

**RENEWAL DANDRUFF MOISTURIZING- selenium sulfide 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.*

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**DRUG FACTS**

**ACTIVE INGREDIENT**

SELENIUM SULFIDE 1%

**PURPOSE**

ANTI-DANDRUFF/ANTI-SEBORRHEIC DERMATITIS

**USES**

FOR THE RELIEF OF FLAKING AND ITCHING ASSOCIATED WITH DANDRUFF AND SEBORRHEIC DERMATITIS AND TO HELP PREVENT THE CHANCE OF RECURRENCE

**WARNINGS**

FOR EXTERNAL USE ONLY

**ASK A DOCTOR BEFORE USE IF YOU HAVE**

SEBORRHEIC DERMATITIS IN AREAS OTHER THAN THE SCALP

**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**

- SHAKE WELL, APPLY SHAMPOO AND RINSE THOROUGHLY
- FOR BEST RESULTS, USE AT LEAST TWICE A WEEK OR AS DIRECTED BY A DOCTOR

**OTHER INFORMATION**

STORE AT ROOM TEMPERATURE

**INACTIVE INGREDIENTS**

WATER (AQUA), SODIUM LAURETH SULFATE, DISTEARYL PHTHALIC ACID AMIDE, SODIUM CHLORIDE, COCAMIDOPROPYL BETAINE, TITANIUM DIOXIDE, SODIUM STEAROYL LACTYLATE, FRAGRANCE (PARFUM), DIMETHICONE, CITRIC ACID, DMDM HYDANTOIN, SODIUM CITRATE, ALOE BARBADENSIS LEAF JUICE, HYDROXYPROPYL METHYLCELLULOSE, BLUE 1 (CI 42090)

**LABEL COPY**



**RENEWAL DANDRUFF MOISTURIZING**

selenium sulfide liquid

**Product Information**

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:11822-6221
<b>Route of Administration</b>	TOPICAL		

**Active Ingredient/Active Moiety**

Ingredient Name	Basis of Strength	Strength
SELENIUM SULFIDE (UNII: Z69D9E381Q) (SELENIUM SULFIDE - UNII:Z69D9E381Q)	SELENIUM SULFIDE	10 mg in 1 mL

**Inactive Ingredients**

Ingredient Name	Strength
WATER (UNII: 059QF0K00R)	
SODIUM LAURETH SULFATE (UNII: BPV390UAP0)	
DISTEARYL PHTHALAMIC ACID (UNII: 5552GSZ9LI)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
COCAMIDOPROPYL BETAINE (UNII: 5OCF3O11KX)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
SODIUM STEAROYL LACTYLATE (UNII: IN99IT31LN)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
DMDM HYDANTOIN (UNII: BYR0546TOW)	
SODIUM CITRATE (UNII: 1Q73Q2JULR)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
HYPROMELLOSES (UNII: 3NXW29V3WO)	
FD&C BLUE NO. 1 (UNII: HBR47K3TBD)	

**Packaging**

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11822-6221-1	325 mL in 1 BOTTLE, PLASTIC		

**Marketing Information**

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part358H	03/03/2015	

**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(11822-6221)