

ZINC THERAPY- pyrithione zinc liquid
D3 Development, 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.

Zinc Therapy Soap

Drug Facts

Active ingredient

Pyrithione zinc 2%

Purposes

Dandruff, Seborrheic dermatitis

Uses

Controls, reduces, and helps stop the symptoms of dandruff and seborrheic dermatitis.

Warnings

For external use only

Ask a doctor before use if you have seborrheic dermatitis that covers a large portion of the body.

When using this product avoid contact with 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 before each use
- Use on affected areas in place of your regular soap
- For best results use at least twice a week or as directed by a doctor
- Work up a lather using warm water and massage into affected areas
- Rinse well

Other information

- Store at room temperature
- Lot number and expiration date can be found on the bottom of this container

Inactive ingredients

Water, potassium cocoate, potassium oleate, olive oil, vegetable glycerin, coco glucoside, salt, oatmeal, citric acid

Questions?

1-800-827-3730

www.dermaharmony.com

Distributed by:

D3 Development, Inc., Portland, ME 04101

Made in the USA

dermaharmony

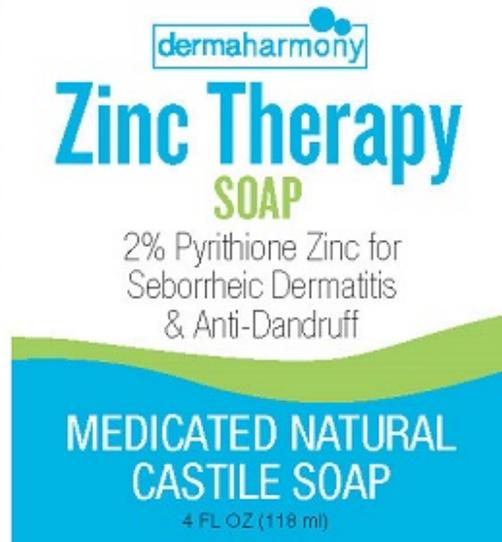
Zinc Therapy SOAP

2% Pyrrhione Zinc for Seborrheic Dermatitis & Anti-Dandruff
 MEDICATED NATURAL CASTILE SOAP

4 FL OZ (118 ml)

Drug Facts (continued)	
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ZINC THERAPY

pyrrhione zinc liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:718 19-007
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
PYRRHIONE ZINC (UNII: R953O2RHZ5) (PYRRHIONE ZINC - UNII:R953O2RHZ5)	PYRRHIONE ZINC	2 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
WATER (UNII: 059QF0KO0R)	
POTASSIUM COCOATE (UNII: F8U72V8ZXP)	
GLYCERIN (UNII: PDC6A3C0OX)	
OATMEAL (UNII: 8PI54V663Y)	
POTASSIUM OLEATE (UNII: 74WHF607EU)	
OLIVE OIL (UNII: 6UYK2W1W1E)	
COCO GLUCOSIDE (UNII: ICS790225B)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:71819-007-04	118 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	01/23/2018	
2	NDC:71819-007-12	355 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	01/23/2018	
3	NDC:71819-007-64	1893 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	01/23/2018	
4	NDC:71819-007-99	3785 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	01/23/2018	
5	NDC:71819-007-10	296 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/02/2019	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part358H	01/23/2018	

Labeler - D3 Development, Inc. (043195877)

Revised: 7/2019

D3 Development, Inc.