**WEGMANS ORANGE SCENT - triclosan soap**
**Sun Products Corporation**

*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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**Active ingredient**

Triclosan 0.10%

**Use** helps fight germs on hands when used as a hand soap

**For external use only**

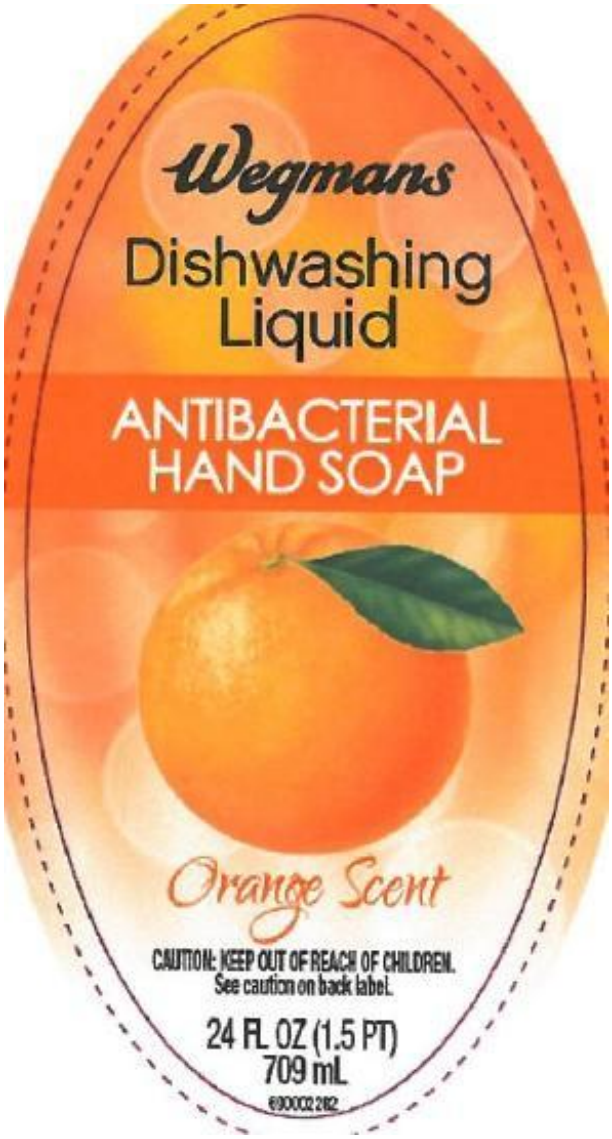
**Keep out of reach of children.** If swallowed, get medical help or contact a Poison Control Center right away. In case of eye contact, rinse thoroughly with water.

**Directions** wash hands and rinse

Antibacterial hand soap

**Inactive ingredients** Water, Urea, Sodium Laureth Sulfate, Sodium Dodecylbenzenesulfonate, Alcohol Denat., Lauramine Oxide, Magnesium Chloride, Sodium Methyl 2-Sulfopalmitate, Fragrance, Sodium Metabisulfite, Disodium 2-Sulfopalmitate, Methyl Palmitate, Tetrasodium EDTA, Methylchloroisothiazolinone, Methylisothiazolinone, Benzophenone-4, FD&C Yellow No. 5, D&C Red No. 33

**Questions?** 1-800-Wegmans (934-6267) Ext 5920 9am-4pm EST workdays


Wegmans
Dishwashing Liquid
ANTIBACTERIAL HAND SOAP
Orange Scent
CAUTION: KEEP OUT OF REACH OF CHILDREN.
See caution on back label.
24 FL OZ (1.5 PT)
709 mL

LBLFR


LBLBK

# WEGMANS ORANGE SCENT

triclosan soap

| Product Information | | | |
|---|---|---|---|
| **Product Type** | HUMAN OTC DRUG | **Item Code (Source)** | NDC:63691-026 |
| **Route of Administration** | TOPICAL | | |

| Active Ingredient/Active Moiety | | |
|---|---|---|
| **Ingredient Name** | **Basis of Strength** | **Strength** |
| **TRICLOSAN** (UNII: 4NM5039Y5X) (TRICLOSAN - UNII:4NM5039Y5X) | TRICLOSAN | 0.1 g in 100 mL |

| Inactive Ingredients | |
|---|---|
| **Ingredient Name** | **Strength** |
| **WATER** (UNII: 059QF0KO0R) | |

| | |
|---|---|
| **UREA** (UNII: 8W8T17847W) | |
| **SODIUM LAURETH SULFATE** (UNII: BPV390UAP0) | |
| **SODIUM DODECYLBENZENESULFONATE** (UNII: 554127163Y) | |
| **ALCOHOL** (UNII: 3K9958V90M) | |
| **LAURAMINE OXIDE** (UNII: 4F6FC4MI8W) | |
| **MAGNESIUM CHLORIDE** (UNII: 02F3473H9O) | |
| **SODIUM METABISULFITE** (UNII: 4VON5FNS3C) | |
| **DISODIUM 2-SULFOPALMITATE** (UNII: VS9295575T) | |
| **METHYL PALMITATE** (UNII: DPY8VCM98I) | |
| **EDETATE SODIUM** (UNII: MP1J8420LU) | |
| **METHYLCHLOROISOTHIAZOLINONE** (UNII: DEL7T5QRPN) | |
| **METHYLISOTHIAZOLINONE** (UNII: 229D0E1QFA) | |
| **SULISOBENZONE** (UNII: 1W6L629B4K) | |
| **FD&C YELLOW NO. 5** (UNII: I753WB2F1M) | |
| **D&C RED NO. 33** (UNII: 9DBA0SBB0L) | |

## Packaging

| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
|---|---|---|---|---|
| **1** | NDC:63691-026-10 | 709 mL in 1 BOTTLE | | |

## Marketing Information

| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|---|---|---|---|
| OTC monograph not final | part333E | 07/01/2014 | |

## Labeler - Sun Products Corporation (070931480)

Revised: 12/2014 Sun Products Corporation