
- for maximum dandruff control, use every time you shampoo.

Inactive ingredients

Water, stearyl alcohol, cetyl alcohol, stearamidopropyl dimethylamine, glutamic acid, dimethicone, fragrance, phenoxyethanol, benzyl alcohol, sodium chloride, citric acid, honey extract,methylchloroisothiazolinone, methylisothiazolinone.

Questions (or comments)?

1-800-723-9569

Dist. by PROCTER & GAMBLE,
CINCINNATI, OH 45202

PRINCIPAL DISPLAY PANEL - TUBE LABEL

head &

shoulders ®
pyrithione zinc dandruff conditioner

Clinical

Dandruff

Defense

Dry Scalp

Rescue

with Manuka Honey

100% DANDRUFF PROTECTION SATISFACTION GUARANTEED*

9.1 FL OZ (270 mL)

HEAD AND SHOULDERS CLINICAL DANDRUFF DEFENSE DRY SCALP RESCUE

pyrithione zinc lotion

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:69423-568
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
	Basis of	

Ingredient Name	Basis of Strength	Strength
PYRITHIONE ZINC (UNII: R953O2RHZ5) (PYRITHIONE ZINC - UNII:R953O2RHZ5)	PYRITHIONE ZINC	0.75 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)	
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	
STEARAMIDOPROPYL DIMETHYLAMINE (UNII: K7VEI00UFR)	
GLUTAMIC ACID (UNII: 3KX376GY7L)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
BENZYL ALCOHOL (UNII: LKG8494WBH)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
HONEY (UNII: Y9H1V576FH)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
STEARYL ALCOHOL (UNII: 2KR89I4H1Y)	
CETYL ALCOHOL (UNII: 936JST6JCN)	
WATER (UNII: 059QF0KO0R)	
DIMETHICONE (UNII: 92RU3N3Y1O)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69423-568-27	270 mL in 1 TUBE; Type 0: Not a Combination Product	01/11/2021	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M032	01/11/2021	

Labeler - The Procter & Gamble Manufacturing Company (004238200)

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