DRY SCALP CARE- pyrithione zinc shampoo HEB

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.

HEB 296.007 296AT-AV

Claims

DANDRUFF 2 IN 1

DRY SCALP RESCUE

Dandruff Care for Great Looing Hair.

Helps relieve scalp dryness, itch, irritation and helps prevent flakes.

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 occurs, 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 ingredients

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, Prunus amygdalus dulcis (sweet almond) oil, methylchloroisothiazolinone, methylisothiazolinone

Questions

Call 1-888-593-0593

Adverse Reactions

MADE WITH PRIDE AND CARE FOR H-E-B, SAN ANTONIO, TX 78204 100% GUARANTEE promise

If you aren't completely pleased with thiw produc,

we'll be happy to replace it or refund your mone.

You have our word on it

principal display panel

H-E-B

Dandruff

2 IN 1

SHAMPOO + CONDITIONER

With Pyrithione Zinc

Dry Scalp Rescue

WITH MOISTURIZERS

RELIEVES DRY SCALP AND ITCH

Leaves Hair

Healthy Looking &

Fights Dry Scalp



DRY SCALP CARE

pyrithione zinc shampoo

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:37808-296

Route of Administration TOPICAL

Active Ingredient/Active Moiety Ingredient Name Basis of Strength PYRITHIONE ZINC (UNII: R95302RHZ5) (PYRITHIONE ZINC - UNII:R95302RHZ5) PYRITHIONE ZINC (UNII: R95302RHZ5) 14 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: 50CF3011KX)					
SODIUM BENZOATE (UNII: OJ245FE5EU)					
GUAR HYDROXYPROPYLTRIMONIUM CHLORIDE (1.7 SUBSTITUENTS PER SACCHARIDE) (UNII: B16G315W7A)					
MAGNESIUM CARBONATE HYDROXIDE (UNII: YQO029V1L4)					
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)					
ALMOND OIL (UNII: 66YXD4DKO9)					
METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)					
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)					

F	Packaging						
#	Item Code	Package Description	Marketing Start Date	Marketing End Date			
1	NDC:37808- 296-35	700 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	05/15/2009				
2	NDC:37808- 296-15	420 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	05/15/2009				

Marketing Information						
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date			
OTC monograph final	part358H	05/15/2009				

Labeler - HEB (007924756)

Registrant - Vi-Jon, LLC (790752542)

EstablishmentNameAddressID/FEIBusiness OperationsVi-Jon, LLC790752542manufacture(37808-296)

Establishment						
Name	Address	ID/FEI	Business Operations			
Vi-Jon, LLC		088520668	manufacture(37808-296)			

Revised: 4/2022 HEB