

DOVE- pyrithione zinc liquid
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.

Dove Dermacare Scalp Aloe & Restore Anti-Dandruff Conditioner

**DOVE DERMACARE SCALP ALOE & RESTORE ANTI-DANDRUFF CONDITIONER -
pyrithione zinc liquid**

Dove Dermacare Scalp Aloe & Restore Anti-Dandruff Conditioner

Drug Facts

Active ingredient

Pyrithione Zinc (0.5%)

Purpose

Anti-dandruff

Use

- Helps prevent and control recurrence of itching, flaking and irritation associated with dandruff.

Warning

- 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 this and all drugs out of reach of children.**
If swallowed, get medical help or contact a Poison Control Center right away.

Directions

For best results use at least twice per week or as directed by a doctor. Massage into scalp and hair. Rinse.

Inactive Ingredients

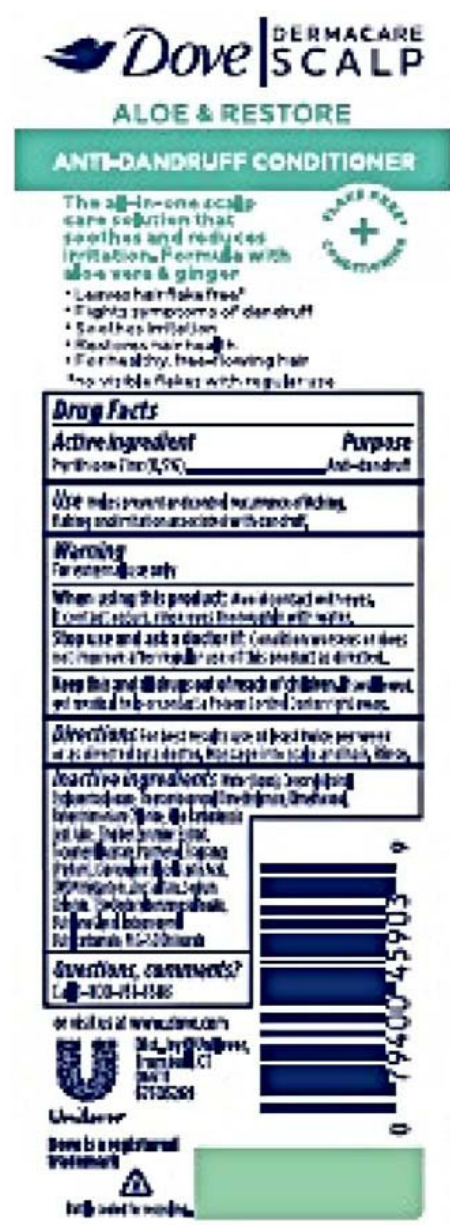
Water(Aqua), Cetearyl Alcohol, Cyclopentasiloxne, Stearamidopropyl Dimethylamine,

Dimethiconol, Behentrimonium Chloride, Aloe Barbadensis Leaf Juice, Zingiber Zerumbet Extract, Tocopheryl Acetate, Panthenol, Fragrance(Parfum), Dipropylene Glycol, Lactic Acid, DMDM Hydantoin, Zinc Sulfate, Sodium Chloride, TEA-Dodecylbenzenesulfonate, Butylene Glycol, Iodopropynyl Butylcarbamate, PEG-150 Distearate.

Questions?

Call 1-800-761-3683

Packaging



DOVE
pyrithione zinc liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:64942-1620
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
PYRITHIONE ZINC (UNII: R953O2RHZ5) (PYRITHIONE ZINC - UNII:R953O2RHZ5)	PYRITHIONE ZINC	0.5 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
WATER (UNII: 059QF0KO0R)	
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)	
CYCLOMETHICONE 5 (UNII: 0THT5PCI0R)	
STEARAMIDOPROPYL DIMETHYLAMINE (UNII: K7VEI00UFR)	
DIMETHICONOL (40 CST) (UNII: 343C7U75XW)	
BEHENTRIMONIUM CHLORIDE (UNII: X7GNG3S47T)	
.ALPHA.-TOCOPHEROL ACETATE, D- (UNII: A7E6112E4N)	
DIPROPYLENE GLYCOL (UNII: E107L85C40)	
LACTIC ACID, UNSPECIFIED FORM (UNII: 33X04XA5AT)	
DMDM HYDANTOIN (UNII: BYR0546TOW)	
ZINC SULFATE, UNSPECIFIED FORM (UNII: 89DS0H96TB)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
TRIETHANOLAMINE DODECYLBENZENESULFONATE (UNII: 8HM7ZD48HN)	
BUTYLENE GLYCOL (UNII: 3XUS85KORA)	
IODOPROPYNYL BUTYLCARBAMATE (UNII: 603P14DHEB)	
PEG-150 DISTEARATE (UNII: 6F36Q0I0AC)	
PANTHENOL (UNII: WW9CM0O67Z)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:64942-1620-1	355 mL in 1 CONTAINER; Type 0: Not a Combination Product	01/01/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part358H	01/01/2020	

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

