EVERYDAY CLEAN DANDRUFF- pyrithione zinc shampoo
Value Merchandisers

Everyday Clean Dandruff Shampoo
153.008/153AZ rev1, BA-BD

Claims

Best Choice®

EVERYDAY CLEAN

Dandruff Shampoo

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 occures, 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, massage onto scalp, rinse, repeat if desired

- for best results use at least twice a week or as directed by a doctor

Inactive ingIngredient

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

Questions?

Call 1-888-593-0593

Disclaimer

This product is not manufactured or distributed by Procter & Gamble, distributor of Head & Shoulders® Classic Clean Dandruff Shampoo.

Adverse Reactions

PRODULY DISTRIBUTED BY:

ASSOCIATED WHOLESALE GROCERS, INC.

KANSAS CITY, KS 66106

Scan here for more product information.

Call 1-844-282-1112 for more product information.

Best Choice®

100% Guaranteed

www.bestchoicebrand.com

Principal display panel

Best Choice®

COMPARE TO THE ACTIVE INGREDIENT IN HEAD & SHOULDERS®*

EVERYDAY CLEAN

Dandruff Shampoo

Pyrithione Zinc

Dandruff Shampoo

12.5 FL OZ (370 mL)

EVERYDAY CLEAN DANDRUFF

pyrithione zinc shampoo

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:63941-153
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 LAURYL SULFATE (UNII: 368GB5141J)	
SODIUM LAURETH SULFATE (UNII: BPV390UAP0)	
GLYCOL DISTEARATE (UNII: 13W7MDN21W)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
ZINC CARBONATE (UNII: EQR32Y7H0M)	
SODIUM XYLENESULFONATE (UNII: G4LZF950UR)	
AMODIMETHICONE (800 CST) (UNII: 363Z2T48P7)	
COCAMIDOPROPYL BETAINE (UNII: 5OCF3O11KX)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
GUAR HYDROXYPROPYLTRIMONIUM CHLORIDE (1.7 SUBSTITUENTS PER SACCHARIDE) (UNII: B16G315W7A)	
MAGNESIUM CARBONATE HYDROXIDE (UNII: YQO029V1L4)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
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:63941-153-15	420 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	03/14/1999	
2	NDC:63941-153-32	370 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	11/15/2023	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M032	03/14/1999	

Labeler - Value Merchandisers (868703513)

Registrant - Vi-Jon, LLC (790752542)

Establishment

Name	Address	ID/FEI	Business Operations
Vi-Jon, LLC		088520668	manufacture(63941-153)

Establishment

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Vi-Jon, LLC		790752542	manufacture(63941-153)

Revised: 11/2023 Value Merchandisers