

**NIOXIN PRO CLINICAL SCALP RECOVERY MOISTURIZING CONDITIONER-
pyrithione zinc lotion
Wella Operations US LLC**

Drug Facts

Active Ingredient

Pyrithione zinc 0.5%

Purpose

Anti-dandruff

Uses

- Helps prevent recurrence of flaking and itching associated with dandruff

Warnings

For external use only.

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.

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.
- Apply to wet scalp and hair, leave for 1-3 minutes. Rinse well.

Inactive ingredients water, stearyl alcohol, cetyl alcohol, stearamidopropyl dimethylamine, dimethicone, glutamic acid, benzyl alcohol, phenoxyethanol, fragrance, methylparaben, mentha arvensis leaf oil, mentha piperita (peppermint) oil, citric acid, propylene glycol, linalool, hexyl cinnamal, camellia sinensis leaf extract, limonene, sodium polynaphthalenesulfonate, geraniol, DMDM hydantoin, cellulose gum, yeast extract, niacinamide, panthenol, biotin, lecithin, tocopheryl acetate, ethoxydiglycol, maltodextrin, glucose, propylparaben, lactic acid, sodium benzoate, equisetum arvense extract, rosmarinus officinalis (rosemary) leaf extract, salvia officinalis (sage) leaf extract, urtica dioica (nettle) extract, achillea millefolium extract, betula alba leaf extract, potassium sorbate, inositol, calcium pantothenate, tartaric acid, caramel.

Questions?

1-800-935-5273

NIOXIN SCALP RECOVERY™

MOISTURIZING CONDITIONER

HELPS TO REDUCE DANDRUFF AND PROVIDE SMOOTHNESS. AS PART OF A COMPLETE SCALP AND HAIR CARE SYSTEM, THIS CONDITIONER LEAVES HAIR NOURISHED,

MOISTURIZED AND PROVIDES RESILIENCE AGAINST BREAKAGE.

MADE IN USA WITH IMPORTED PARTS

Dist. Wella Operations US LLC, Calabasas, CA 91302

www.nioxin.com

nioxin®

PRO CLINICAL

SCALP RECOVERY™

moisturizing conditioner

FOR ANTI-DANDRUFF

FOR ITCHY, FLAKY SCALP

COMBATS DANDRUFF FROM THE 1ST USE

WITH PYRITHIONE ZINC

CLINICALLY & DERMATOLOGICALLY TESTED

200 mL (6.7 FL OZ)

nioxin
— PRO CLINICAL —

SCALP RECOVERY™

*moisturizing
conditioner*

FOR
ANTI-DANDRUFF
FOR ITCHY,
FLAKY SCALP
CORRECTS DANDRUFF
FROM THE 1ST USE
WITH
PYRITHIONE ZINC

CLINICALLY
DERMATOLOGICALLY
TESTED

200mL | 6.7FL OZ

9976815396

nioxin
— PRO CLINICAL —

SCALP RECOVERY*

*moisturizing
conditioner*

FOR
ANTI-DANDRUFF

FOR ITCHY,
FLAKY SCALP

COMBATS DANDRUFF
FROM THE 1ST USE

WITH
PYRITHIONE ZINC

CLINICALLY &
DERMATOLOGICALLY
TESTED

1L | 33.8FL OZ

9926R015398

EAN / UPC Bar Code
Reading direction

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Calabasas, CA 91302
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99268015403

CONDITIONER

pyrithione zinc lotion

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:82157-008
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
PYRITHIONE ZINC (UNII: R953O2RHZ5) (PYRITHIONE ZINC - UNII:R953O2RHZ5)	PYRITHIONE ZINC	0.5 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
DIMETHICONE (UNII: 92RU3N3Y1O)	
MENTHA PIPERITA (PEPPERMINT) OIL (UNII: AV092KU4JH)	
NIACINAMIDE (UNII: 25X51I8RD4)	
STEARYL ALCOHOL (UNII: 2KR89I4H1Y)	
GLUTAMIC ACID (UNII: 3KX376GY7L)	
CALCIUM PANTOTHENATE (UNII: 568ET80C3D)	
CETYL ALCOHOL (UNII: 936JST6JCN)	
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
EQUISETUM ARVENSE BRANCH (UNII: 1L0VKZ185E)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
SAGE (UNII: 065C5D077J)	
CARAMEL (UNII: T9D99G2B1R)	
STEARAMIDOPROPYL DIMETHYLAMINE (UNII: K7VEI00UFR)	
BENZYL ALCOHOL (UNII: LKG8494WBH)	
PANTHENOL (UNII: WW9CM0O67Z)	
LACTIC ACID (UNII: 33X04XA5AT)	
MALTODEXTRIN (UNII: 7CVR7L4A2D)	
GERANIOL (UNII: L837108USY)	
DMDM HYDANTOIN (UNII: BYR0546TOW)	
BIOTIN (UNII: 6S06U10H04)	
.ALPHA.-TOCOPHEROL ACETATE, D- (UNII: A7E6112E4N)	
ETHOXYDIGLYCOL (UNII: A1A1I8X02B)	
ROSEMARY (UNII: IJ67X351P9)	
INOSITOL (UNII: 4L6452S749)	
WATER (UNII: 059QF0KO0R)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
CITRIC ACID (UNII: 2968PHW8QP)	
LINALOOL, (+/-)- (UNII: D81QY6I88E)	
CAMELLIA SINENSIS LEAF (UNII: W2ZU1RY8B0)	
LIMONENE, (+/-)- (UNII: 9MC3I34447)	

BETULA PUBESCENS LEAF (UNII: 84SOH00300)	
MENTHA ARVENSIS LEAF OIL (UNII: 1AEY1M553N)	
.ALPHA.-HEXYLCINNAMALDEHYDE (UNII: 7X6O37OK2I)	
FORMALDEHYDE/SODIUM NAPHTHALENESULFONATE COPOLYMER (3000 MW) (UNII: 90D834OZUI)	
YEAST, UNSPECIFIED (UNII: 3NY3SM6B8U)	
CELLULOSE GUM (UNII: K679OBS311)	
LECITHIN, SOYBEAN (UNII: 1DI56QDM62)	
GLUCOSE (UNII: 5SL0G7R0OK)	
ACHILLEA MILLEFOLIUM (UNII: 2FXJ6SW4PK)	
URTICA DIOICA LEAF (UNII: X6M0DRN46Q)	
TARTARIC ACID (UNII: W4888I119H)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:82157-008-20	200 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	12/19/2024	
2	NDC:82157-008-10	1000 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	12/19/2024	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M032	12/19/2024	

Labeler - Wella Operations US LLC (117781338)

Revised: 12/2024

Wella Operations US LLC