HEAD AND SHOULDERS OCEAN LIFT- pyrithione zinc lotion/shampoo
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 ®

Ocean Lift

Drug Facts

Active ingredient

Pyrithione zinc 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

- for best results use at least twice a week or as directed by a doctor.
- for maximum dandruff control, use every time you shampoo.
- shake before use.
- wet hair, massage onto scalp, rinse, repeat if desired.

Inactive ingredients

Water, sodium lauryl sulfate, sodium laureth sulfate, glycol distearate, zinc carbonate, sodium chloride, fragrance, sodium xylenesulfonate, cocamidopropyl betaine, dimethicone, sodium benzoate, guar hydroxypropyltrimonium chloride, magnesium carbonate hydroxide, methylchloroisothiazolinone, methylisothiazolinone, blue 1, yellow 5.

Questions (or comments)?

1-800-723-9569

Dist. by PROCTER & GAMBLE,
CINCINNATI, OH 45202

PRINCIPAL DISPLAY PANEL - 700 mL Bottle Label

3 Action

FORMULA

head &

shoulders ®

pyrithione zinc **dandruff shampoo**

ocean lift

for an invigorating scent

FLAKE FREE.*

UP TO 100%

GUARANTEED^

23.7 FL OZ

(700 mL)

HEAD AND SHOULDERS OCEAN LIFT

pyrithione zinc lotion/shampoo

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:37000-100
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
PYRITHIONE ZINC (UNII: R953O2RHZ5) (PYRITHIONE ZINC - UNII:R953O2RHZ5)	PYRITHIONE ZINC	1 g in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
SODIUM LAURYL SULFATE (UNII: 368GB5141J)	

SODIUM LAURETH-3 SULFATE (UNII: BPV390UAP0)	
GLYCOL DISTEARATE (UNII: 13W7MDN21W)	
ZINC CARBONATE (UNII: EQR32Y7H0M)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
SODIUM XYLENESULFONATE (UNII: G4LZF950UR)	
COCAMIDOPROPYL BETAINE (UNII: 5OCF3O11KX)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
MAGNESIUM CARBONATE HYDROXIDE (UNII: YQO029V1L4)	
METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)	
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:37000-100-70	700 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	09/01/2013	
2	NDC:37000-100-40	400 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	12/14/2014	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part358H	09/01/2013	

Labeler - Procter & Gamble Manufacturing Company (004238200)

Establishment

Name	Address	ID/FEI	Business Operations
Arch Chemicals, Inc.		002220804	api manufacture(37000-100)

Revised: 10/2016 Procter & Gamble Manufacturing Company