HEAD AND SHOULDERS HAIR AND SCALP CONDITIONER CLASSIC CLEAN- pyrithione zinc lotion
The Procter & Gamble Manufacturing Company

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

Head and Shoulders ® Hair and Scalp Conditioner

Classic Clean

Drug Facts

Active ingredient

Pyrithione zinc 0.5%

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, rinsewell.

Inactive ingredients

Water, stearyl alcohol, cetyl alcohol, stearamidopropyl dimethylamine, dimethicone,

fragrance, glutamic acid, phenoxyethanol, benzyl alcohol, citric acid, sodium chloride, methylchloroisothiazolinone, methylisothiazolinone.

Questions (or comments)?

1-800-723-9569

Dist. by PROCTER & GAMBLE,
CINCINNATI, OH 45202

PRINCIPAL DISPLAY PANEL - 325 mL Tube Label

head &
shoulders **®**

pyrithione zinc **dandruff conditioner**

classic clean

DAILY HAIR & SCALP

CONDITIONER

10.9 FL OZ (325 mL)

HEAD AND SHOULDERS HAIR AND SCALP CONDITIONER CLASSIC CLEAN

pyrithione zinc lotion

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:69423-171
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
PYRITHIONE ZINC (UNII: R953O2RHZ5) (PYRITHIONE ZINC - UNII:R953O2RHZ5)	PYRITHIONE ZINC	0.5 g in 100 mL

Inactive Ingredients

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

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69423-171-10	10 mL in 1 POUCH; Type 0: Not a Combination Product	10/01/2016	
2	NDC:69423-171-40	400 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	10/01/2016	11/26/2020
3	NDC:69423-171-32	325 mL in 1 TUBE; Type 0: Not a Combination Product	11/29/2018	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	M032	10/01/2016	

Labeler - The Procter & Gamble Manufacturing Company (004238200)

Revised: 2/2023 The Procter & Gamble Manufacturing Company