

**DOVE DERMACARE SCALP DRYNESS AND ITCH RELIEF ANTI-DANDRUFF-
pyrithione zinc shampoo
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 Dryness and Itch Relief Anti-Dandruff Shampoo

Drug Facts

Active Ingredient

Pyrithione Zinc (1.0%)

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), Sodium Laureth Sulfate, Cocamidopropyl Betaine, Sodium Chloride, Cocos Nucifera (Coconut) Oil, Butyrospermum Parkii (Shea) Butter, Carbomer, Fragrance (Parfum), Dimethiconol, DMDM Hydantoin, Citric Acid, Glycerin, PPG-9, Zinc Sulfate, Guar

Hydroxypropyltrimonium Chloride, TEA-Dodecylbenzenesulfonate, Butylene Glycol, Iodopropynyl Butylcarbamate, Methylchloroisoithiazolinone, Methylisothiazolinone

Questions, comments?

Call 1-800-761-3683

Gently cleanses & soothes itchy, dry scalp

For smooth, less frizzy hair

FLAKE FREE* + pH BALANCED

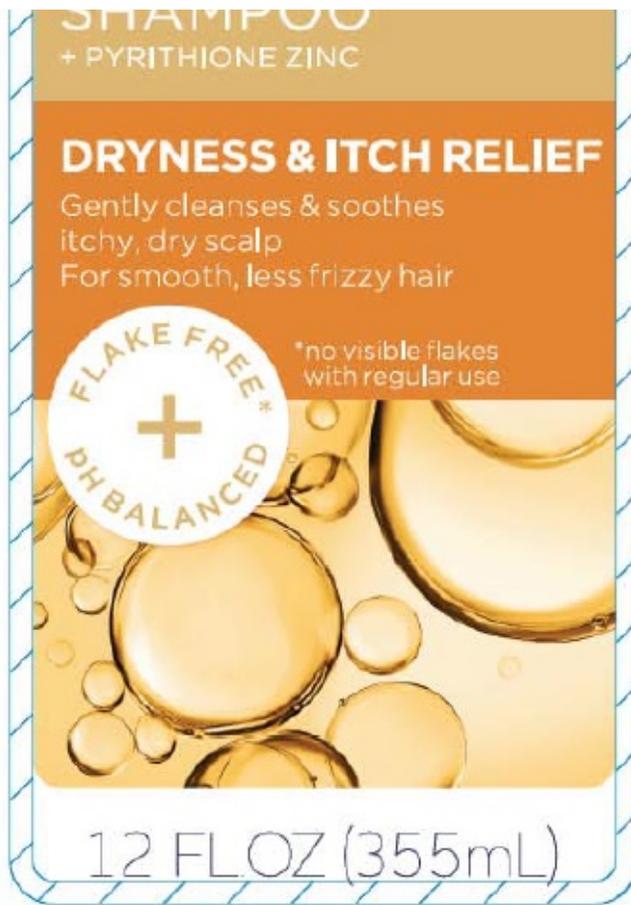
The solution that nourishes the scalp and relieves it from irritation

- Leaves hair flake free*
- Hydrates dry scalp
- Soothes itchy scalp
- Coconut & Shea butter indulgent scent
- For smooth, less frizzy hair

*No visible flakes with regular use

Packaging





DOVE DERMACARE SCALP DRYNESS AND ITCH RELIEF ANTI-DANDRUFF

pyrithione zinc shampoo

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:64942-1487
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 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
SODIUM LAURETH-3 SULFATE (UNII: BPV390UAP0)	
COCAMIDOPROPYL BETAINE (UNII: 5OCF3011KX)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
COCONUT OIL (UNII: Q9L0073W7L)	
SHEA BUTTER (UNII: K49155WL9Y)	
CARBOMER HOMOPOLYMER TYPE C (ALLYL PENTAERYTHRITOL CROSSLINKED) (UNII:)	

4Q93RCW27E)	
DIMETHICONOL (40 CST) (UNII: 343C7U75XW)	
DMDM HYDANTOIN (UNII: BYR0546TOW)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
GLYCERIN (UNII: PDC6A3C0OX)	
PPG-9 (UNII: I29VQH0G0B)	
ZINC SULFATE, UNSPECIFIED FORM (UNII: 89DS0H96TB)	
GUAR HYDROXYPROPYLTRIMONIUM CHLORIDE (1.7 SUBSTITUENTS PER SACCHARIDE) (UNII: B16G315W7A)	
TRIETHANOLAMINE DODECYLBENZENESULFONATE (UNII: 8HM7ZD48HN)	
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	
IODOPROPYNYL BUTYLCARBAMATE (UNII: 603P14DHEB)	
METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)	
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:64942-1487-1	355 mL in 1 CONTAINER; Type 0: Not a Combination Product	11/10/2016	
2	NDC:64942-1487-2	53 mL in 1 CONTAINER; Type 0: Not a Combination Product	10/15/2018	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part358H	11/10/2016	

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

Revised: 3/2021

Conopco, Inc. d/b/a/ Unilever