EPOCH AVA PUHI MONI- pyrithione zinc shampoo NSE Products, Inc.

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.

epoch_® Ava puhi moni[®] Anti-Dandruff Shampoo

Drug Facts

Active Ingredients

Zinc Pyrithione 1%

Purpose

Anti-dandruff

Use

Controls the symptoms of dandruff.

Warning

For external use only

- **Keep out of the reach of children.** If swallowed, get medical help or contact a Poison Control Center right away.
- **Avoid contact with eyes.** If contact occurs, rinse eyes thoroughly with water.
- If condition worsens or does not improve after regular use of the product as directed, consult a
 doctor.

Directions

- Wet hair, lather thoroughly
- Leave in 1–2 minutes and rinse well
- For best results, use daily or at least twice a week (or as directed by a doctor).

Inactive Ingredients

Water (Aqua), Ammonium Lauryl Sulfate, Ammonium Laureth Sulfate, Acrylates/Aminoacrylates/C10-30 Alkyl PEG-20 Itaconate Copolymer, Cocamide MEA, Cocamide MIPA, Dimethicone, Zingiber Zerumbet Juice, Clematis Vitalba Leaf Extract, Mahonia Aquifolium Root Extract, Salvia Officinalis (Sage) Leaf Extract, Urtica Dioica (Nettle) Extract, Panthenol, Butylene Glycol, Sodium Chloride, Citric Acid, Tetrasodium EDTA, Fragrance (Parfum), Methylchloroisothiazolinone, Methylisothiazolinone, Chlorophyllin-Copper Complex (CI 75810).

Questions?

PRINCIPAL DISPLAY PANEL - 250 ml Bottle Label

epoch_®

Ava puhi moni®

Anti-Dandruff Shampoo

the "true ava puhi" 250 ml e (8.4 fl. oz.)



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Questions? 1-888-742-7626

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EPOCH AVA PUHI MONI

pyrithione zinc shampoo

Product Informa	tıo	n
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Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:62839-2810

Route of Administration TOPICAL

Active Ingredient/Active Moiety		
Ingredient Name	Basis of Strength	Strength
Pyrithione Zinc (UNII: R953O2RHZ5) (Pyrithione Zinc - UNII:R953O2RHZ5)	Pyrithione Zinc	10 mL in 1000 mL

Inactive Ingredients	
Ingredient Name	Strength
Water (UNII: 059QF0KO0R)	
Ammonium Lauryl Sulfate (UNII: Q7AO2R1M0B)	
Coco Monoethanolamide (UNII: C80684146D)	
Dimethicone (UNII: 92RU3N3Y1O)	
Clematis Vitalba Leaf (UNII: 526T95850X)	
Mahonia Aquifolium Root (UNII: 746TB9VNDP)	
Sage (UNII: 065C5D077J)	
Urtica Dioica (UNII: 710 FLW4U46)	
Panthenol (UNII: WV9CM0O67Z)	
Butylene Glycol (UNII: 3XUS85K0RA)	
Sodium Chloride (UNII: 451W47IQ8X)	
Citric Acid Monohydrate (UNII: 2968PHW8QP)	
Edetate Sodium (UNII: MP1J8420LU)	
Methylchloroisothiazolinone (UNII: DEL7T5QRPN)	
Methylisothiazolinone (UNII: 229 D0 E1QFA)	
Sodium Copper Chlorophyllin (UNII: 1D276TYV9O)	

Packaging			
# Item Code	Package Description	Marketing Start Date	Marketing End Date
1 NDC:62839-2810-1	250 mL in 1 BOTTLE, PLASTIC		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC MONOGRAPH FINAL	part358H	10/01/2009	

Labeler - NSE Products, Inc. (966817975)

Revised: 6/2011 NSE Products, Inc.