EQUALINE CLASSIC CLEAN 2 IN 1- pyrithione zinc 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

PYRITHIONE ZINC 1%

PURPOSE

ANTI-DANDRUFF

USES

TO HELP PREVENT RECURRENCE OF FLAKING AND ITCHING ASSOCIATED WITH DANDRUFF.

WARNINGS

FOR EXTERNAL USE ONLY.

WHEN USING THIS PRODUCT

AVOID CONTACT WITH EYES. IF CONTACT OCCURS, RINSE WITH WATER.

STOP USING THIS PRODUCT AND ASK 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 MAXIMUM DANDRUFF CONTROL, USE EVERY TIME YOU SHAMPOO. WET HAIR, MASSAGE ONTO SCALP AND RINSE. REPEAT IF DESIRED.

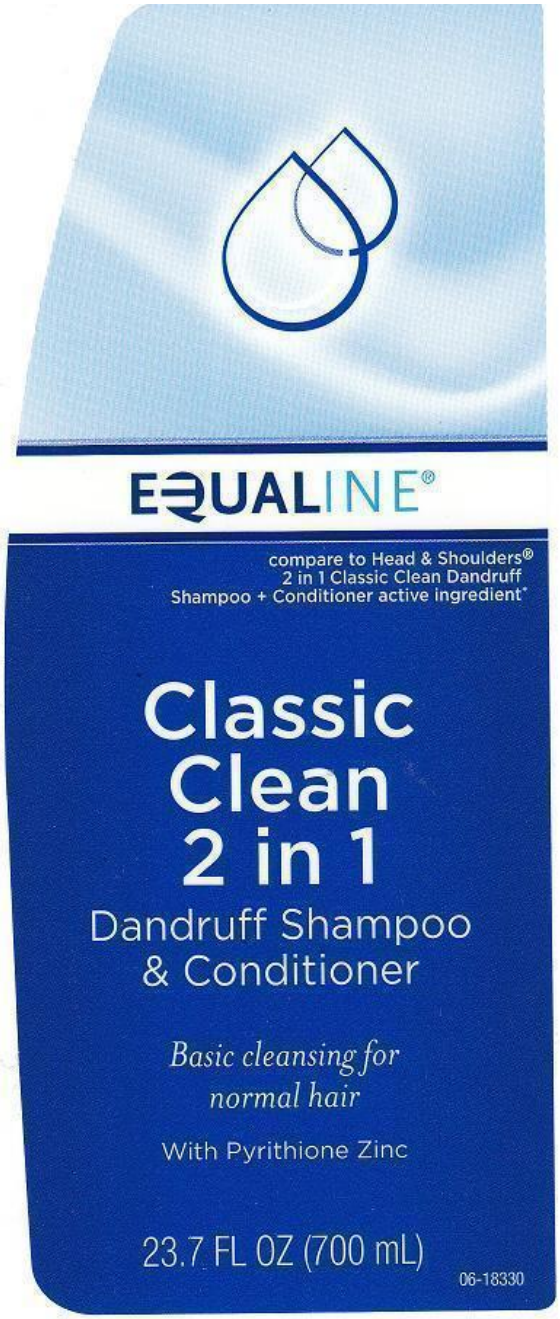
INACTIVE INGREDIENTS:

