MY ESSENTIALS CITRUS 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, Sodium Dodecylbenzenesulfonate, Palm Kernelamide DEA, Sodium Methyl 2-Sulfopalmitate, Urea, Sodium Chloride, Disodium 2-Sulfopalmitate, Methyl Palmitate, Fragrance, Sodium Citrate, Tetrasodium EDTA, Methylchloroisothiazolinone, Methylisothiazolinone, Benzophenone-4, D&C Orange No. 4

Questions? 1-866-322-2439

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MY ESSENTIALS CITRUS SCENT

triclosan soap

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:63691-022
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)	

SODIUM DODECYLBENZENESULFONATE (UNII: 554127163Y)	
SODIUM PALM KERNELATE (UNII: 6H91L1NXTW)	
UREA (UNII: 8W8T17847W)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
DISODIUM 2-SULFOPALMITATE (UNII: VS9295575T)	
METHYL PALMITATE (UNII: DPY8VCM98I)	
SODIUM CITRATE (UNII: 1Q73Q2JULR)	
EDETATE SODIUM (UNII: MP1J8420LU)	
METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)	
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	
SULISOBENZONE (UNII: 1W6L629B4K)	
D&C ORANGE NO. 4 (UNII: Q1LIY3BO0U)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:63691-022-04	739 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