RENEWAL ANTI-DANDRUFF- salicylic acid liquid
RITE AID CORPORATION

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.

DRUG FACTS

ACTIVE INGREDIENT

SALICYLIC ACID 2%

PURPOSE

ANTI-DANDRUFF/ ANTI-SEBORRHEIC DERMATITIS/ ANTI-PSORIASIS

USES

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 EYES. IF CONTACT OCCURS, RINSE EYES THOROUGHLY WITH WATER
- FOR USE ON COLOR TREATED OR PERMED HAIR, RINSE THOROUGHLY

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

IN CASE OF ACCIDENTAL INGESTION, GET MEDICAL HELP OR CONTACT A POISON CONTROL CENTER IMMEDIATELY

DIRECTIONS

- FOR BEST RESULTS, USE TWICE A WEEK OR AS DIRECTED BY A DOCTOR
- WET HAIR, MASSAGE ONTO SCALP FOR SEVERAL MINUTES, AND THEN RINSE
- REPEAT IF DESIRED

INACTIVE INGREDIENTS

WATER (AQUA), SODIUM LAURETH SULFATE, ACRYLATES CROSSPOLYMER-4, COCAMIDE MEA, SODIUM COCOYL SARCOSINATE, GLYCOL DISTEARATE, POLYQUATERNIUM-7, FRARANCE (PARFUM), SODIUM CHLORIDE, SODIUM HYDROXIDE,

TETRASODIUM EDTA, BHT, METHYLCHLOROISOTHIAZOLINONE, METHYLISOTHIAZOLINONE, BLUE 1 (CI 42090)

QUESTIONS OR COMMENTS?

1-866-695-3030

LABEL COPY

RENEWAL ANTI-DANDRUFF

salicylic acid liquid

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:11822-4500
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
SALICYLIC ACID (UNII: O414PZ4LPZ) (SALICYLIC ACID - UNII:O414PZ4LPZ)	SALICYLIC ACID	20 mg in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
SODIUM LAURETH SULFATE (UNII: BPV390UAP0)	
CARBOMER COPOLYMER TYPE A (ALLYL PENTAERYTHRITOL CROSSLINKED) (UNII: 71DD5V995L)	
COCO MONOETHANOLAMIDE (UNII: C80684146D)	
SODIUM COCOYL SARCOSINATE (UNII: 1R9DUY89CZ)	
GLYCOL DISTEARATE (UNII: 13W7MDN21W)	
POLYQUATERNIUM-7 (70/30 ACRYLAMIDE/DADMAC; 1600000 MW) (UNII: 0L414VCS5Y)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
EDETATE SODIUM (UNII: MP1J8420LU)	
BUTYLATED HYDROXYTOLUENE (UNII: 1P9D0Z171K)	
METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)	
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:11822-4500-7	207 mL in 1 BOTTLE, PLASTIC		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part333D	12/08/2014	

Labeler - RITE AID CORPORATION (014578892)

Registrant - APOLLO HEALTH AND BEAUTY CARE (201901209)

Establishment

Name	Address	ID/FEI	Business Operations
APOLLO HEALTH AND BEAUTY CARE		201901209	manufacture(11822-4500)

Revised: 12/2014 RITE AID CORPORATION