

## **NIOXIN SCALP RECOVERY MEDICATING CLEANSER- pyrithione zinc shampoo** **Wella Operations US LLC**

*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.*

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### **Nioxin Scalp Recovery Medicating Cleanser**

#### **Drug Facts**

##### **Active ingredient**

Pyrithione zinc 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**

- For best results use at least twice a week or as directed by a doctor.
- Use daily for maximum dandruff control.
- Massage on to wet scalp and hair. Rinse. Repeat if desired.

**Inactive ingredients** water, sodium laureth sulfate, PEG-3 distearate, sodium laureth-8 sulfate, sodium chloride, cocamidopropyl betaine, acrylates/steareth-20 methacrylate copolymer, magnesium laureth-8 sulfate, magnesium laureth sulfate, sodium oleth sulfate, fragrance, sodium benzoate, citric acid, DMDM hydantoin, mentha arvensis leaf oil, mentha piperita (peppermint) oil, magnesium oleth sulfate, sodium hydroxide, polyquaternium-10, propylene glycol, benzoic acid, linalool, hexyl cinnamal, camellia sinensis leaf extract, sodium polynaphthalenesulfonate, benzyl alcohol, limonene, geraniol, cellulose gum, yeast extract, niacinamide, tocopheryl acetate, panthenol, biotin, lecithin, ethoxydiglycol, maltodextrin, methylparaben, glucose, propylparaben, lactic acid, achillea millefolium extract, betula alba leaf extract, urtica dioica (nettle) extract, salvia officinalis (sage) leaf extract, rosmarinus officinalis (rosemary) leaf extract, equisetum arvense extract, potassium sorbate, calcium pantothenate, inositol, tartaric acid, caramel.

**Questions?**

**1-800-935-5273**

**NIOXIN SCALP RECOVERY®**

**MEDICATING CLEANSER**

IS AN ANTIDANDRUFF SHAMPOO WITH PYRITHIONE ZINC THAT HELPS PREVENT AND RELIEVE THE SYMPTOMS OF DANDRUFF, SUCH AS SCALP ITCHING AND FLAKING.

**MADE IN U.S.A. of U.S. and/or imported ingredients**

Dist. Wella Operations US LLC, Calabasas, CA 91302

**[www.nioxin.com](http://www.nioxin.com)**

1-800-935-5273

**NIOXIN®**

**PYRITHIONE ZINC MEDICATING CLEANSER**

STEP 1

SCALP RECOVERY®

**FOR ITCHY, FLAKY SCALP**

COMBATS DANDRUFF FROM THE 1ST USE

**200 mL (6.76 FL OZ)**



**NIOXIN**  
**PYRITHIONE ZINC  
MEDICATING  
CLEANSER**  
**STEP 1**

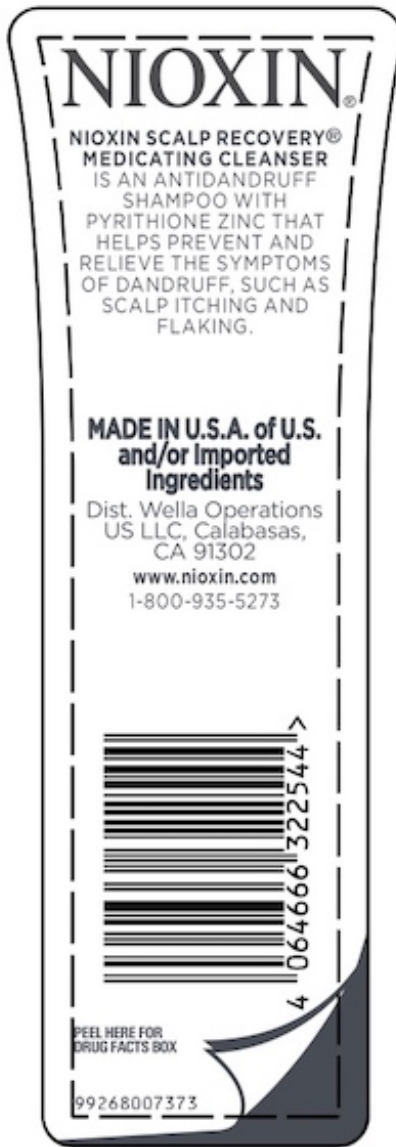
**SCALP RECOVERY®**

**FOR ITCHY  
FLAKY SCALP**  
COMBATS DANDRUFF  
FROM THE 1<sup>ST</sup> USE



**200 mL (6.76 FL. OZ.)**

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## NIOXIN SCALP RECOVERY MEDICATING CLEANSER

pyrithione zinc shampoo

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:82157-003
<b>Route of Administration</b>	TOPICAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>PYRITHIONE ZINC</b> (UNII: R953O2RHZ5) (PYRITHIONE ZINC - UNII:R953O2RHZ5)	PYRITHIONE ZINC	1 g in 100 mL

### Inactive Ingredients

Ingredient Name	Strength
<b>POLYQUATERNIUM-10 (400 MPA.S AT 2%)</b> (UNII: HB1401PQFS)	

