

PYRITHIONE ZINC - pyrithione zinc liquid

CVS Pharmacy

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.

drug facts

| Active ingredient | Purpose |
|---------------------------|--|
| Pyrithione Zinc (1%)..... | Anti-dandruff Anti-seborrheic dermatitis |

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Use Controls the symptoms of dandruff and seborrheic dermatitis

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.
- Wet hair thoroughly. - Massage a liberal amount into your scalp.
- Leave lather on scalp for several minutes - Rinse and repeat.

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 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.

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Inactive Ingredients Water, Ammonium Lauryl Sulfate, Sodium Laureth Sulfate, Cocoamphodlacetate, Glycol Stearate, Divinydimethicone.Dimethicone Copolymer, C12-13 Pareth-3, C12-13 Pareth-23, Hydrolyzed Wheat Protein PG-Propyl Silanetriol, BHT, Tocopharyl Acetate, Amodimethicone, Castor Isostearate Succinate, Guar Hydroxypropyltrimonium Chloride, Sodium Methyl Cocoyl Tauratem, Methylparaben, Propylparaben, Acrylates Copolymer, Dichlorophenyl Inidazoidioxoian, Blue 1, Fragrance

FRONT LABEL

CVS[®]
pharmacy

NEW!

**daily
therapeutic
shampoo +
conditioner**

2 in 1

**Dandruff Control
Shampoo and
Conditioner**

**Fast-acting
gentle formula**

- Vitamin E & triple conditioners
- Safe for color treated hair
- For daily use

1% Pyrithione Zinc

*Compare to Neutrogena[®]
T/Gel[®] Daily Control

8.5 FL OZ (250 mL)

BACK LABEL

Drug Facts

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|----------------------|---|
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Inactive Ingredients Water, Ammonium Lauryl Sulfate, Sodium Laureth Sulfate, Cocamidopropyl Betaine, Disodium Cocoamphodiacetate, Glycol Stearate, DivinylDimethicone/Dimethicone Copolymer, C12-13 Parath-3, C12-13 Parath-23, Hydrolyzed Wheat Protein PG-Propyl Silanetriol, BHT, Tocopheryl Acetate, Amoldimethicone, Castor Isostearate Succinate, Guar Hydroxypropyltrimonium Chloride, Sodium Methyl Cocoyl Taurate, Methylparaben, Propylparaben, Acrylates Copolymer, Dichlorophenyl Imidazolidioxolan, Blue 1, Fragrance.

*This product is not manufactured or distributed by Neutrogena[®] Corp., distributor of Neutrogena[®] T/Gel Daily[®] Control.

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PYRITHIONE ZINC

pyrithione zinc liquid

Product Information

| | | | |
|-------------------------|----------------|--------------------|---------------|
| Product Type | HUMAN OTC DRUG | Item Code (Source) | NDC:59779-016 |
| Route of Administration | TOPICAL | | |

Active Ingredient/Active Moiety

| Ingredient Name | Basis of Strength | Strength |
|--|-------------------|---------------|
| PYRITHIONE ZINC (UNII: R953O2RHZ5) (PYRITHIONE ZINC - UNII:R953O2RHZ5) | PYRITHIONE ZINC | 29 mg in 1 mL |

Inactive Ingredients

| Ingredient Name | Strength |
|--|----------|
| WATER (UNII: 059QF0KO0R) | |
| AMMONIUM LAURYL SULFATE (UNII: Q7AO2R1M0B) | |
| SODIUM LAURETH SULFATE (UNII: BPV390UAP0) | |
| COCAMIDOPROPYL BETAINE (UNII: 5OCF3O11KX) | |
| BUTYLATED HYDROXYTOLUENE (UNII: 1P9D0Z171K) | |
| .ALPHA.-TOCOPHEROL ACETATE, D- (UNII: A7E6112E4N) | |
| POLYOXYL 100 STEARATE (UNII: YD01N1999R) | |
| METHYLPARABEN (UNII: A2I8C7HI9T) | |
| PROPYLPARABEN (UNII: Z8IX2SC1OH) | |
| COCAMIDOPROPYL BETAINE (UNII: 5OCF3O11KX) | |
| CARBOMER 934 (UNII: Z135WT9208) | |
| SODIUM LAURETH-2 SULFATE (UNII: ZZQ59TY3KG) | |
| DIMETHICONE (UNII: 92RU3N3Y1O) | |
| TRIMETHYL OCTADECYL AMMONIUM CHLORIDE (UNII: CZ70647U92) | |
| DIAZOLIDINYL UREA (UNII: H5RIZ3MPW4) | |

Packaging

| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
|---|------------------|---------------------|----------------------|--------------------|
| 1 | NDC:59779-016-10 | 250 mL in 1 BOTTLE | | |

Marketing Information

| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|---------------------|--|----------------------|--------------------|
| OTC monograph final | part358H | 07/16/2010 | |

Labeler - CVS Pharmacy (062312574)

Registrant - Pharma Pac, LLC (140807475)

Establishment

| Name | Address | ID/FEI | Business Operations |
|-----------------|---------|-----------|---------------------|
| Pharma Pac, LLC | | 140807475 | manufacture |

Revised: 7/2010

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