BLADE HOWL ANTI-DANDRUFF- pyrithione zinc shampoo
CVS PHARMACY 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.

Drug Facts Box

Active Ingredient

Pyrithione Zinc 1%

Purpose
Antidandruff

Uses

To help 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 using this product 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

In case of accidental ingestion, get medical help or contact a Poison Control Center immediately.

Directions

For maximum dandruff control, use every time you shampoo. Wet hair, massage onto scalp and rinse . Repeat if desired.

Inactive Ingredients
Water (Aqua), Sodium Laureth Sulfate, Acrylates Copolymer, Polyquaternium-7, Cocamide MEA, Glycol Distearate, Cocamidopropyl Betaine, Fragrance (Parfum), Dimethicone, Sodium Chloride, Tetrasodium EDTA, Sodium Hydroxide, Citric Acid DMDM Hydantoin, Blue 1 (CI 42090), Red 33 (CI 17200).

Package Front and Back Labels

blade.jpg

BLADE HOWL ANTI-DANDRUFF

pyrithione zinc shampoo

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:59779-422
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 mL in 100 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
SODIUM LAURETH SULFATE (UNII: BPV390UAP0)	
AMMONIO METHACRYLATE COPOLYMER TYPE A (UNII: 8GQS4E66YY)	
POLYQUATERNIUM-7 (70/30 ACRYLAMIDE/DADMAC; 1600 KD) (UNII: 0L414VCS5Y)	
COCO MONOETHANOLAMIDE (UNII: C80684146D)	
GLYCOL DISTEARATE (UNII: 13W7MDN21W)	
COCAMIDOPROPYL BETAINE (UNII: 5OCF3O11KX)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
EDETATE SODIUM (UNII: MP1J8420LU)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
DMDM HYDANTOIN (UNII: BYR0546TOW)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
D&C RED NO. 33 (UNII: 9DBA0SBB0L)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:59779-422-12	355 mL in 1 BOTTLE, PLASTIC		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part358H	11/03/2010	

Labeler - CVS PHARMACY INC (062312574)

Registrant - APOLLO HEALTH AND BEAUTY CARE (201901209)

Establishment

Name	Address	ID/FEI	Business Operations
APOLLO HEALTH AND BEAUTY CARE		201901209	manufacture

Revised: 11/2010 CVS PHARMACY INC