**EQUALINE MEDICATED DANDRUFF- selenium sulfide shampoo**
**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.*

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**DRUG FACTS**

**ACTIVE INGREDIENT**

SELENIUM SULFIDE 1%

**PURPOSE**

ANTIDANDRUFF

**USES**

FOR RELIEF OF FLAKING AND ITCHING ASSOCIATED 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 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.

**QUESTIONS? COMMENTS?**

1-877-932-7943

**INACTIVE INGREDIENTS**

WATER (AQUA), AMMONIUM LAURYL SULFATE, TEA-LAURYL SULFATE, AMMONIUM

LAURETH SULFATE, COCAMIDOPROPYL BETAINE, MAGNESIUM ALUMINUM SILICATE, FRAGRANCE (PARFUM), MENTHOL, COCAMIDE DEA, DMDM HYDANTOIN, CITRIC ACID, HYDROXYPROPYL METHYLCELLULOSE, SODIUM CITRATE, SODIUM CHLORIDE, BLUE 1 (CI 42090), RED 33 (CI 17200).

**LABEL COPY**


## EQUALINE MEDICATED DANDRUFF

selenium sulfide shampoo

| Product Information | | | |
|---|---|---|---|
| **Product Type** | HUMAN OTC DRUG | **Item Code (Source)** | NDC:41163-617 |
| **Route of Administration** | TOPICAL | | |

| Active Ingredient/Active Moiety | | |
|---|---|---|
| **Ingredient Name** | **Basis of Strength** | **Strength** |
| **SELENIUM SULFIDE** (UNII: Z69D9E381Q) (SELENIUM - UNII:H6241UJ22B) | SELENIUM SULFIDE | 1 mL in 100 mL |

| Inactive Ingredients | |
|---|---|
| **Ingredient Name** | **Strength** |
| **WATER** (UNII: 059QF0KO0R) | |
| **AMMONIUM LAURYL SULFATE** (UNII: Q7AO2R1M0B) | |
| **TROLAMINE LAURYL SULFATE** (UNII: E8458C1KAA) | |
| **AMMONIUM LAURETH-2 SULFATE** (UNII: 698O4Z48G6) | |
| **COCAMIDOPROPYL BETAINE** (UNII: 5OCF3O11KX) | |
| **MAGNESIUM ALUMINUM SILICATE** (UNII: 6M3P64V0NC) | |
| **MENTHOL** (UNII: L7T10EIP3A) | |
| **COCO DIETHANOLAMIDE** (UNII: 92005F972D) | |
| **DMDM HYDANTOIN** (UNII: BYR0546TOW) | |
| **CITRIC ACID MONOHYDRATE** (UNII: 2968PHW8QP) | |
| **HYPROMELLOSE 2208 (100 MPA.S)** (UNII: B1QE5P712K) | |
| **SODIUM CITRATE** (UNII: 1Q73Q2JULR) | |
| **SODIUM CHLORIDE** (UNII: 451W47IQ8X) | |
| **FD&C BLUE NO. 1** (UNII: H3R47K3TBD) | |
| **D&C RED NO. 33** (UNII: 9DBA0SBB0L) | |

| Packaging | | | | |
|---|---|---|---|---|
| **#** | **Item Code** | **Package Description** | **Marketing Start Date** | **Marketing End Date** |
| **1** | NDC:41163-617-11 | 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 | 04/30/2012 | |

## 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 |

Revised: 5/2012 SUPERVALU INC.