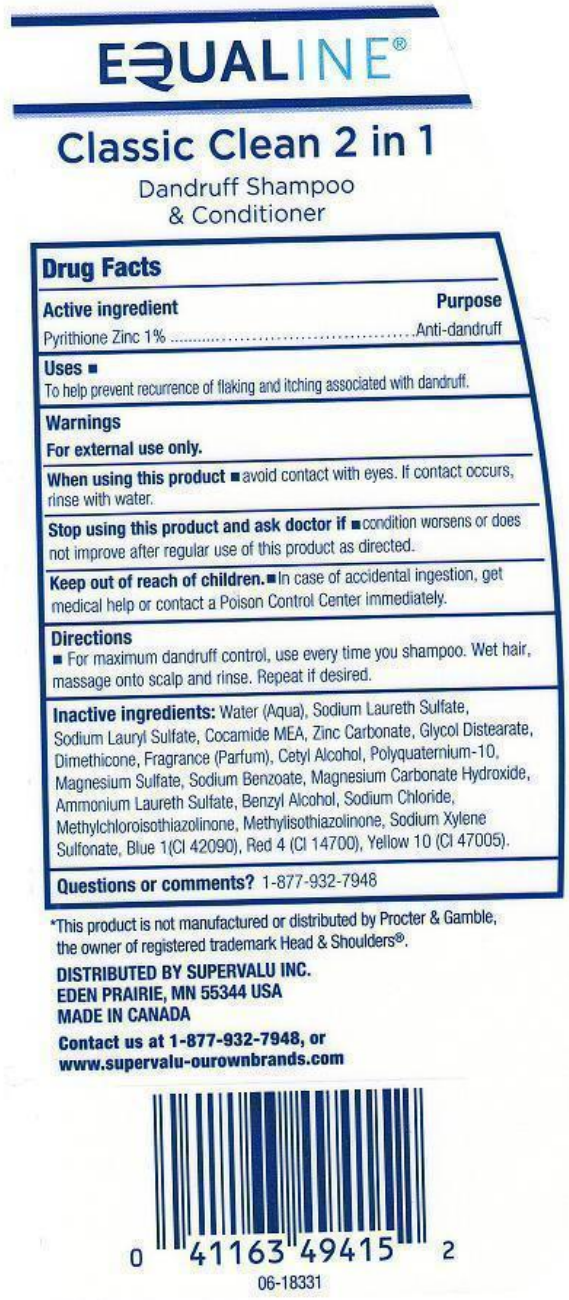
WATER (AQUA), SODIUM LAURETH SULFATE, SODIUM LAURYL SULFATE, COCAMIDE MEA, ZINC CARBONATE, GLYCOL DISTEARATE, DIMETHICONE, FRAGRANCE (PARFUM), CETYL ALCOHOL, POLYQUATERNIUM-10, MAGNESIUM SULFATE, SODIUM BENZOATE, MAGNESIUM CARBONATE HYDROXIDE, AMMONIUM LAURETH SULFATE, BENZYL ALCOHOL, SODIUM CHLORIDE, METHYLCHLOROISOTHIAZOLINONE, METHYLISOTHIAZOLINONE, SODIUM XYLENE SULFONATE, BLUE 1 (CI 42090), RED 4 (CI 14700), YELLOW 10 (CI 47005).

QUESTIONS OR COMMENTS?

1-877-932-7948

LABEL COPY





EQUALINE CLASSIC CLEAN 2 IN 1

pyrithione zinc liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:41163-417
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
PYRITHIONE ZINC (UNII: R953O2RHZ5) (PYRITHIONE ZINC - UNII:R953O2RHZ5)	PYRITHIONE ZINC	10 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
SODIUM LAURETH SULFATE (UNII: BPV390UAP0)	
SODIUM LAURYL SULFATE (UNII: 368GB5141J)	
COCO MONOETHANOLAMIDE (UNII: C80684146D)	
ZINC CARBONATE (UNII: EQR32Y7H0M)	
GLYCOL DISTEARATE (UNII: 13W7MDN21W)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
CETYL ALCOHOL (UNII: 936JST6JCN)	
POLYQUATERNIUM-10 (400 CPS AT 2%) (UNII: HB1401PQFS)	
MAGNESIUM SULFATE (UNII: DE08037SAB)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
MAGNESIUM CARBONATE HYDROXIDE (UNII: YQO029V1L4)	
AMMONIUM LAURETH-3 SULFATE (UNII: 896SJ235FN)	
BENZYL ALCOHOL (UNII: LKG8494WBH)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)	
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	
SODIUM XYLENESULFONATE (UNII: G4LZF950UR)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
FD&C RED NO. 4 (UNII: X3W0AM1JLX)	
D&C YELLOW NO. 10 (UNII: 35SW5USQ3G)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:41163-417-24	700 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(41163-417)