AXE SIGNATURE DAILY DANDRUFF DEFENSE ANTI-DANDRUFF- pyrithione zinc shampoo
Conopco, Inc. d/b/a Unilever

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.

Axe Signature Daily Dandruff Defense Anti-Dandruff Shampoo

Drug Facts

Active ingredient

With Pyrithione Zinc (1.0% WW)

Purpose

anti-dandruff

Uses

- helps control scalp itching and flaking 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 out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

Directions

- Squeeze. Lather for several minutes. Rinse.
- Use at least twice a week or as directed by a doctor.

Inactive Ingredients

Water, Sodium Laureth Sulfate, Cocamidopropyl Betaine, Sodium Chloride, Fragrance (Parfum), Carbomer, Dimethiconol, DMDM Hydantoin, Citric Acid, PPG-9, Guar Hydroxypropyltrimonium Chloride, TEA-Dodecylbenzenesulfonate, Butylene Glycol, Iodopropynyl Butylcarbamate, Methylchloroisothiazolinone, Methylisothiazolinone

Questions?

1-800-450-7580

Consult a doctor prior to use in children under 2 years of age.

Packaging

AXE SIGNATURE DAILY DANDRUFF DEFENSE ANTI-DANDRUFF pyrithione zinc shampoo

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:64942-1483
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 LAURETH-3 SULFATE (UNII: BPV390UAP0)	
COCAMIDOPROPYL BETAINE (UNII: 5OCF3O11KX)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
CARBOMER HOMOPOLYMER TYPE C (ALLYL PENTAERYTHRITOL CROSSLINKED) (UNII: 4Q93RCW27E)	
DIMETHICONOL (40 CST) (UNII: 343C7U75XW)	
DMDM HYDANTOIN (UNII: BYR0546TOW)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
PPG-9 (UNII: I29VQH0G0B)	
GUAR HYDROXYPROPYLTRIMONIUM CHLORIDE (1.7 SUBSTITUENTS PER SACCHARIDE) (UNII: B16G315W7A)	
TRIETHANOLAMINE DODECYLBENZENESULFONATE (UNII: 8HM7ZD48HN)	
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	
IODOPROPYNYL BUTYLCARBAMATE (UNII: 603P14DHEB)	
METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)	
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:64942-1483-1	355 mL in 1 CONTAINER; Type 0: Not a Combination Product	11/10/2016	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part358H	11/10/2016	

Labeler - Conopco, Inc. d/b/a Unilever (001375088)