

SATINIQUE ANTI DANDRUFF- pyrrithione zinc cream

Access Business Group LLC

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.

Satinique Anti-Dandruff shampoo

Drug Facts

Active Ingredients

Pyrrithione Zinc 0.95%

Purpose

Anti Dandruff

Uses

- Controls symptoms of dandruff

Warnings

For external use only.

Avoid contact with eyes. If contact occurs, rinse eyes thoroughly with water.

If condition worsens or does not improve after regular use, consult a doctor.

Keep out of reach of children.

If swallowed, get medical help or contact a Poison Control Center right away.

Directions

- Shake well before use.
- Wet hair. Apply shampoo and work into a lather. Rinse thoroughly.
- For best result, use at least twice a week or as directed by a doctor.

Inactive Ingredients

Water, Sodium Laureth Sulfate, Decyl Glucoside, Glycol Distearate, Disodium Laureth Sulfosuccinate, Glycereth-26, Ceramide 2, Ceramide-3, C10-40 isoalkylamidopropylethyldimonium ethosulfate, Creatine, Fragrance, Dimethicone, Sodium Chloride, C12-15 Alkyl Lactate, Hydroxypropyl Methylcellulose, Guar Hydroxypropyltrimonium Chloride, Zinc Chloride, Hectorite, Hydroxyethylcellulose, Tetrasodium EDTA, Citric Acid, Tocopheryl Acetate, Limnanthes Alba (Meadowfoam) Seed Oil, Propylene Glycol, Aloe Barbadosensis Leaf Juice, Butyl Avodate, PPG-12-Buteth-15, Triethanolamine, Sodium PCA, Sodium Lactate, Arginine, Aspartic Acid, PCA, Glycine, Alanine, Methylchlororisorothiazolinone, Serine, Valine, Behenic Acid, Cholesterol, Isoleucine, Proline, Threonine, Methylisothiazolinone, Histidine, Phenylalanine, Perilla Ocymoides Leaf Extract, Green 3, Blue 1.

Package Labeling:

FRONT PANEL

SATINIQUE



.0992 between logo and icon

3.9522 from base of bottle to bottom of icon

ANTI-DANDRUFF SHAMPOO for dry, irritated scalp

280 mL (9.4 fl.oz.)

.75" from base of bottle

3.9522 from base of bottle to bottom of icon

NO PRINT

BACK PANEL

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Dist. by Amway Corp., 7575 Fulton, Ada, MI 49355
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Amway™

1033393



11-0670



SATINIQUE ANTI DANDRUFF

pyrithione zinc cream

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:10056-006
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
PYRITHIONE ZINC (UNII: R953O2RHZ5) (PYRITHIONE ZINC - UNII:R953O2RHZ5)	PYRITHIONE ZINC	95 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
SODIUM LAURETH-3 SULFATE (UNII: BPV390UAP0)	
DECYL GLUCOSIDE (UNII: Z17H97EA6Y)	
GLYCOL DISTEARATE (UNII: 13W7MDN21W)	
DISODIUM LAURETH SULFO SUCCINATE (UNII: D6DH1DTN7E)	
GLYCERETH-26 (UNII: NNE56F2N14)	
CERAMIDE 2 (UNII: C04977SRJ5)	
CERAMIDE 3 (UNII: 4370DF050B)	
CREATINE (UNII: MU72812GK0)	
DIMETHICONE (UNII: 92RU3N3Y10)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
C12-15 ALKYL LACTATE (UNII: GC844VRD7E)	
HYPROMELLOSES (UNII: 3NXW29V3WO)	
GUAR HYDROXYPROPYLTRIMONIUM CHLORIDE (1.7 SUBSTITUENTS PER SACCHARIDE) (UNII: B16G315W7A)	
ZINC CHLORIDE (UNII: 86Q357L16B)	
HECTORITE (UNII: 08X4KI73EZ)	
EDETATE SODIUM (UNII: MP1J8420LU)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
MEADOWFOAM SEED OIL (UNII: 412ZHA4T4Y)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
PPG-12-BUTETH-16 (UNII: 58CG7042J1)	
TROLAMINE (UNII: 9O3K93S3TK)	
SODIUM PYRROLIDONE CARBOXYLATE (UNII: 469OTG57A2)	
SODIUM LACTATE (UNII: TU7HW0W0QT)	
ARGININE (UNII: 94ZLA3W45F)	
ASPARTIC ACID (UNII: 30KYC7MIA)	
PIDOLIC ACID (UNII: SZB8301W42)	
GLYCINE (UNII: TE7660XO1C)	

ALANINE (UNII: OF5P57N2ZX)
METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)
SERINE (UNII: 452VLY9402)
VALINE (UNII: HG18B9YRS7)
BEHENIC ACID (UNII: H390488X0A)
CHOLESTEROL (UNII: 97C5T2UQ7J)
ISOLEUCINE (UNII: 04Y7590D77)
PROLINE (UNII: 9DLQ4CIU6V)
THREONINE (UNII: 2ZD004190S)
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)
HISTIDINE (UNII: 4QD397987E)
PHENYLALANINE (UNII: 47E5O17Y3R)
PERILLA FRUTESCENS LEAF (UNII: T4L5881Y68)
FD&C GREEN NO. 3 (UNII: 3P3ONR6O1S)
FD&C BLUE NO. 1 (UNII: HBR47K3TBD)

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:10056-006-01	280 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/29/2017	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part358H	03/29/2017	

Labeler - Access Business Group LLC (839830713)

Revised: 1/2019

Access Business Group LLC