

HOME 360 ULTRA DISH DETERGENT 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

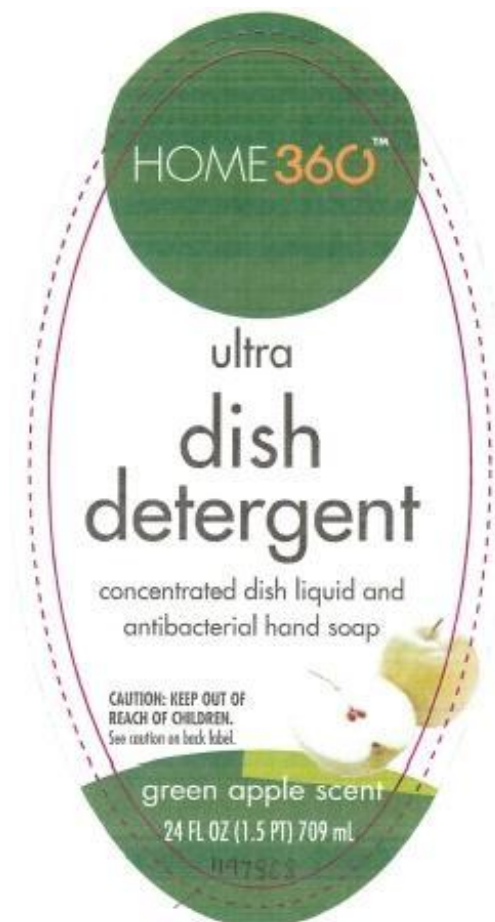
Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away. In case of eye contact, rinse thoroughly with water.

Directions wash hands and rinse

Antibacterial hand soap

Inactive ingredients Water, TEA-Dodecylbenzenesulfonate, Sodium Laureth Sulfate, Sodium Methyl 2-Sulfopalmitate, Lauramine Oxide, Sodium Lauryl Sulfate, Alcohol Denat., Disodium 2-Sulfopalmitate, Methyl Palmitate, Fragrance, Tetrasodium EDTA, Methylchloroisothiazolinone, Methylisothiazolinone, Benzophenone-4, Yellow No. 5, Blue No. 1

Questions? 1-800-776-6702





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HOME 360 ULTRA DISH DETERGENT GREEN APPLE SCENT			
triclosan soap			
Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:63691-013
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)			
TRIETHANOLAMINE DODECYLBENZENESULFONATE (UNII: 8HM7ZD48HN)			
SODIUM LAURYL SULFATE (UNII: 368GB5141J)			
SODIUM LAURETH SULFATE (UNII: BPV390UAP0)			
ALCOHOL (UNII: 3K9958V90M)			

LAURAMINE O XIDE (UNII: 4F6FC4MI8W)	
DISODIUM 2-SULFOPALMITATE (UNII: VS9295575T)	
METHYL PALMITATE (UNII: DPY8VCM98I)	
EDETATE SODIUM (UNII: MP1J8420LU)	
METHYLCHLORO ISOTHIAZOLINONE (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:63691-013-10	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	08/01/2013	

Labeler - Sun Products Corporation (070931480)

Revised: 12/2014

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