

NIOXIN PRO CLINICAL SCALP RECOVERY SCALP SOOTHING SERUM- pyrithione zinc lotion

Wella Operations US LLC

Drug Facts

Active Ingredient

Pyrithione zinc 0.1%

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

- Apply to affected areas one to four times daily or as directed by a doctor.

Inactive ingredients water, cetyl alcohol, stearamidopropyl dimethylamine, stearyl alcohol, distearyldimonium chloride, hydroxyethylcellulose, benzyl alcohol, phenoxyethanol, PEG-2M, dimethicone, fragrance, cetearyl alcohol, methylparaben, glyceryl stearate, oleyl alcohol, mentha arvensis leaf oil, mentha piperita (peppermint) oil, citric acid, polysorbate 60, propylene glycol, camellia sinensis leaf extract, linalool, hexyl cinnamal, limonene, silica, geraniol, benzyl salicylate, alpha-isomethyl ionone, sodium polynaphthalenesulfonate, yeast extract, disodium phosphate, niacinamide, panthenol, biotin, lecithin, tocopheryl acetate, ethoxydiglycol, DMDM hydantoin, maltodextrin, cellulose gum, sodium phosphate, BHT, glucose, propylparaben, lactic acid, sodium benzoate, betula alba leaf extract, achillea millefolium extract, utica dioica (nettle) extract, salvia officinalis (sage) leaf extract, rosmarinus officinalis (rosemary) leaf extract, equisetum arvense extract, potassium sorbate, calcium pantothenate, inositol, caramel, tartaric acid

Questions?

1-800-935-5273

NIOXIN SCALP RECOVERY™

SCALP SOOTHING SERUM

WITH PYRITHIONE ZINC IS A LEAVE-ON TREATMENT THAT HELPS RELIEVE DANDRUFF.

Made in USA WITH IMPORTED PARTS

Dist Wella Operations US LLC. Calabasas, CA 91302

www.nioxin.com

NIOXIN®

PRO CLINICAL

SCALP RECOVERY

scalp soothing serum

FOR ANTI-DANDRUFF

FOR ITCHY FLAKY SCALP

COMBATS DANDRUFF FROM THE 1ST USE

WITH PYRITHIONE ZINC

CLINICALLY & DERMATOLOGICALLY TESTED

100 mL (3.3 FL OZ)

nioxin[®]
— PRO CLINICAL —

SCALP RECOVERY™

*scalp soothing
serum*

FOR
ANTI-DANDRUFF

FOR ITCHY,
FLAKY SCALP

COMBATS DANDRUFF
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99268015400

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DRUG FACTS CONTINUED PEEL HERE

99268015405

Drug Facts (continued)

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Directions
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Drug Facts (continued)

Inactive ingredients
 water, cetyl alcohol, stearamidopropyl dimethylamine, stearyl alcohol, distearyldimonium chloride, hydroxyethylcellulose, benzyl alcohol, phenoxyethanol, PEG-20, dimethicone, fragrance, cetearyl alcohol, methylparaben, glyceryl stearate, oleyl alcohol, mentha arvensis leaf oil, mentha piperita (peppermint) oil, citric acid, polysorbate 60, propylene glycol, camellia sinensis leaf extract, tirakool, hexyl cinnamal limonene, silica, geraniol, benzyl salicylate, alpha-isomethyl ionone, sodium polyacrylate, sodium polyacrylate, sodium phosphate, niacinamide, panthenol, biotin, lecithin, tocopheryl acetate, ethoxydiglycol, DMDM hydantoin, maltodextrin, cellulose gum, sodium phosphate, BHT, glucose, propylparaben, lactic acid, sodium benzoate, betula alba leaf extract, achillea millefolium extract, urtica dioica (nettle) extract, salvia officinalis (sage) leaf extract, rosmarinus officinalis (rosemary) leaf extract, equisetum arvense extract, potassium sorbate, calcium panthothenate, inositol, caramel, tartaric acid.

