

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

nioxin PRO CLINICAL SCALP RECOVERY moisturizing conditioner

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, phenoxyethanol, dimethicone, glutamic acid, fragrance, mentha arvensis leaf oil, citric acid, propylene glycol, mentha piperita oil, linalool, hexyl cinnamal, camellia sinensis leaf extract, limonene, sodium polynaphthalenesulfonate, geraniol, faex extract/yeast extract/extrait de levure, cellulose gum, biotin, lecithin, niacinamide, panthenol, tocopheryl acetate, ethoxydiglycol, methylisothiazolinone, methylparaben, glucose, propylparaben, lactic acid, sodium benzoate, achillea millefolium extract, betula alba leaf extract, equisetum arvense extract, rosmarinus officinalis (rosemary) leaf extract, salvia officinalis (sage) leaf extract, urtica dioica (nettle) extract, potassium sorbate, calcium pantothenate, inositol, tartaric acid, caramel.

Questions?

1-800-935-5273

NIOXIN SCALP RECOVERY™

MOISTURIZING CONDITIONER

Helps reduce dandruff and protect hair against breakage, leaving hair nourished and moisturized.

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

9926815396

SCALP RECOVERY™

moisturizing conditioner

FOR ANTI-DANDRUFF

FOR ITCHY, FLAKY SCALP

COMBATS DANDRUFF FROM THE 1ST USE

WITH PYRITHIONE ZINC

CLINICALLY & DERMATOLOGICALLY TESTED

1L | 33.8 FL OZ

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

CONDITIONER

pyrithione zinc lotion

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:82157-012
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)	
NIACINAMIDE (UNII: 25X5118RD4)	
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	
BETULA ALBA LEAF (UNII: 84SOH0O3OO)	
CALCIUM PANTOTHENATE (UNII: 568ET80C3D)	
TARTARIC ACID (UNII: W4888I119H)	
MENTHA ARVENSIS LEAF OIL (UNII: 1AEY1M553N)	
CITRIC ACID (UNII: 2968PHW8QP)	
HEXAMETHYLINDANOPYRAN (UNII: 14170060AT)	
BIOTIN (UNII: 6SO6U10H04)	
URTICA DIOICA LEAF (UNII: X6M0DRN46Q)	
INOSITOL (UNII: 4L6452S749)	
WATER (UNII: 059QF0KO0R)	
LECITHIN, SOYBEAN (UNII: 1DI56QDM62)	
PANTHENOL (UNII: WW9CM0O67Z)	
ETHOXYDIGLYCOL (UNII: A1A1I8X02B)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
FORMALDEHYDE/SODIUM NAPHTHALENESULFONATE COPOLYMER (3000 MW) (UNII: 90D834OZUI)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
LINALOOL, (+/-)- (UNII: D81QY6I88E)	
HEXYL CINNAMAL (UNII: 7X6O37OK2I)	
CAMELLIA SINENSIS LEAF (UNII: W2ZU1RY8B0)	
GERANIOL (UNII: L837108USY)	
GLUCOSE (UNII: 5SL0G7R0OK)	
LACTIC ACID (UNII: 33X04XA5AT)	
SAGE (UNII: 065C5D077J)	
CETYL ALCOHOL (UNII: 936JST6JCN)	
STEARAMIDOPROPYL DIMETHYLAMINE (UNII: K7VEI00UFR)	
MENTHA PIPERITA (PEPPERMINT) OIL (UNII: AV092KU4JH)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
STEARYL ALCOHOL (UNII: 2KR89I4H1Y)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	

GLUTAMIC ACID (UNII: 3KX376GY7L)	
LIMONENE, (+/-)- (UNII: 9MC3I34447)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)	
CELLULOSE GUM (UNII: K679OBS311)	
EQUISETUM ARVENSE BRANCH (UNII: 1L0VKZ185E)	
.ALPHA.-TOCOPHEROL ACETATE, D- (UNII: A7E6112E4N)	
ROSEMARY (UNII: IJ67X351P9)	
CARAMEL (UNII: T9D99G2B1R)	
ACHILLEA MILLEFOLIUM (UNII: 2FXJ6SW4PK)	
YEAST, UNSPECIFIED (UNII: 3NY3SM6B8U)	

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:82157-012-20	200 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	04/01/2025	
2	NDC:82157-012-10	1000 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	04/01/2025	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M032	04/01/2025	

Labeler - Wella Operations US LLC (117781338)

Revised: 11/2025

Wella Operations US LLC