DENOREX EXTRA STRENGTH- salicylic acid shampoo Neoteric Cosmetics, Inc.

Denorex Extra Strength

Drug Facts

Active Ingredients:

Salicylic acid 3%

Purpose

Controls the symptoms of dandruff, seborrheic dermatitis and psoriasis

Uses

- Reduces and helps eliminate scalp itching, flaking and scaling associated with dandruff, seborrheic dermatitis and psoriasis
- Helps prevent recurrence of 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

- Shake well
- apply to wet hair
- gently massage into hair and scalp to work up a lather
- rinse thoroughly and repeat

• for best results, use at least twice weekly or as directed by a doctor

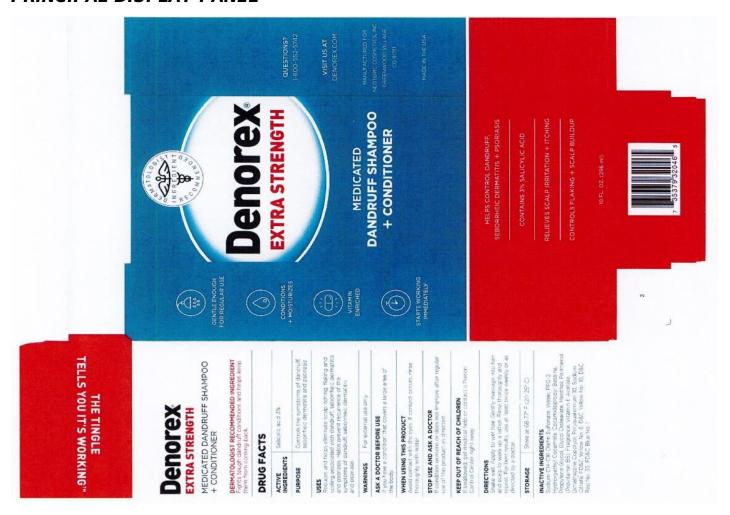
Other information

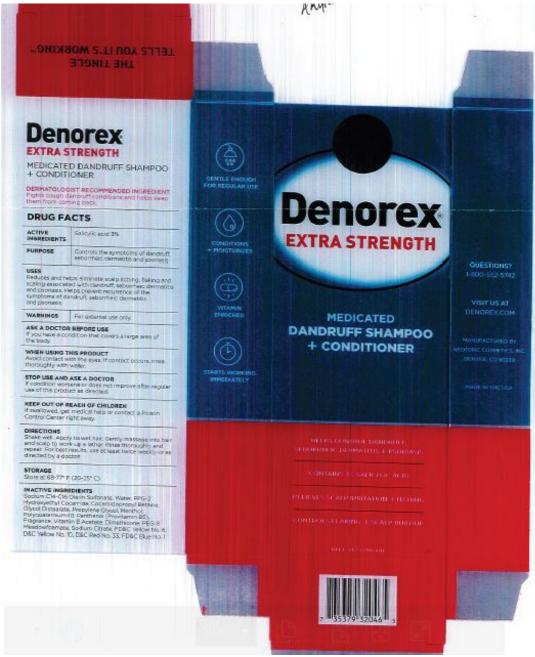
Store at 20º-25º C (68º - 77º F)

Inactive ingredients

cocamidopropyl betaine, D&C Red No. 33, D&C Yellow No. 10, dimethicone PEG-8 meadowfoamate, FD&C Blue No. 1, FD&C Yellow No. 6, fragrance, glycol distearate, menthol, panthenol, polyquaternium-10, PPG-2 hydroxyethyl cocamide, propylene glycol, sodium C14-C16 olefin sulfonate, sodium citrate, vitamin E acetate, water

PRINCIPAL DISPLAY PANEL





EXTRA STRENGTH

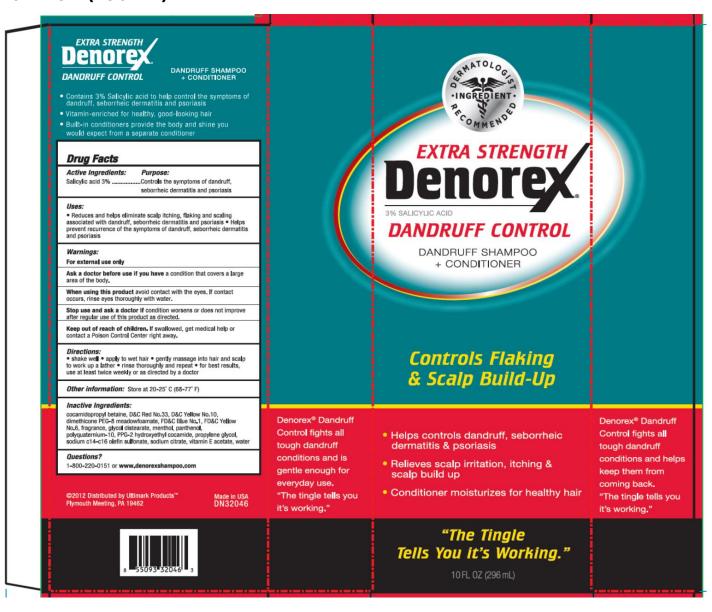
DENOREX

3 % Salicylic Acid

DANDRUFF Control

SHAMPOO +

10 FL OZ (296 mL)



DENOREX EXTRA STRENGTH

salicylic acid shampoo

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:62673-058
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety			
	Ingredient Name	Basis of Strength	Strength
	SALICYLIC ACID (UNII: O414PZ4LPZ) (SALICYLIC ACID - UNII:O414PZ4LPZ)	SALICYLIC ACID	0.03 g in 1 mL

Inactive Ingredients		
Ingredient Name	Strength	
WATER (UNII: 059QF0KO0R)		
SODIUM C14-16 OLEFIN SULFONATE (UNII: O9W3D3YF5U)		
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)		
GLYCOL DISTEARATE (UNII: 13W7MDN21W)		
MENTHOL (UNII: L7T10EIP3A)		
COCAMIDOPROPYL BETAINE (UNII: 50CF3011KX)		
PANTHENOL (UNII: WW9CM0O67Z)		
.ALPHATOCOPHEROL ACETATE (UNII: 9E8X80D2L0)		
SODIUM CITRATE (UNII: 1Q73Q2JULR)		
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)		
D&C YELLOW NO. 10 (UNII: 35SW5USQ3G)		
D&C RED NO. 33 (UNII: 9DBA0SBB0L)		
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)		

l	Packaging				
	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
	1	NDC:62673-058- 10	296 mL in 1 BOTTLE; Type 0: Not a Combination Product	12/13/2017	

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC Monograph Drug	M032	11/01/2009		

Labeler - Neoteric Cosmetics, Inc. (790615181)

Registrant - Pharma Nobis, LLC (118564114)

Establishment				
Name	Address	ID/FEI	Business Operations	
Pharma Nobis, LLC		118564114	analysis(62673-058), manufacture(62673-058), pack(62673-058), label(62673-058)	

Revised: 12/2024 Neoteric Cosmetics, Inc.