UP (AND) UP GREEN APPLE 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, 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, Yellow No. 5, Blue No.1

Questions?call 1-800-910-6874

LBLFR

LBLBK

UP (AND) UP GREEN APPLE SCENT

triclosan soap

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:63691-019
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 DODECYLBENZENESULFONATE (UNII: 554127163Y)	

SODIUM LAURETH SULFATE (UNII: BPV390UAP0)	
ALCOHOL (UNII: 3K9958V90M)	
MAGNESIUM CHLORIDE (UNII: 02F3473H9O)	
LAURAMINE OXIDE (UNII: 4F6FC4MI8W)	
DISODIUM 2-SULFOPALMITATE (UNII: VS9295575T)	
METHYL PALMITATE (UNII: DPY8VCM98I)	
SODIUM METABISULFITE (UNII: 4VON5FNS3C)	
EDETATE SODIUM (UNII: MP1J8420LU)	
METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)	
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	
SULISOBENZONE (UNII: 1W6L629B4K)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:63691-019-10	709 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/01/2011	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	01/01/2011	

Labeler - Sun Products Corporation (070931480)

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