

VANICREAM Z-BAR- pyrithione zinc soap
Pharmaceutical Specialties, Inc.

Vanicream Z-Bar

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
- Use on affected areas in place of your regular soap
- Work up a rich lather using warm water and massage gently into affected areas
- Rinse well

Other information

Store at room temperature

Inactive ingredients sodium cocoyl isethionate, stearic acid, coconut acid, sodium isethionate, water, petrolatum, sorbitol, cetearyl alcohol, propanediol, cetareth-20, simethicone, glyceryl stearate, PEG-30 stearate, sorbic acid

Questions? 1-800-325-8232 www.vanicream.com

Dist. by PHARMACEUTICAL SPECIALTIES, INC. ROCHESTER, MN 55901

1-800-325-8232 www.vanicream.com

Made in the USA with US and select imported materials

VANICREAM™ Z-Bar

Pyrrhithione Zinc 2%

NDC 45334-392-35

DERMATOLOGIST TESTED

Seborrheic Dermatitis & Anti-dandruff Medicated Cleansing Bar

for Sensitive Skin

Net Wt 3.5 oz (99g)



VANICREAM Z-BAR

pyrithione zinc soap

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:45334-392
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 g

Inactive Ingredients

Ingredient Name	Strength
SODIUM COCOYL ISETHIONATE (UNII: 518XTE8493)	
PROPANEDIOL (UNII: 5965N8W85T)	
POLYOXYL 20 CETOSTEARYL ETHER (UNII: YRC528SWUY)	
PEG-30 STEARATE (UNII: 1U8KB35S20)	
WATER (UNII: 059QF0KO0R)	
COCONUT ACID (UNII: 40U37V505D)	
DIMETHICONE, UNSPECIFIED (UNII: 92RU3N3Y10)	
SORBIC ACID (UNII: X045WJ989B)	
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)	
GLYCERYL MONOSTEARATE (UNII: 230OU9XXE4)	
SORBITOL (UNII: 506T60A25R)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
PETROLATUM (UNII: 4T6H12BN9U)	
SODIUM ISETHIONATE (UNII: 3R36J71C17)	

Product Characteristics

Color	white	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:45334-392-35	99 g in 1 CARTON; Type 0: Not a Combination Product	10/01/2024	

Marketing Information

Marketing	Application Number or Monograph	Marketing Start	Marketing End
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Category	Citation	Date	Date
OTC Monograph Drug	M032	10/01/2024	

Labeler - Pharmaceutical Specialties, Inc. (076499557)

Revised: 10/2024

Pharmaceutical Specialties, Inc.