NEUTROGENA SCALP THERAPY ANTI-DANDRUFF DAILY CONTROL- salicylic acid shampoo
Johnson & Johnson Consumer Inc.

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.

NEUTROGENA ® Scalp Therapy ™ Anti-Dandruff Daily Control Shampoo

Drug Facts

Active ingredient

Salicylic Acid (1.8%)

Purposes

Anti-dandruff, Anti-seborrheic dermatitis, Anti-psoriasis

Use

Controls the symptoms of dandruff, seborrheic dermatitis and psoriasis.

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 the 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.
- Wet hair thoroughly.
- Massage shampoo into your scalp.
- Lather, leaving on your hair and scalp for a few minutes.
- Rinse and repeat.

Other information

Store at room temperature.

Inactive ingredients

Water, Sodium C14-16 Olefin Sulfonate, Cocamidopropyl Betaine, Sodium Chloride, Polyquaternium-22, Sodium Citrate, Hexylene Glycol, Sodium Lauroyl Sarcosinate, Fragrance, Linoleamidopropyl PG-Dimonium Chloride Phosphate, Citric Acid, Sodium Hydroxide

Questions?

Call **800-256-4247** or **215-273-8755** (collec **t)** or visit www.neutrogena.com

Distributed by:
JOHNSON & JOHNSON CONSUMER INC.
Skillman, NJ 08558

PRINCIPAL DISPLAY PANEL - 354 mL Bottle Carton

NEUTROGENA

Scalp Therapy ™

Anti-Dandruff

MEDICATED

DAILY CONTROL

1.8% SALICYLIC ACID

SHAMPOO

Neutrogena®

DERMATOLOGIST RECOMMENDED BRAND

12 FL OZ (354 mL)

NEUTROGENA SCALP THERAPY ANTI-DANDRUFF DAILY CONTROL

salicylic acid shampoo

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:69968-0722
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
SALICYLIC ACID (UNII: O414PZ4LPZ) (SALICYLIC ACID - UNII:O414PZ4LPZ)	SALICYLIC ACID	18 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
SODIUM C14-16 OLEFIN SULFONATE (UNII: O9W3D3YF5U)	
COCAMIDOPROPYL BETAINE (UNII: 5OCF3O11KX)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
SODIUM CITRATE, UNSPECIFIED FORM (UNII: 1Q73Q2JULR)	
SODIUM LAUROYL SARCOSINATE (UNII: 632GS99618)	
HEXYLENE GLYCOL (UNII: KEH0A3F75J)	
LINOLEAMIDOPROPYL PROPYLENE GLYCOL-DIMONIUM CHLORIDE PHOSPHATE (UNII: 5Q87K461JO)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69968-	354 mL in 1 BOTTLE, PLASTIC; Type 0: Not a	12/06/2021	

1	0722-1	Combination Product	12/06/2021	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part358H	12/06/2021	

Labeler - Johnson & Johnson Consumer Inc. (118772437)

Revised: 1/2023

Johnson & Johnson Consumer Inc.