<b>FORMALDEHYDE/SODIUM NAPHTHALENESULFONATE COPOLYMER (3000 MW)</b> (UNII: 90D834OZUI)	
<b>CITRIC ACID MONOHYDRATE</b> (UNII: 2968PHW8QP)	
<b>PROPYLENE GLYCOL</b> (UNII: 6DC9Q167V3)	
<b>LIMONENE, (+/-)-</b> (UNII: 9MC3I34447)	
<b>SODIUM OLEYL SULFATE</b> (UNII: 46F21X4586)	
<b>COCAMIDOPROPYL BETAINE</b> (UNII: 5OCF3O11KX)	
<b>ETHYL ACRYLATE/METHACRYLIC ACID/STEARETH-20 METHACRYLATE COPOLYMER</b> (UNII: EPA1872R1N)	
<b>MAGNESIUM LAURETH-8 SULFATE</b> (UNII: 2OTJ9LF5UA)	
<b>MAGNESIUM LAURETH SULFATE</b> (UNII: UKW9G007TZ)	
<b>SODIUM HYDROXIDE</b> (UNII: 55X04QC32I)	
<b>BIOTIN</b> (UNII: 6SO6U10H04)	
<b>SODIUM LAURETH-8 SULFATE</b> (UNII: YP8U3694P0)	
<b>GERANIOL</b> (UNII: L837108USY)	
<b>.ALPHA.-TOCOPHEROL ACETATE</b> (UNII: 9E8X80D2L0)	
<b>DIETHYLENE GLYCOL MONOETHYL ETHER</b> (UNII: A1A18X02B)	
<b>ROSEMARY</b> (UNII: IJ67X351P9)	
<b>INOSITOL</b> (UNII: 4L6452S749)	
<b>LACTIC ACID</b> (UNII: 33X04XA5AT)	
<b>BETULA PUBESCENS LEAF</b> (UNII: 84SOH00300)	
<b>POTASSIUM SORBATE</b> (UNII: 1VPU26JZZ4)	
<b>SAGE</b> (UNII: 065C5D077J)	
<b>ACHILLEA MILLEFOLIUM</b> (UNII: 2FXJ6SW4PK)	
<b>CALCIUM PANTOTHENATE</b> (UNII: 568ET80C3D)	
<b>BENZOIC ACID</b> (UNII: 8SKN0B0MIM)	
<b>.ALPHA.-HEXYLCINNAMALDEHYDE</b> (UNII: 7X6O37OK2I)	
<b>CARBOXYMETHYLCELLULOSE SODIUM, UNSPECIFIED FORM</b> (UNII: K679OBS311)	
<b>MALTODEXTRIN</b> (UNII: 7CVR7L4A2D)	
<b>URTICA DIOICA LEAF</b> (UNII: X6M0DRN46Q)	
<b>EQUISETUM ARVENSE BRANCH</b> (UNII: 1L0VKZ185E)	
<b>TARTARIC ACID</b> (UNII: W4888I119H)	
<b>CARAMEL</b> (UNII: T9D99G2B1R)	
<b>PROPYLPARABEN</b> (UNII: Z8IX2SC1OH)	
<b>LECITHIN, SOYBEAN</b> (UNII: 1DI56QDM62)	
<b>NIACINAMIDE</b> (UNII: 25X51I8RD4)	
<b>PANTHENOL</b> (UNII: WW9CM0067Z)	
<b>METHYLPARABEN</b> (UNII: A2I8C7HI9T)	
<b>DEXTROSE, UNSPECIFIED FORM</b> (UNII: IY9XDZ35W2)	
<b>SODIUM CHLORIDE</b> (UNII: 451W47IQ8X)	
<b>MAGNESIUM OLETH SULFATE</b> (UNII: LCQ4WP3DTL)	
<b>LINALOOL, (+/-)-</b> (UNII: D81QY6I88E)	
<b>WATER</b> (UNII: 059QF0KO0R)	
<b>SODIUM LAURETH-3 SULFATE</b> (UNII: BPV390UAP0)	
<b>PEG-3 DISTEARATE</b> (UNII: 8420ECX438)	
<b>SODIUM BENZOATE</b> (UNII: OJ245FE5EU)	
<b>DMDM HYDANTOIN</b> (UNII: BYR0546TOW)	
<b>PEPPERMINT OIL</b> (UNII: AV092KU4JH)	
<b>MENTHA ARVENSIS LEAF OIL</b> (UNII: 1AEY1M553N)	
<b>BENZYL ALCOHOL</b> (UNII: LKG8494WBH)	
<b>GREEN TEA LEAF</b> (UNII: W2ZU1RY8B0)	
<b>YEAST, UNSPECIFIED</b> (UNII: 3NY3SM6B8U)	

**Packaging**

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:82157-003-20	200 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	12/01/2022	

**Marketing Information**

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	M032	12/01/2022	

**Labeler** - Wella Operations US LLC (117781338)

Revised: 12/2022

Wella Operations US LLC