VANICREAM- pyrithione zinc shampoo
Pharmaceutical Specialties, Inc.

VANICREAM™ Dandruff Shampoo

Drug Facts

Active Ingredient

Pyrithione zinc 2%

Purposes

Anti-dandruff,

Anti-seborrheic dermatitis

Uses

Controls and reduces the symptoms of dandruff and seborrheic dermatitis

Warnings

For external use only

Ask a doctor before use if you have a condition that covers a large area of the body.

When using this product avoid contact with the 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 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 a week or as directed by a doctor
- Shake before use
- Wet hair and massage gently into scalp
- Leave lather on scalp for several minutes
- Rinse well, repeat if needed

Other information

Store at room temperature

Inactive ingredients purified water, lauryl glucoside, coco-glucoside, acrylates copolymer, panthenol, pentylene glycol, glycerin, sucrose cocoate, disodium cocoyl glutamate, sodium cocoyl glycinate, behenyl alcohol, sucrose stearate, cetyl palmitate, sodium cocoyl glutamate, 1,2-hexanediol, sodium hydroxide, caprylyl glycol, sodium chloride

Questions?

1-800-325-8232

www.vanicream.com

Dist. by

PHARMACEUTICAL SPECIALTIES, INC.

ROCHESTER, MN 55901 • Made in U.S.A.

www.vanicream.com • 1-800-325-8232

NDC 45334-206-08

Formerly known as Free & Clear™

VANICREAM™

Pyrithione Zinc 2%

Dandruff Shampoo

for Sensitive Skin

DERMATOLOGIST TESTED

Free of dyes, fragrance, masking fragrance, lanolin, protein, parabens, & formaldehyde releasers

Sulfate-free • Betaine-free • Gluten-free

8 fl oz (237 mL)

VANICREAM

pyrithione zinc shampoo

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:45334-206
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
PYRITHIONE ZINC (UNII: R953O2RHZ5) (PYRITHIONE ZINC - UNII:R953O2RHZ5)	PYRITHIONE ZINC	2 g in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**

1,2-HEXANEDIOL (UNII: TR046Y3K1G)	
METHACRYLIC ACID - ETHYL ACRYLATE COPOLYMER (4500 MPA.S) (UNII: T967IEU43C)	
SUCROSE COCOATE (UNII: 3H18P0UK73)	
SUCROSE STEARATE (UNII: 274KW0O50M)	
CAPRYLYL GLYCOL (UNII: 00YIU5438U)	
COCO GLUCOSIDE (UNII: ICS790225B)	
SODIUM COCOYL GLYCINATE (UNII: XLU9KH03XM)	
WATER (UNII: 059QF0KO0R)	
DOCOSANOL (UNII: 9G1OE216XY)	
PENTYLENE GLYCOL (UNII: 50C1307PZG)	
GLYCERIN (UNII: PDC6A3C0OX)	
DEXPANTHENOL (UNII: 1O6C93RI7Z)	
LAURYL GLUCOSIDE (UNII: 76LN7P7UCU)	
DISODIUM COCOYL GLUTAMATE (UNII: MBK0CP8F5A)	
CETYL PALMITATE (UNII: 5ZA2S6B08X)	
SODIUM COCOYL GLUTAMATE (UNII: BMT4RCZ3HG)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:45334-206-08	237 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	01/03/2022	
2	NDC:45334-206-07	7.4 mL in 1 TUBE; Type 0: Not a Combination Product	01/03/2022	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M032	01/03/2022	

Labeler - Pharmaceutical Specialties, Inc. (076499557)

Establishment

Name	Address	ID/FEI	Business Operations
Pharmaceutical Specialties, Inc.		076499557	manufacture(45334-206)

Revised: 10/2023 Pharmaceutical Specialties, Inc.