REXALL ALOE- selenium sulfide liquid
DOLGENCORP 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.

DRUG FACTS

ACTIVE INGREDIENT

SELENIUM SULFIDE 1%

PURPOSE

ANTI-DANDRUFF, ANTI-SEBORRHEIC DERMATITIS

USES

FOR RELIEF OF FLAKING AND ITCHING ASSOCATED WITH DANDRUFF AND SEBORRHEIC DERMATITIS AND TO HELP PREVENT THE CHANCE OF RE-OCCURENCE

WARNINGS

FOR EXTERNAL USE ONLY

ASK A DOCTOR BEFORE USING IF YOU HAVE

SEBORRHEIC DERMATITIS IN AREAS OTHER THAN THE SCALP

WHEN USING THIS PRODUCT

AVOID CONTACT WITH EYES. IF PRODUCT GETS INTO EYES, RINSE THOROUGHLY WITH WATER. FOR USE ON COLOR TREATED OR PERMED HAIR, RINSE THOROUGHLY

STOP USING THIS PRODUCT 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

INACTIVE INGREDIENTS

WATER (AQUA), SODIUM LAURETH SULFATE, DISTEARYL PHTHALIC ACID AMIDE, SODIUM CHLRODIE, COCAMIDOPROPYL BETAINE, SODIUM STEAROYL LACTYLATE, FRAGRANCE (PARFUM), DIMETHICONE, CITRIC ACID, DMDM HYDANTOIN, SODIUM CITRATE, ALOE BARBADENSIS LEAF JUICE, HYDROXYPROPYL METHYLCELLULOSE,

TITANIUM DIOXIDE, BLUE 1 (CI 42090)

QUESTIONS OR COMMENTS?

1-877-932-7948

LABEL COPY

REXALL ALOE

selenium sulfide liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:55910-621
Route of Administration	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: 059QF0KO0R)	
SODIUM LAURETH SULFATE (UNII: BPV390UAP0)	
DISTEARYL PHTHALAMIC ACID (UNII: 5552GSZ9LI)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
COCAMIDOPROPYL BETAINE (UNII: 5OCF3O11KX)	
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)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:55910-621-12	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	01/15/2015	

Labeler - DOLGENCORP INC (068331990)

Registrant - APOLLO HEALTH AND BEAUTY CARE (201901209)

Establishment

Name	Address	ID/FEI	Business Operations
APOLLO HEALTH AND BEAUTY CARE		201901209	manufacture(55910-621)

Revised: 1/2015 DOLGENCORP INC