

EQUATE EVERYDAY CLEAN- pyritione zinc liquid
Wal-Mart Stores Inc

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Drug Facts

Active ingredient

Pyrithione Zinc 1%

Purpose

Anti-dandruff

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 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.

- 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, Sodium Chloride, Acrylates Copolymer, Glycol Distearate, Cocamidopropyl Betaine, Cocamide MEA, Laureth-4, Fragrance (Parfum), Sodium Hydroxide, Tetrasodium EDTA, Methylchloroisothiazolinone, Methylisothiazolinone, Blue1 (CI 42090), Red 33 (CI 17200).

Label Copy

equate[®]

Compare to Head & Shoulders[®] Active Ingredient*

Everyday Clean

Pyrithione Zinc Dandruff Shampoo

For Normal Hair

Helps relieve itching and flaking associated with dandruff

33.8 FL OZ (1 L)

06-19166

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Questions or comments?
1-888-287-1915

Satisfaction guaranteed - for questions or comments please call 1-888-287-1915.

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EQUATE EVERYDAY CLEAN

pyrithione zinc liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:49035-427
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 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
SODIUM LAURETH SULFATE (UNII: BPV390UAP0)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
METHACRYLIC ACID - METHYL METHACRYLATE COPOLYMER (1:1) (UNII: 74G4R6TH13)	
GLYCOL DISTEARATE (UNII: 13W7MDN21W)	
COCAMIDO PROPYL BETAINE (UNII: 5OCF3O11KX)	
COCO MONOETHANOLAMIDE (UNII: C80684146D)	
LAURETH-4 (UNII: 6HQ855798J)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
EDETATE SODIUM (UNII: MP1J8420LU)	
METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)	
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	
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:49035-427-33	1000 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	04/13/2017	
2	NDC:49035-427-13	420 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	04/13/2017	
3	NDC:49035-427-24	701 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	04/13/2017	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part358H	04/13/2017	

Labeler - Wal-Mart Stores Inc (051957769)

Registrant - Apollo Health and Beauty Care Inc. (201901209)

Establishment

Name	Address	ID/FEI	Business Operations
Apollo Health and Beauty Care Inc.		201901209	manufacture(49035-427)