

HEAD AND SHOULDERS SUPREME SCALP MOISTURIZER WITH HEMP SEED-pyrrithione 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 ® Supreme Scalp Moisturizer Hemp Seed Essential Oil

Drug Facts

Active ingredient

Pyrrithione 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.

Inactive ingredients

Water, cannabis sativa seed oil, cetyl alcohol, stearamidopropyl dimethylamine, stearyl alcohol, hydroxyethylcellulose, quaternium-18, phenoxyethanol, fragrance, benzyl alcohol, PEG-2M, cetearyl alcohol, methylparaben, dimethicone, glyceryl stearate, oleyl alcohol, propylparaben, citric acid, polysorbate 60.

Questions (or comments)?

1-800-723-9569

Dist. by PROCTER & GAMBLE,
CINCINNATI, OH 45202

PRINCIPAL DISPLAY PANEL - 50 mL bottle label

Head & Shoulders

PYRITHIONE ZINC DANDRUFF TREATMENT

SUPREME SCALP MOISTURIZER

HEMP OIL ESSENTIAL OIL

NOURISHES DRY SCALP,

HELPS STRENGTHEN HAIR

1.69 FL OZ (50 mL)

SUPREME
FAST ABSORBING - WEIGHTLESS - NOURISHING

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head & shoulders
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MADE IN U.S.A. of U.S.
and/or Imported Ingredients
Dist. by PROCTER & GAMBLE,
CINCINNATI, OH 45202
www.headandshoulders.com
Patents: www.pg.com/patents

P&G
www.pg.com

91436361

0 37000 67961 5

**HEAD AND SHOULDERS SUPREME SCALP MOISTURIZER WITH
HEMP SEED**

pyrithione zinc lotion

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:69423-569
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
STEARAMIDOPROPYL DIMETHYLAMINE (UNII: K7VEI00UFR)	
HYDROXYETHYL CELLULOSE, UNSPECIFIED (UNII: T4V6TWG28D)	
POLYSORBATE 60 (UNII: CAL22UVI4M)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
POLYETHYLENE OXIDE 100000 (UNII: V46Y6OJ5QB)	
OLEYL ALCOHOL (UNII: 172F2WN8DV)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
QUATERNIUM-18 (UNII: O7757NO1VL)	
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)	
STEARYL ALCOHOL (UNII: 2KR89I4H1Y)	
CETYL ALCOHOL (UNII: 936JST6JCN)	
WATER (UNII: 059QF0KO0R)	
CANNABIS SATIVA SEED OIL (UNII: 69VJ1LPN1S)	
BENZYL ALCOHOL (UNII: LKG8494WBH)	
GLYCERYL MONOSTEARATE (UNII: 230OU9XXE4)	
DIMETHICONE (UNII: 92RU3N3Y1O)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69423-569-50	50 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	02/01/2021	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part358H	02/01/2021	

Revised: 7/2021

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