

HEAD AND SHOULDERS HEADSTRONG DAILY DANDRUFF- pyrithione zinc shampoo
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 ®
#HEADSTRONG DAILY DANDRUFF SHAMPOO

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 laureth sulfate, cocamide MEA, sodium xylenesulfonate, zinc carbonate, glycol distearate, sodium lauryl sulfate, cocamidopropyl betaine, sodium chloride, fragrance, dimethicone, guar hydroxypropyltrimonium chloride, sodium benzoate, stearyl alcohol, magnesium carbonate hydroxide, cetyl alcohol, polyquaternium-76, prunus amygdalus dulcis (sweet almond) oil, methylchloroisoithiazolinone, methylisothiazolinone.

Questions (or comments)?

1-800-723-9569

Dist. by PROCTER & GAMBLE,
CINCINNATI, OH 45202

PRINCIPAL DISPLAY PANEL - 950 mL Bottle Label

**head &
shoulders**®

pyrithione zinc dandruff shampoo

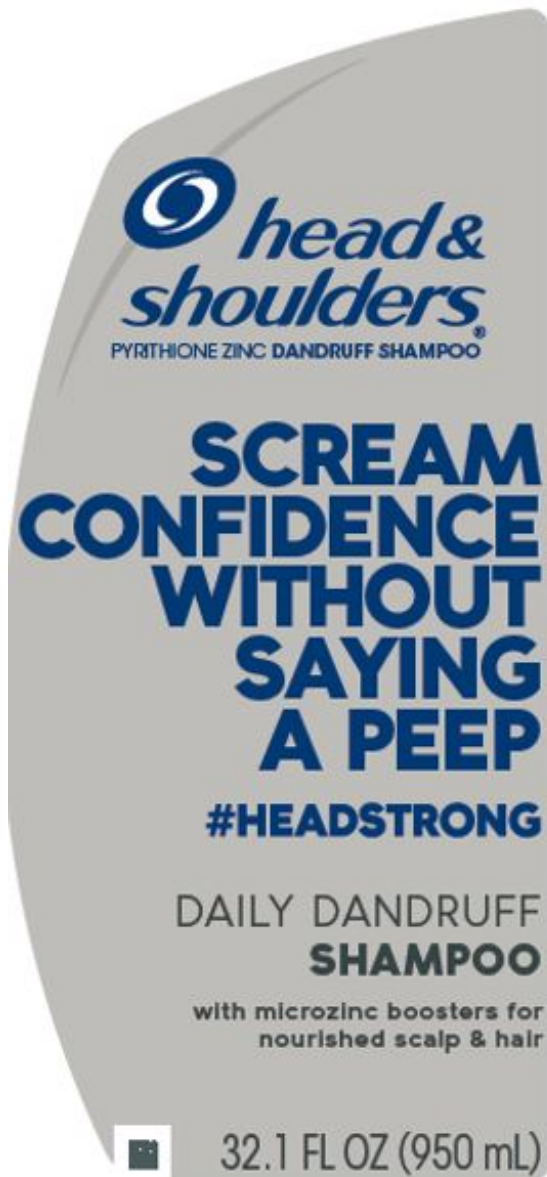
#HEADSTRONG

Daily Dandruff Shampoo

with microzinc boosters for

nourished scalp & hair

32.1 FL OZ (950 mL)



HEAD AND SHOULDERS HEADSTRONG DAILY DANDRUFF
pyrithione zinc shampoo

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:69423-323
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
GUAR HYDROXYPROPYLTRIMONIUM CHLORIDE (1.7 SUBSTITUENTS PER SACCHARIDE) (UNII: B16G315W7A)	
WATER (UNII: 059QF0K00R)	
SODIUM LAURETH-3 SULFATE (UNII: BPV390UAP0)	
SODIUM LAURYL SULFATE (UNII: 368GB5141J)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
COCO MONOETHANOLAMIDE (UNII: C80684146D)	
ZINC CARBONATE (UNII: EQR32Y7H0M)	
GLYCOL DISTEARATE (UNII: 13W7MDN21W)	
SODIUM XYLENESULFONATE (UNII: G4LZF950UR)	
CETYL ALCOHOL (UNII: 936JST6JCN)	
SODIUM CHLORIDE (UNII: 451W471Q8X)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
MAGNESIUM CARBONATE HYDROXIDE (UNII: YQO029V1L4)	
ALMOND OIL (UNII: 66YXD4DKO9)	
METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)	
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	
STEARYL ALCOHOL (UNII: 2KR89I4HIY)	
COCAMIDOPROPYL BETAINE (UNII: 5OCF3O11KX)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69423-323-40	400 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	12/31/2018	
2	NDC:69423-323-95	950 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	12/31/2018	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part358H	12/31/2018	

Revised: 2/2020

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