WALMART- pyrithione zinc lotion/shampoo Wal-Mart Stores, Inc

NDC- 79903-899, Equate Smooth & Silky

Pyrithione Zinc

For the relief of itching, redness and flaking associated with dandruff

For External use

avoid contact with eyes. If contact occurs, rinse thoroughly with water.

Stop use and ask a doctor ifcondition worsens or does not improve after regular use of this product.

Keep out of reach of children. If swallowed, call a poison control centre or get medical help right away.

- Adults and children 2 years and older.
- Shake well.
- Apply evenly to scalp. leave on for several minutes and then rinse off.
- Use at least twice per week or as directed by a doctor or pharmacist.

Water, Stearyl Alcohol, Behentrimonium Chloride, Cetyl Alcohol, Cocos Nucifera (Coconut) Oil,

Dimethicone, Fragrance, PPG-3 Benzyl Ether Ethylhexanoate, Lauryl PEG/PPG-18/18 Methicone,

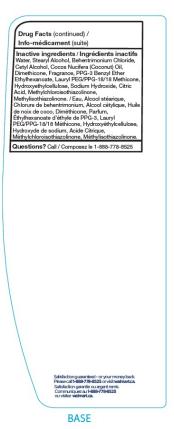
Hydroxyethylcellulose, Sodium Hydroxide, Citric Acid, Methylchloroisothiazolinone, Methylisothiazolinone.

Store at room temperature

Relieves itching, redness and flakes associated with Dandruff.







WALMART

pyrithione zinc lotion/shampoo

Product Type HUMAN OTC DRUG Item Code (Source) NDC:79903-899

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name

Basis of Strength

PYRITHIONE ZINC (UNII: R953O2RHZ5) (PYRITHIONE ZINC - UNII:R953O2RHZ5)

PYRITHIONE ZINC 300 mg in 100 mL

Inactive Ingredients	
Ingredient Name	Strength
BEHENTRIMONIUM CHLORIDE (UNII: X7GNG3S47T)	
FRAGRANCE CLEAN ORC0600327 (UNII: 329LCV5BTF)	
METHYLCHLOROISOTHIAZOLINONE/METHYLISOTHIAZOLINONE MIXTURE (UNII: 1509QS218W)	
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
LAURYL PEG/PPG-18/18 METHICONE (UNII: ZJ5S27D9NX)	
CETYL HYDROXYETHYLCELLULOSE (350000 MW) (UNII: T7SWE4S2TT)	
STEARYL ALCOHOL (UNII: 2KR89I4H1Y)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
PPG-3 BENZYL ETHER ETHYLHEXANOATE (UNII: 3N703GY99W)	

WATER (UNII: 059QF0KO0R)	
CETYL ALCOHOL (UNII: 936JST6JCN)	
COCONUT OIL (UNII: Q9L0O73W7L)	
DIMETHICONE (UNII: 92RU3N3Y1O)	

I	Packaging					
	#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
	1	NDC:79903- 899-70	700 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	11/04/2022		

Marketing Information					
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date		
OTC Monograph Drug	M032	11/04/2022			

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(79903-899)			

Revised: 1/2025 Wal-Mart Stores, Inc