

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, rinse well.

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 & shoulders

classic clean
pyrithione zinc
dandruff conditioner

PROVEN
PROTECTION
FLUORENOLONE

- 1) INTENSIVE MOISTURIZATION - for soft, manageable hair
- 2) SCALP PROTECTION - soothes itchy, dry scalp - GUARANTEED*
- 3) HEALTHY HAIR - grows from a healthy scalp

#1 DERMATOLOGIST RECOMMENDED

Drug Facts

Active Ingredient

Pyrrithione zinc 1.5%

Purpose

Anti-dandruff

Uses: helps prevent recurrence of itching 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:

- irritation persists or worsens or appears after regular use of this product is discontinued.

Keep this and all drugs out of the 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 liberally after shampooing gently using a hair brush or comb.

Inactive Ingredients: Water, cetyl alcohol, cetyl laurate, cocamidopropyl dimethylamine, dimethylsiloxane, fragrance, glycerin, hydroxyethylcellulose, isopropyl alcohol, isopropyl myristate, isopropyl palmitate, isopropyl stearate, isopropyl myristate, isopropyl palmitate, isopropyl stearate, isopropyl myristate, isopropyl palmitate, isopropyl stearate.

Questions (or comments)? 1-800-723-0500

MADE IN U.S.A. of U.S. and/or imported ingredients.

Dist. by PROCTER & GAMBLE, CINCINNATI, OH 45202

Patent: www.pg.com/patents

P&G

www.pg.com

www.headandshoulders.com

*Surface tension is measured. If you are not satisfied with the product, it is guaranteed to be returned.

†Based on volume of use.

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CLASSIC
CLEAN
DAILY HAIR & SCALP
CONDITIONER



AMERICA'S #1 DANDRUFF
CONDITIONER BRAND**

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