

ZINCON MEDICATED DANDRUFF- pyrithione zinc shampoo
Neoteric Cosmetics, Inc.

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.

Zincon Medicated Dandruff Shampoo 4oz

Drug Facts

Active Ingredient

Pyrithione zinc 1%

(To the right)

Purpose

Anti-dandruff

Uses

- Reduces and helps eliminate scalp itching, flaking and scaling associated with 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 eyes. If contact occurs, rinse thoroughly with water.

Stop use and ask a doctor if:

- Condition worsens
- Condition 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:

- Shake well
- Wet hair, apply shampoo and work into lather
- Rinse thoroughly
- For best results, use at least twice a week or as directed by a doctor
- Do not use on children under 2 years of age, except as directed by a doctor

Inactive Ingredients

Citric acid, cocamide MEA, deionized water, FD & C Blue No. 1, Fragrance, glycol distearate, magnesium aluminum silicate, methylchloroisothiazolinone, methylisothiazolinone, propylene glycol, sodium chloride, sodium laureth sulfate, sodium lauryl sulfate.

Questions? 1-800-220-0151

Manufactured for Neoteric Cosmetics, Inc.

Denver, CO 80239

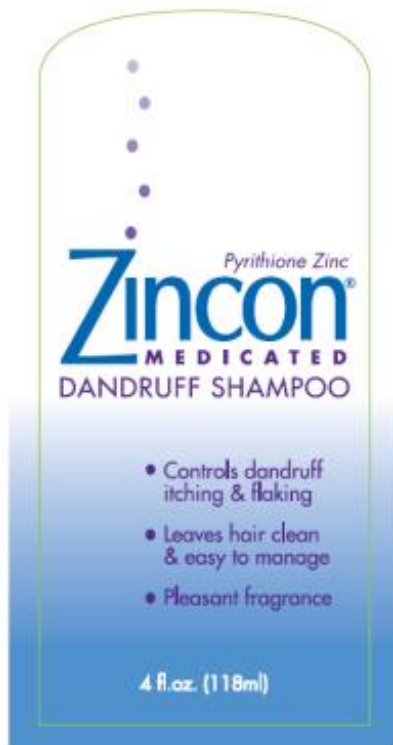
WWW.Neotericcosmetics.com MADE IN USA

PRINCIPAL DISPLAY PANEL

Pyrithione Zinc
ZINCON
MEDICATED
DANDRUFF SHAMPOO

- Controls dandruff itching & flaking
- Leaves hair clean & easy to manage
- Pleasant fragrance

4 fl.oz. (118ml)



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Pyrithione Zinc
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- Controls dandruff itching & flaking

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8 fl.oz. (236ml)



ZINCON MEDICATED DANDRUFF

pyrithione zinc shampoo

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:626 73-0 16
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
PYRITHIONE ZINC (UNII: R953O2RHZ5) (PYRITHIONE ZINC - UNII:R953O2RHZ5)	PYRITHIONE ZINC	0.01 g in 1 mL

Inactive Ingredients

Ingredient Name	Strength
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	

COCO MONOETHANOLAMIDE (UNII: C80684146D)
WATER (UNII: 059QF0KO0R)
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)
GLYCOL DISTEARATE (UNII: 13W7MDN21W)
MAGNESIUM ALUMINUM SILICATE (UNII: 6M3P64V0NC)
METHYLCHLORO ISOTHIAZOLINONE (UNII: DEL7T5QRPN)
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)
SODIUM CHLORIDE (UNII: 451W47IQ8X)
SODIUM LAURETH SULFATE (UNII: BPV390UAP0)
SODIUM LAURYL SULFATE (UNII: 368GB5141J)

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:62673-016-94	118 mL in 1 BOTTLE; Type 0: Not a Combination Product	12/13/2017	
2	NDC:62673-016-98	236 mL in 1 BOTTLE; Type 0: Not a Combination Product	12/13/2017	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC MONOGRAPH FINAL	part358H	12/13/2017	

Labeler - Neoteric Cosmetics, Inc. (790615181)

Revised: 12/2017

Neoteric Cosmetics, Inc.