

EQUALINE - triclosan 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

TRICLOSAN 0.46%

PURPOSE

ANTIBACTERIAL

USES

FOR WASHING TO DECREASE BACTERIA ON THE SKIN.

WARNINGS

FOR EXTERNAL USE ONLY.

WHEN USING THIS PRODUCT

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

STOP USING THIS PRODUCT AND ASK A DOCTOR IF

IRRITATION AND REDNESS DEVELOPS AND LASTS MORE THAN 7 DAYS.

KEEP OUT OF REACH OF CHILDREN

IN CASE OF ACCIDENTAL INGESTION, GET MEDICAL HELP OR CONTACT A POISON CONTROL CENTER IMMEDIATELY.

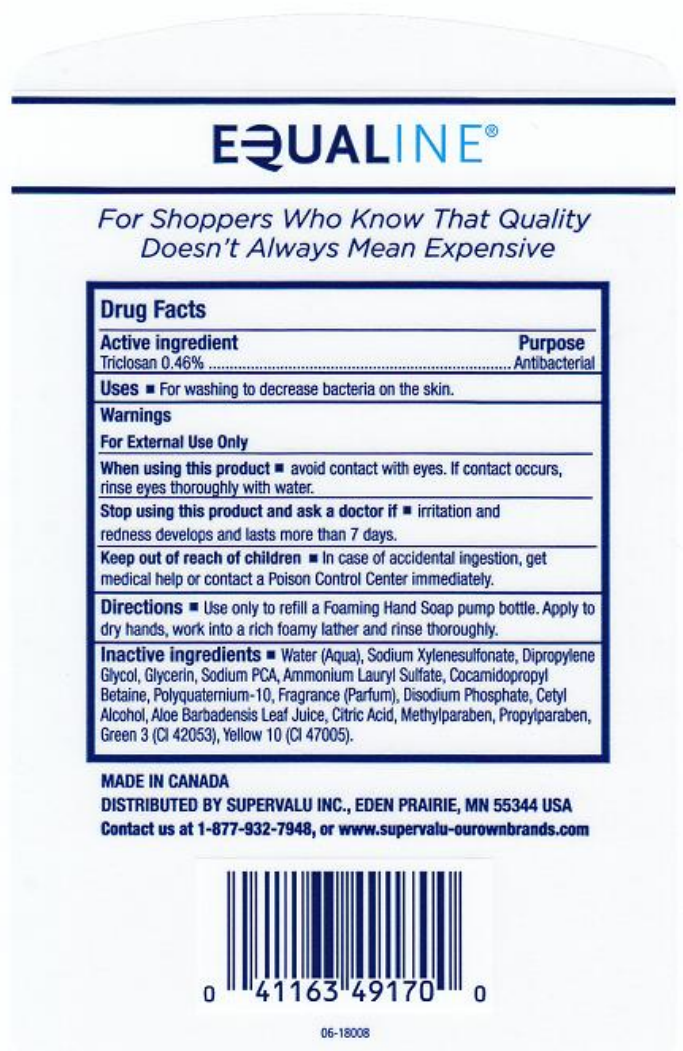
DIRECTIONS

USE ONLY TO REFILL A FOAMING HAND SOAP PUMP BOTTLE. APPLY TO DRY HANDS, WORK INTO A RICH FOAMY LATHER AND RINSE THOROUGHLY.

INACTIVE INGREDIENTS

WATER (AQUA), SODIUM XYLENESULFONATE, DIPROPYLENE GLYCOL, GLYCERIN, SODIUM PCA, AMMONIUM LAURYL SULFATE, COCAMIDOPROPYL BETAINE, POLYQUATERNIUM-10, FRAGRANCE (PARFUM), DISODIUM PHOSPHATE, CETYL ALCOHOL, ALOE BARBADENSIS LEAF JUICE, CITRIC ACID, METHYLPARABEN, PROPYLPARABEN, GREEN 3 (CI 42053), YELLOW 10 (CI 47005).

LABEL COPY



EQUALINE

triclosan liquid

Product Information

| | | | |
|--------------------------------|----------------|---------------------------|---------------|
| Product Type | HUMAN OTC DRUG | Item Code (Source) | NDC:41163-163 |
| Route of Administration | TOPICAL | | |

Active Ingredient/Active Moiety

| Ingredient Name | Basis of Strength | Strength |
|--|-------------------|-------------------|
| TRICLOSAN (UNII: 4NM5039Y5X) (TRICLOSAN - UNII:4NM5039Y5X) | TRICLOSAN | 0.46 mg in 100 mL |

Inactive Ingredients

| Ingredient Name | Strength |
|---|----------|
| WATER (UNII: 059QF0K00R) | |
| SODIUM XYLENESULFONATE (UNII: G4LZF950UR) | |
| DIPROPYLENE GLYCOL (UNII: E107L85C40) | |

| | |
|--|--|
| GLYCERIN (UNII: PDC6A3C00X) | |
| SODIUM PYRROLIDONE CARBOXYLATE (UNII: 469OTG57A2) | |
| AMMONIUM LAURYL SULFATE (UNII: Q7AO2R1M0B) | |
| COCAMIDOPROPYL BETAINE (UNII: 5OCF3O11KX) | |
| POLYQUATERNIUM-10 (400 CPS AT 2%) (UNII: HB1401PQFS) | |
| SODIUM PHOSPHATE (UNII: SE337SVY37) | |
| CETYL ALCOHOL (UNII: 936JST6JCN) | |
| ALOE VERA LEAF (UNII: ZY81Z83H0X) | |
| CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP) | |
| METHYLPARABEN (UNII: A218C7H9T) | |
| PROPYLPARABEN (UNII: Z8IX2SC1OH) | |
| FD&C GREEN NO. 3 (UNII: 3P3ONR6O1S) | |
| D&C YELLOW NO. 10 (UNII: 35SW5USQ3G) | |

Packaging

| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
|---|------------------|-----------------------------|----------------------|--------------------|
| 1 | NDC:41163-163-32 | 946 mL in 1 BOTTLE, PLASTIC | | |

Marketing Information

| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|-------------------------|--|----------------------|--------------------|
| OTC monograph not final | part333E | 07/02/2012 | |

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: 7/2012

SUPERVALU INC.