HEAD AND SHOULDERS ROYAL OILS DAILY MOISTURE SCALP- pyrithione zinc cream
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 ® Royal Oils Daily Moisture Scalp Cream

Drug Facts

Active ingredient

Pyrithione zinc 0.1%

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

• Apply to affected areas one to four times daily or as directed by a doctor.

• Twist the cap open. Place tip directly onto scalp (damp or dry)

• Gently squeeze the bottle while moving tip fromfront to back of your scalp. Repeat to cover your entire scalp.

• Massage for even distribution. Do not rinse.

Inactive ingredients

Water, cocos nucifera (coconut) oil, cetyl alcohol, stearamidopropyl dimethylamine, fragrance, stearyl alcohol, quaternium-18, hydroxyethylcellulose, phenoxyethanol, benzyl alcohol, dimethicone, PEG-2M, cetearyl alcohol, methylparaben, propylparaben, oleyl alcohol, glyceryl stearate, citric acid, polysorbate 60.

Questions (or comments)?

1-800-723-9569

Dist. by PROCTER & GAMBLE,
CINCINNATI, OH 45202

PRINCIPAL DISPLAY PANEL - 150 mL Bottle Label

head &
shoulders ®
pyrithione zinc **dandruff treatment**

ROYAL OILS

DAILY MOISTURE

SCALP CREAM

WITH COCONUT OIL

MOISTURIZES SCALP TO

PREVENT DRYNESS AND ITCH*

SULFATE, DYE &

MINERAL OIL FREE

5.0 FL OZ
(150 mL)

HEAD AND SHOULDERS ROYAL OILS DAILY MOISTURE SCALP

pyrithione zinc cream

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:69423-307
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
PYRITHIONE ZINC (UNII: R953O2RHZ5) (PYRITHIONE ZINC - UNII:R953O2RHZ5)	PYRITHIONE ZINC	0.1 g in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**
POLYSORBATE 60 (UNII: CAL22UVI4M)	
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
OLEYL ALCOHOL (UNII: 172F2WN8DV)	

POLYETHYLENE OXIDE 100000 (UNII: V46Y6OJ5QB)	
WATER (UNII: 059QF0KO0R)	
STEARYL ALCOHOL (UNII: 2KR89I4H1Y)	
CETYL ALCOHOL (UNII: 936JST6JCN)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
BENZYL ALCOHOL (UNII: LKG8494WBH)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
STEARAMIDOPROPYL DIMETHYLAMINE (UNII: K7VEI00UFR)	
QUATERNIUM-18 (UNII: O7757NO1VL)	
COCONUT OIL (UNII: Q9L0O73W7L)	
GLYCOL STEARATE (UNII: 0324G66D0E)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69423-307-15	150 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	08/30/2018	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part358H	08/30/2018	

Labeler - The Procter & Gamble Manufacturing Company (004238200)

Revised: 4/2019 The Procter & Gamble Manufacturing Company