EXCHANGE SELECT ULTRA 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

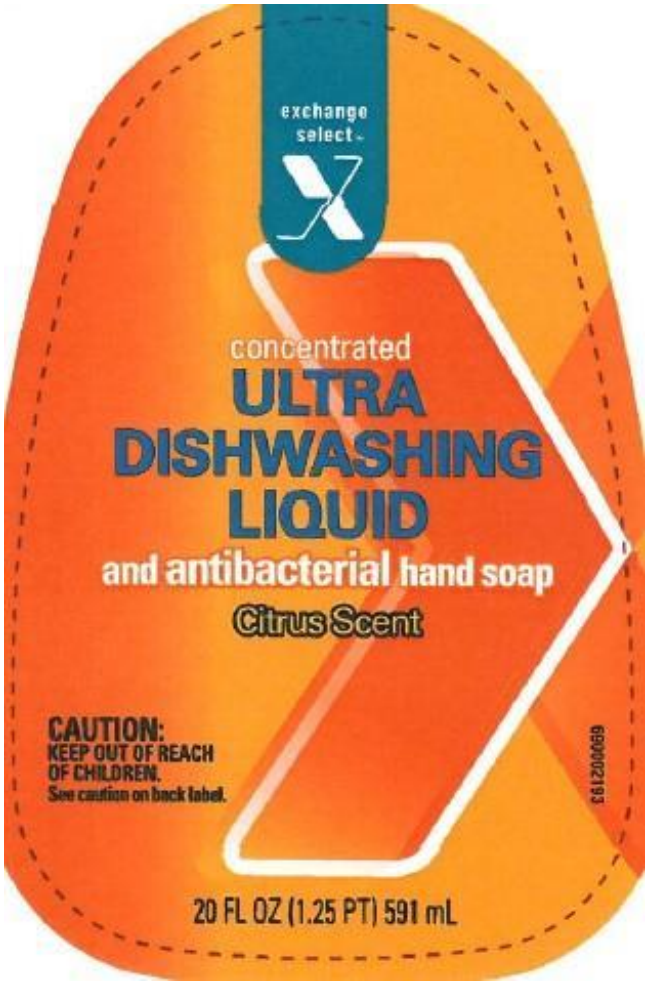
Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away. In case of eye contact, rinse thoroughly with water.

Directions wash hands and rinse

Antibacterial hand soap

Inactive ingredients Water, Sodium Dodecylbenzenesulfonate, Sodium Laureth Sulfate, Sodium Methyl 2-Sulfopalmitate, Magnesium Sulfate, Sodium Xylenesulfonate, Alcohol Denat., Fragrance, Disodium 2-Sulfopalmitate, Methyl Palmitate, Tetrasodium EDTA, Methylchloroisothiazolinone, Methylisothiazolinone, Benzophenone-4, D&C Orange No. 4, Sodium Citrate

Questions? 1-800-776-6702



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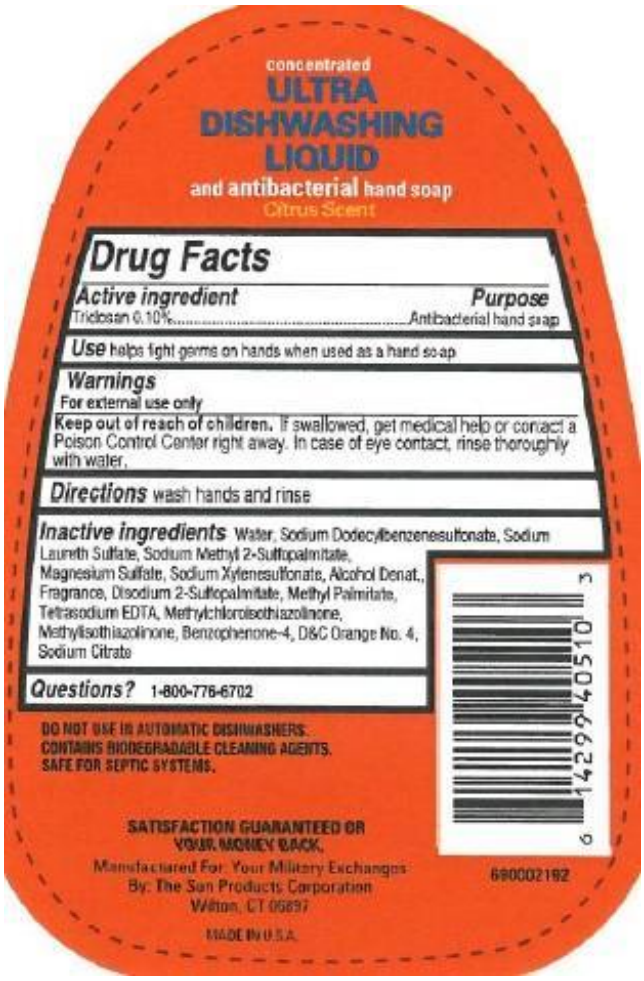

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EXCHANGE SELECT ULTRA CITRUS SCENT

triclosan soap

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:63691-024
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 LAURETH SULFATE (UNII: BPV390UAP0)	
MAGNESIUM SULFATE (UNII: DE08037SAB)	
SODIUM XYLENESULFONATE (UNII: G4LZF950UR)	
ALCOHOL (UNII: 3K9958V90M)	
DISODIUM 2-SULFOPALMITATE (UNII: VS9295575T)	
METHYL PALMITATE (UNII: DPY8VCM98I)	
EDETATE SODIUM (UNII: MP1J8420LU)	
METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)	
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	
SULISOBENZONE (UNII: 1W6L629B4K)	
D&C ORANGE NO. 4 (UNII: Q1LIY3BO0U)	
SODIUM CITRATE (UNII: 1Q73Q2JULR)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:63691-024-12	591 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)

Registrant - Military Exchange (827856670)

Establishment

Name	Address	ID/FEI	Business Operations
Sun Products Corporation - Salt Lake City		096752865	manufacture(63691-024)

Establishment

Name	Address	ID/FEI	Business Operations
Sun Products Corporation - Bowling Green		809709314	manufacture(63691-024)

Revised: 12/2014 Sun Products Corporation