

WEGMANS ULTRA DISHWASHING 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.

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, TEA-Dodecylbenzenesulfonate, Sodium Laureth Sulfate, Sodium Methyl 2-Sulfopalmitate, Lauramine Oxide, Sodium Lauryl Sulfate, Alcohol Denat., Disodium 2-Sulfopalmitate, Methyl Palmitate, Fragrance, Tetrasodium EDTA, Methylchloroisoithiazolinone, Methylisoithiazolinone, Benzophenone-4, Yellow No. 5, Red No. 33

Questions or comments

about this product?

1-800 Wegmans (934-6267) Ext 5920

9am - 4pm EST weekdays

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LBLFR



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WEGMANS ULTRA DISHWASHING ORANGE SCENT

triclosan soap

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:63691-011
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
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WATER (UNII: 059QF0KO0R)	
TRIETHANOLAMINE DODECYLBENZENESULFONATE (UNII: 8HM7ZD48HN)	
SODIUM LAURYL SULFATE (UNII: 368GB5141J)	
SODIUM LAURETH SULFATE (UNII: BPV390UAP0)	
ALCOHOL (UNII: 3K9958V90M)	
LAURAMINE OXIDE (UNII: 4F6FC4M18W)	
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:63691-011-10	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	12/01/2010	

Labeler - Sun Products Corporation (070931480)