

**HEAD AND SHOULDERS DEEP MOISTURE HAIR AND SCALP CONDITIONER-
pyrithione zinc lotion
The Procter & Gamble Manufacturing Company**

Head and Shoulders ® Deep Moisture 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, glutamic acid, dimethicone, fragrance, phenoxyethanol, benzyl alcohol, caffeine, citric acid, sodium

chloride, persea gratissima (avocado) oil, 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**

deep moisture

DAILY HAIR & SCALP

CONDITIONER

enriched moisture for scalp & hair

infused with avocado oil

10.6 FL OZ (315 mL)

head & shoulders deep moisture
pyrithione zinc
dandruff conditioner

PROVEN PROTECTION
FLAKE-FREE HAIR CARE

- 1) INTENSE MOISTURIZATION - for more hydrated hair from root to tip
- 2) SCALP PROTECTION - from flaking, itch, & dryness - "GLASS WIPES"
- 3) BEAUTIFUL HAIR - grows from a healthy scalp

SAFE FOR COLOR TREATED HAIR*

Drug Facts

Active Ingredient	Purpose
Pyrithione zinc 1.5%	Anti-dandruff

Use 5 to help prevent recurrence of flaking and itch by consistent use with each use.

Warnings

For external use only.

When using to prevent dandruff, use 5 to 7 times a week. If contact occurs, rinse eyes thoroughly with water.

Stop use and ask a doctor if:

- condition worsens or does not improve after use of this product for 2 weeks.
- keep this and all drugs in a cool, dry place at 70-80°F. If needed, get medical help or contact a Poison Control Center right away.

Directions

- for best results use 5 to 7 times a week or as directed by a doctor.
- apply to wet hair after shampooing thoroughly with shampoo, then wet.

Inactive Ingredients Water, stearyl alcohol, cetyl alcohol, stearamidopropyl dimethylamine, glutaric acid, dimethylsiloxane, hexoic acid, glycerin, methanol, benzyl alcohol, caffeine, citric acid, sodium chloride, potassium citrate (citric acid salt), methyl t. Monothiuronium sulfonium, methyl methacrylate.

Questions for comments? 1-800-4-723-9540

MADE IN U.S.A. or U.S. and for important ingredients list by PROCTER & GAMBLE, CINCINNATI, OH 45202

Patent: www.pg.com/patents

P&G
www.pg.com

www.headandshoulders.com

*On the back of the bottle, a patch test is recommended where this product will be applied. Do a patch test on a small area of skin first. Call 1-800-4-723-9540 for more details.

**Visible flaking, or itch, may occur when associated with dry scalp. Shampoo & conditioner separate in non-conditioning use spots.



Infused With More Moisturizers

head & shoulders
PYRITHIONE ZINC DANDRUFF CONDITIONER

DEEP MOISTURE
DAILY HAIR & SCALP CONDITIONER



ENRICHED MOISTURE FOR SCALP & HAIR
INFUSED WITH AVOCAO OIL

10.6 FL OZ (315 mL)

HEAD AND SHOULDERS DEEP MOISTURE HAIR AND SCALP CONDITIONER			
pyrithione zinc lotion			
Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:69423-197
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
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	
CAFFEINE (UNII: 3G6A5W338E)	
AVOCADO OIL (UNII: 6VNO72PFC1)	
WATER (UNII: 059QF0K00R)	
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)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69423-197-38	380 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	09/11/2017	10/24/2020
2	NDC:69423-197-31	315 mL in 1 TUBE; Type 0: Not a Combination Product	11/29/2018	10/01/2026

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M032	09/11/2017	10/01/2026

Labeler - The Procter & Gamble Manufacturing Company (004238200)

Establishment

Name	Address	ID/FEI	Business Operations
The Procter & Gamble Manufacturing Company		081329183	manufacture(69423-197)

Revised: 12/2025

The Procter & Gamble Manufacturing Company