CARE ONE MOISTURIZING- selenium sulfide liquid
AMERICAN SALES COMPANY

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

ANTIDANDRUFF

USE

CONTROLS FLAKING, SCALING AND ITCHING ASSOCIATED WITH DANDRUFF.

WARNINGS

FOR EXTERNAL USE ONLY.

DO NOT USE

ON SCALP THAT IS BROKEN OR INFLAMED, IF YOU ARE ALLERGIC TO INGREDIENTS IN THIS PRODUCT.

WHEN USING THIS PRODUCT

AVOID CONTACT WITH EYES. IF PRODUCT GETS INTO EYES, 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. SHAMPOO, THEN RINSE THOROUGHLY. FOR BEST RESULTS, USE AT LEAST TWICE A WEEK OR AS DIRECTED BY A DOCTOR.

OTHER INFORMATION

STORE AT ROOM TEMPERATURE.

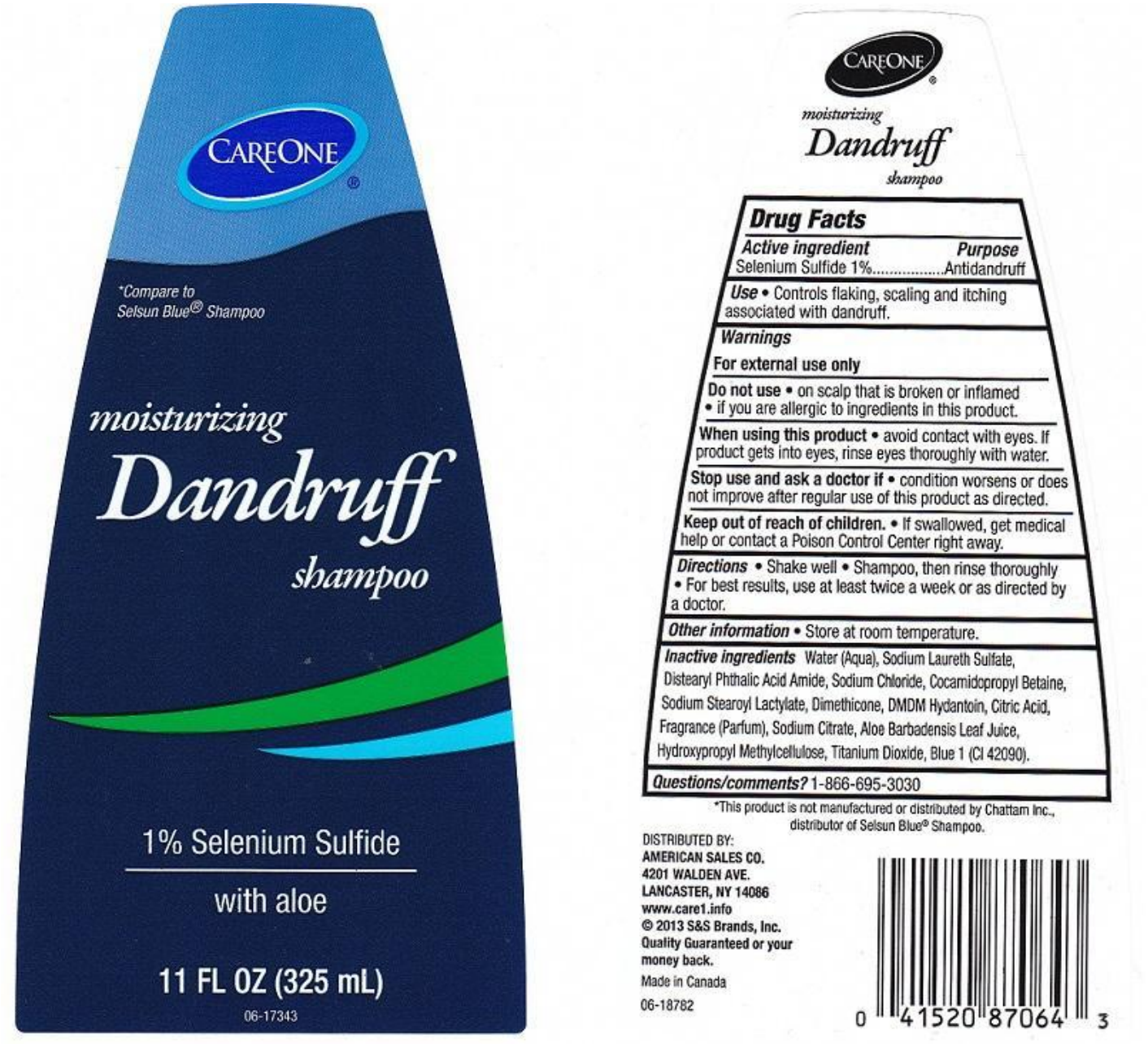
INACTIVE INGREDIENTS:

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

QUESTIONS/COMMENTS?

1-866-695-3030

LABEL COPY



CARE ONE MOISTURIZING
selenium sulfide liquid

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:41520-621
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)	
DMDM HYDANTOIN (UNII: BYR0546TOW)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
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:41520-621-11	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	07/15/2013	

Labeler - AMERICAN SALES COMPANY (809183973)

Registrant - APOLLO HEALTH AND BEAUTY CARE (201901209)

Establishment

Name	Address	ID/FEI	Business Operations
APOLLO HEALTH AND BEAUTY CARE		201901209	manufacture(41520-621)