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pyrithione zinc lotion

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:82157-009
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
PYRITHIONE ZINC (UNII: R953O2RHZ5) (PYRITHIONE ZINC - UNII:R953O2RHZ5)	PYRITHIONE ZINC	0.1 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
BHT (UNII: 1P9D0Z171K)	
ALPHA-ISOMETHYL IONONE (UNII: 9XP4LC555B)	
LINALOOL, (+/-)- (UNII: D81QY6I88E)	
DISODIUM PHOSPHATE (UNII: 22ADO53M6F)	
.ALPHA.-TOCOPHEROL ACETATE, D- (UNII: A7E6112E4N)	
POLYSORBATE 60 (UNII: CAL22UVI4M)	
LIMONENE, (+/-)- (UNII: 9MC3I34447)	
NIACINAMIDE (UNII: 25X51I8RD4)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
MENTHA PIPERITA (PEPPERMINT) OIL (UNII: AV092KU4JH)	
HEXYL CINNAMAL (UNII: 7X6O37OK2I)	
FORMALDEHYDE/SODIUM NAPHTHALENESULFONATE COPOLYMER (3000 MW) (UNII: 90D834OZUI)	
SILICA (UNII: ETJ7Z6XBU4)	
SODIUM PHOSPHATE (UNII: SE337SVY37)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
INOSITOL (UNII: 4L6452S749)	
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)	
PEG-2M (UNII: V46Y6OJ5QB)	
BENZYL SALICYLATE (UNII: WAO5MKN9TU)	
CARAMEL (UNII: T9D99G2B1R)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
MENTHA ARVENSIS LEAF OIL (UNII: 1AEY1M553N)	
STEARAMIDOPROPYL DIMETHYLAMINE (UNII: K7VEI00UFR)	
STEARYL ALCOHOL (UNII: 2KR89I4H1Y)	
GLYCERYL STEARATE (UNII: 230OU9XXE4)	
BIOTIN (UNII: 6SO6U10H04)	
SAGE (UNII: 065C5D077J)	
BETULA PUBESCENS LEAF (UNII: 84SOH00300)	
CALCIUM PANTOTHENATE (UNII: 568ET80C3D)	
HYDROXYETHYLCELLULOSE (UNII: T4V6TWG28D)	
CETYL ALCOHOL (UNII: 936JST6JCN)	
DISTEARYLDIMONIUM CHLORIDE (UNII: OM9573ZX3X)	

PHENOXYETHANOL (UNII: HIE492ZZ3T)
PANTHENOL (UNII: WW9CM0067Z)
LECITHIN, SOYBEAN (UNII: 1DI56QDM62)
ETHOXYDIGLYCOL (UNII: A1A1I8X02B)
DMDM HYDANTOIN (UNII: BYR0546TOW)
MALTODEXTRIN (UNII: 7CVR7L4A2D)
CELLULOSE GUM (UNII: K679OBS311)
GLUCOSE (UNII: 5SL0G7R0OK)
LACTIC ACID (UNII: 33X04XA5AT)
EQUISETUM ARVENSE BRANCH (UNII: 1L0VKZ185E)
ROSEMARY (UNII: IJ67X351P9)
URTICA DIOICA LEAF (UNII: X6M0DRN46Q)
ACHILLEA MILLEFOLIUM (UNII: 2FXJ6SW4PK)
TARTARIC ACID (UNII: W4888I119H)
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)
BENZYL ALCOHOL (UNII: LKG8494WBH)
OLEYL ALCOHOL (UNII: 172F2WN8DV)
CITRIC ACID (UNII: 2968PHW8QP)
CAMELLIA SINENSIS LEAF (UNII: W2ZU1RY8B0)
GERANIOL (UNII: L837108USY)
YEAST, UNSPECIFIED (UNII: 3NY3SM6B8U)
DIMETHICONE (UNII: 92RU3N3Y1O)
CETEARYL ALCOHOL (UNII: 2DMT128M1S)

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:82157-009-01	100 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	12/19/2024	
2	NDC:82157-009-10	1 in 1 CARTON	12/19/2024	
2		100 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		

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)