HEAD AND SHOULDERS HAIR LOSS PREVENTION- pyrithione zinc shampoo
ALL NATURAL DYNAMICS

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 hair loss prevention

Drug Facts

Active ingredient

Pyrithione zinc 1%

Purpose

Anti-dandruff

Uses

helps prevent recurrence of flaking and itching associated with dandruff.

helps prevent hair loss

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

Inactive ingredients

Water, Sodium laureth sulfate, Sodium lauryl sulfate, Coco Monoethanolamide, Zinc carbonate, Glycol distearate, Dimethicone, Guar hydroxypropyltrimonium chloride, Magnesium sulfate, Sodium benzoate, Magnesium carbonate hydroxide, Benzyl alcohol, Tocopheryl acetate, FD&C Yellow No. 5, Methylchloroisothiazolinone, Methylisothiazolinone, FD&C Blue no. 1

Questions (or comments)?

01-800-717-2413

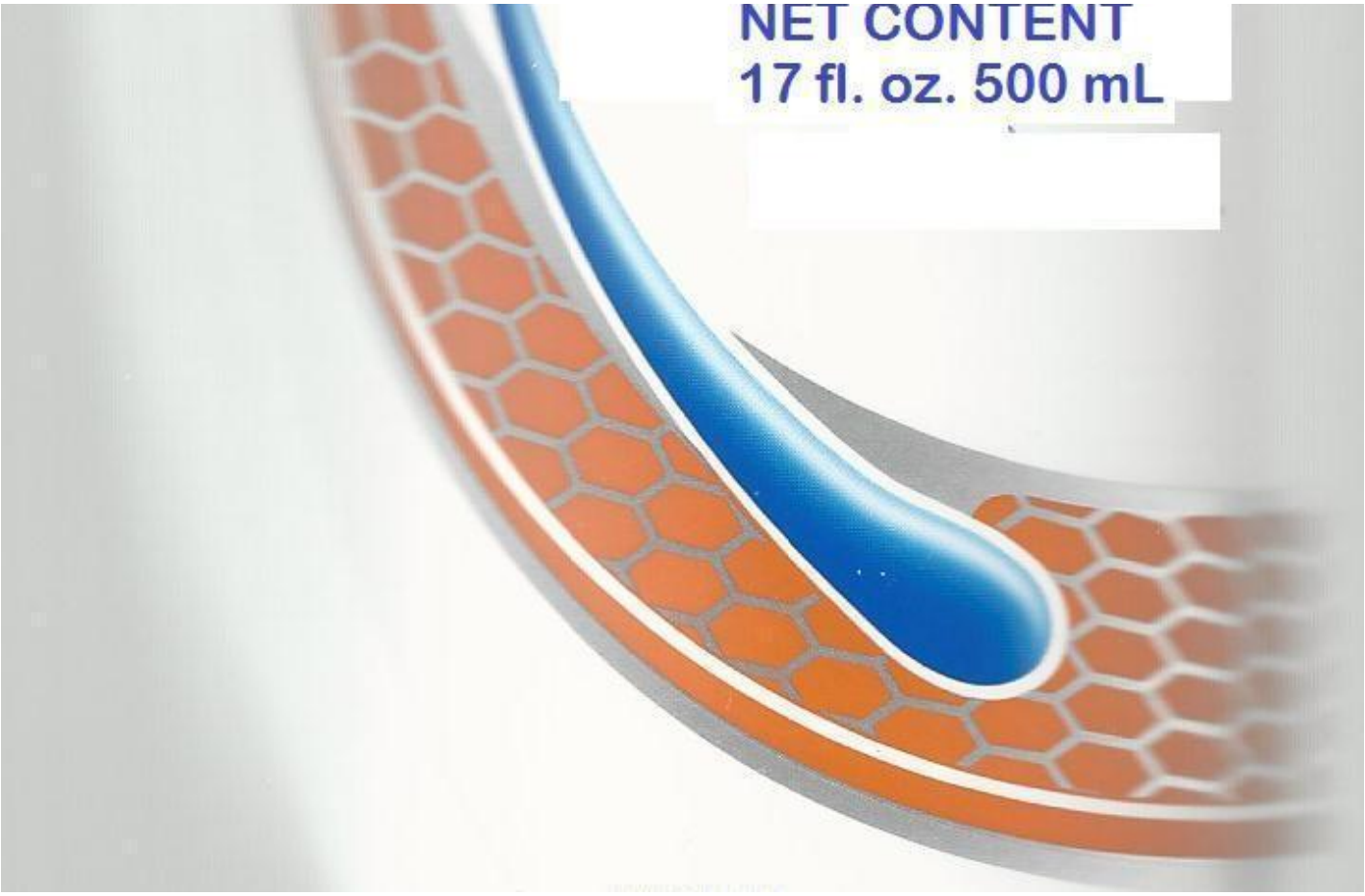
Dist. by

Procter & Gamble Manufactura, S. de R.L. de C.V.

San Andr`s Atoto No. 326, Naucalpan de Juarez,

Estado de Mexico, MX

www.headandshoulders.com



HEAD AND SHOULDERS HAIR LOSS PREVENTION

pyrithione zinc shampoo

Product Information

Product Type	HUMAN OTC DRUG LABEL	**Item Code (Source)**	NDC:51769-140
Route of Administration	TOPICAL	**DEA Schedule**	

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
Pyrithione Zinc (PYRITHIONE ZINC)	Pyrithione Zinc	0.01 g in 1 mL

Inactive Ingredients

Ingredient Name	Strength
Water	
Sodium laureth sulfate	
Sodium lauryl sulfate	
Coco Monoethanolamide	
Zinc carbonate	
Glycol distearate	
Dimethicone	
Guar hydroxypropyltrimonium chloride (1.7 substituents per saccharide)	

Magnesium sulfate	
Sodium benzoate	
magnesium carbonate hydroxide	
Benzyl alcohol	
.ALPHA.-TOCOPHEROL ACETATE	
FD&C Yellow No. 5	
methylchloroisothiazolinone	
methylisothiazolinone	
FD&C Blue no. 1	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:51769-140-50	500 mL in 1 BOTTLE, PLASTIC		
2	NDC:51769-140-40	400 mL in 1 BOTTLE, PLASTIC		
3	NDC:51769-140-20	200 mL in 1 BOTTLE, PLASTIC		
4	NDC:51769-140-11	1000 mL in 1 BOTTLE, PLASTIC		
5	NDC:51769-140-18	1180 mL in 1 BOTTLE, PLASTIC		
6	NDC:51769-140-05	50 mL in 1 BOTTLE, PLASTIC		
7	NDC:51769-140-01	10 mL in 1 POUCH		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part358H	02/01/2013	

Labeler - ALL NATURAL DYNAMICS (962732892)

Establishment

Name	Address	ID/FEI	Business Operations
Procter & Gamble Manufactura, S. de R.L. de C.V.		812807550	manufacture(51769-140)