FRESH MINT DANDRUFF - pyrithione zinc shampoo
Vi Jon

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.

Dandruff Shampoo 309

Active ingredient

Pyrithione zinc 1%

purpose

Anti-dandruff

use

helps prevent recurrence of flaking and itching associated with dandruff

Warnings

For external use only

When using this product

• do not get into 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 as directed.

Keep out of reach of children.

If swallowed, get medical help or contact a Poison Control Center right away.

Directions

• shake well • for maximum dandruff control, use every time you shampoo • wet hair, message 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 lauryl sulfate, sodium laureth sulfate, glycol distearate, sodium chloride, zinc carbonate, sodium xylenesulfonate, amodimethicone, cocamidopropyl betaine, fragrance, sodium benzoate, guar hydroxypropyltrimonium chloride, magnesium carbonate hydroxide, citric acid, methylchloroisothiazolinone, methylisothiazolinone, blue 1, red 33

principal display panel

Dandruff Shampoo
14.2 FL OZ (420 mL)

Image not available.
Manufactured exclusively for private label distribution

FRESH MINT DANDRUFF

pyrithione zinc shampoo

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:11344-309
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 kg in 100 kg

Inactive Ingredients	
Ingredient Name	**Strength**
SODIUM LAURETH SULFATE (UNII: BPV390UAP0)	
SODIUM LAURYL SULFATE (UNII: 368GB5141J)	
GLYCOL DISTEARATE (UNII: 13W7MDN21W)	
ZINC CARBONATE (UNII: EQR32Y7H0M)	
SODIUM XYLENESULFONATE (UNII: G4LZF950UR)	
AMODIMETHICONE (1300 CST) (UNII: 3V7U636DWN)	
COCAMIDOPROPYL BETAINE (UNII: 5OCF3O11KX)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
GUAR HYDROXYPROPYLTRIMONIUM CHLORIDE (1.7 SUBSTITUENTS PER SACCHARIDE) (UNII: B16G315W7A)	
MAGNESIUM CARBONATE (UNII: 0E53J927NA)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)	
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	

FD&C RED NO. 3 (UNII: PN2ZH5LOQY)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11344-309-15	.402 kg in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	08/21/2004	
2	NDC:11344-309-21	.088 kg in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	08/21/2004	
3	NDC:11344-309-35	.700 kg in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	08/21/2004	
4	NDC:11344-309-39	1.0 kg in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	08/21/2004	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part358H	08/21/2004	

Labeler - Vi Jon (150931459)

Registrant - Vi Jon (088520668)

Establishment

Name	Address	ID/FEI	Business Operations
Vi Jon		088520668	manufacture(11344-309)

Revised: 1/2020 Vi Jon