EQUALINE CLINICAL STRENGTH- selenium sulfide liquid
SUPERVALU 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

HELPS PREVENT THE CHANCE OF RECURRENCE OF FLAKING, ITCHING, IRRITATION, SCALING AND REDNESS ASSOCIATED WITH DANDRUFF AND SEBORRHEIC DERMATITIS.

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, RINSE THOROUGHLY. FOR BEST RESULTS, USE AT LEAST TWICE A WEEK OR AS DIRECTED BY A DOCTOR.

INACTIVE INGREDIENTS

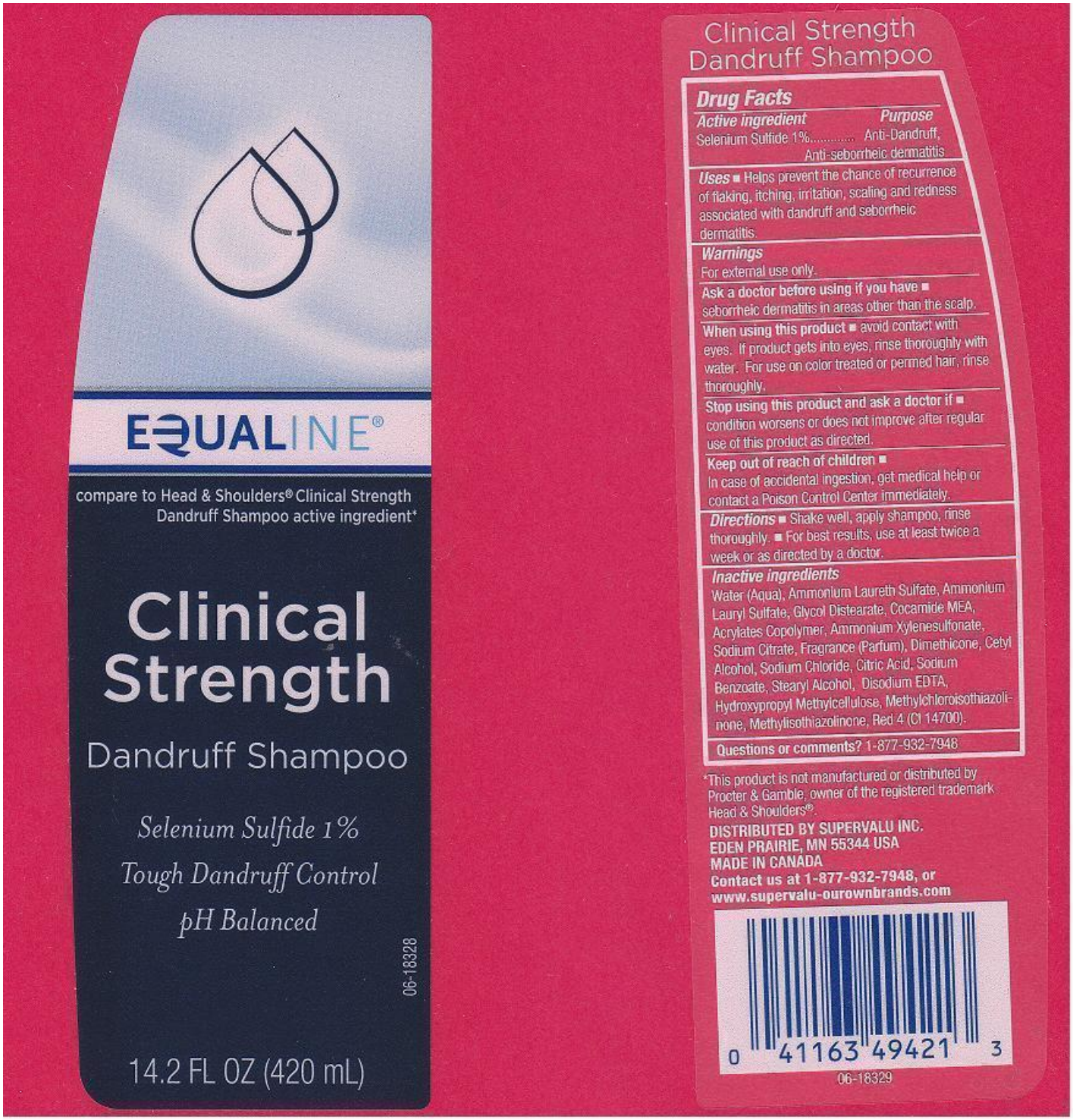
WATER (AQUA), AMMONIUM LAURETH SULFATE, AMMONIUM LAURYL SULFATE, GLYCOL DISTEARATE, COCAMIDE MEA, ACRYLATES COPOLYMER, AMMONIUM XYLENESULFONATE, SODIUM CITRATE, FRAGRANCE (PARFUM), DIMETHICONE, CETYL ALCOHOL, SODIUM CHLORIDE, CITRIC ACID, SODIUM BENZOATE, STEARYL ALCOHOL,

DISODIUM EDTA, HYDROXYPROPYL METHYLCELLULOSE, METHYLCHLOROISOTHIAZOLINONE, METHYLISOTHIAZOLINONE, RED 4 (CI 14700).

QUESTIONS OR COMMENTS?

1-877-932-7948

LABEL COPY



EQUALINE CLINICAL STRENGTH
selenium sulfide liquid

Product Information			
Product Type	HUMAN OTC DRUG LABEL	**Item Code (Source)**	NDC:41163-618
Route of Administration	TOPICAL	**DEA Schedule**	

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
SELENIUM SULFIDE (SELENIUM SULFIDE)	SELENIUM SULFIDE	10 mg in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**
WATER	
AMMONIUM LAURETH-3 SULFATE	
AMMONIUM LAURYL SULFATE	
GLYCOL DISTEARATE	
COCO MONOETHANOLAMIDE	
METHACRYLIC ACID - METHYL METHACRYLATE COPOLYMER (1:1)	
AMMONIUM XYLENESULFONATE	
SODIUM CITRATE	
DIMETHICONE	
CETYL ALCOHOL	
SODIUM CHLORIDE	
CITRIC ACID MONOHYDRATE	
SODIUM BENZOATE	
STEARYL ALCOHOL	
EDETATE DISODIUM	
HYPROMELLOSES	
METHYLCHLOROISOTHIAZOLINONE	
METHYLISOTHIAZOLINONE	
FD&C RED NO. 4	

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:41163-618-14	420 mL in 1 BOTTLE, PLASTIC		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part358H	02/14/2013	

Labeler - SUPERVALU INC. (006961411)

Registrant - APOLLO HEALTH AND BEAUTY CARE (201901209)

Establishment

Name	Address	ID/FEI	Business Operations
APOLLO HEALTH AND BEAUTY CARE		201901209	manufacture(41163-618)

Revised: 2/2013

SUPERVALU INC.