HEAD AND SHOULDERS CONDITIONER SMOOTH AND SILKY- 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 ® Conditioner

Smooth & Silky

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 <u>at least</u> twice a week or as directed by a doctor.
- apply to wet hair after shampooing by gently massaging into hair and scalp, rinse well.

Inactive ingredients

Water, stearyl alcohol, cetyl alcohol, stearamidopropyl dimethylamine, glutamic acid,

dimethicone, fragrance, 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 - 315 mL Tube Label

head &

shoulders ®

pyrithione zinc dandruff conditioner

smooth & silky HAIR & SCALP CONDITIONER

24 hour frizz control from root to tip 10.6 FL OZ (315 mL)



HEAD AND SHOULDERS CONDITIONER SMOOTH AND SILKY

pyrithione zinc lotion

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Prod	uct	INTOFM	ation

Product Type HUMAN OTC DRUG Item Code (Source) NDC:69423-103

Route of Administration TOPICAL

Active Ingredient/Active Moiety Ingredient Name Basis of Strength PYRITHIONE ZINC (UNII: R95302RHZ5) (PYRITHIONE ZINC - UNII: R95302RHZ5) PYRITHIONE ZINC (UNII: R95302RHZ5) 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- 103-40	400 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	11/10/2016	12/22/2019	
2	NDC:69423- 103-68	680 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	11/10/2016	09/08/2019	
3	NDC:69423- 103-90	90 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	11/10/2016	01/15/2021	
4	NDC:69423- 103-65	650 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	09/11/2017	10/03/2020	
5	NDC:69423- 103-38	380 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	01/09/2018	10/31/2020	
6	NDC:69423- 103-10	10 mL in 1 POUCH; Type 0: Not a Combination Product	11/10/2017		
7	NDC:69423- 103-25	250 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	04/14/2018	07/20/2020	
8	NDC:69423- 103-61	610 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	11/29/2018	09/01/2024	
9	NDC:69423- 103-31	315 mL in 1 TUBE; Type 0: Not a Combination Product	11/29/2018		
10	NDC:69423- 103-59	592 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	12/15/2022		

Marketing Information					
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date		
OTC monograph final	M032	11/10/2016			

Labeler - The Procter & Gamble Manufacturing Company (004238200)

Revised: 4/2023 The Procter & Gamble Manufacturing Company