REEVA ANTIBACTERIAL GREEN APPLE- 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

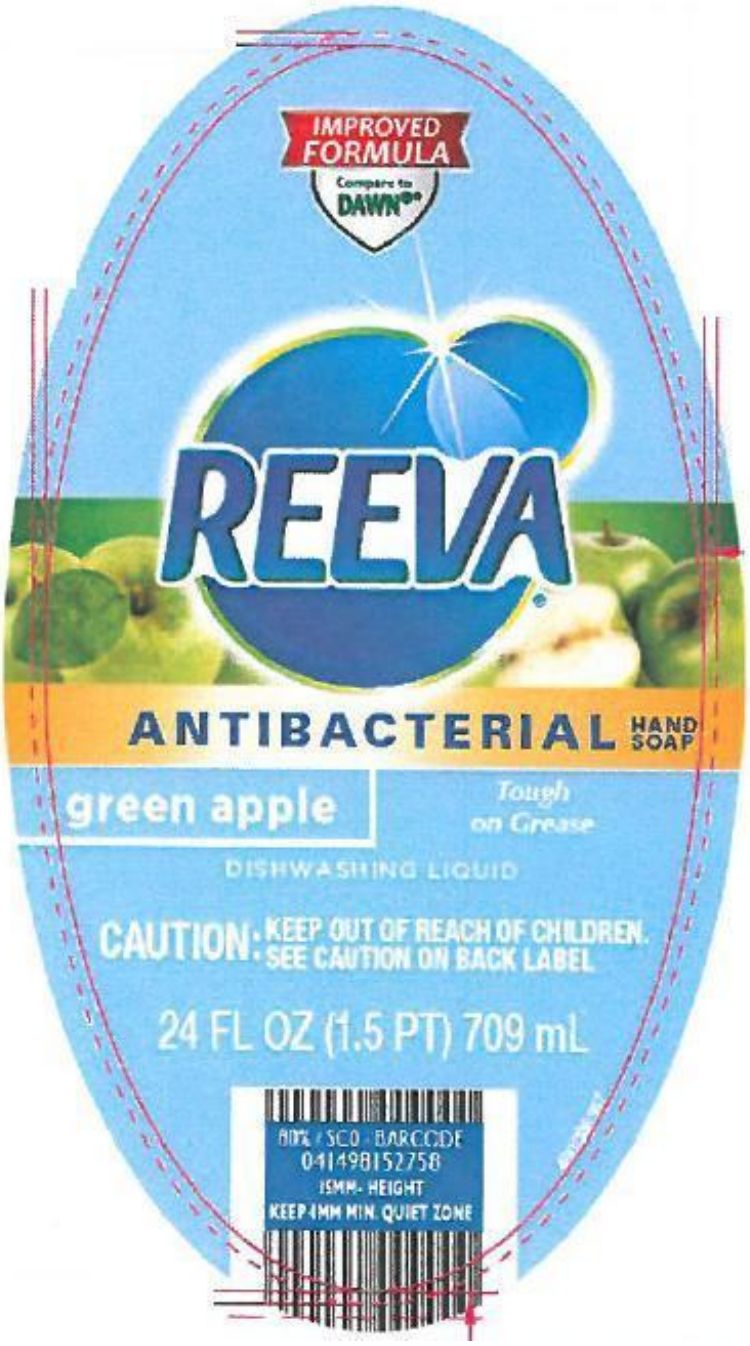
For external use only

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 Lauryl Sulfate, Sodium Laureth Sulfate, Lauramine Oxide, Sodium Dodecylbenzenesulfonate, Sodium Xylene Sulfonate, Alcohol Denat., Fragrance, Sodium Citrate, Methylchloroisothiazolinone, Methylisothiazolinone, Benzophenone-4, Blue No. 1, Yellow No. 5


IMPROVED FORMULA
Compare to DAWN®
REEVA
ANTIBACTERIAL HAND SOAP
green apple
Tough on Grease
DISHWASHING LIQUID
CAUTION: KEEP OUT OF REACH OF CHILDREN.
SEE CAUTION ON BACK LABEL
24 FL OZ (1.5 PT) 709 mL
80% / SCO - BARCODE
041498152758
15MM - HEIGHT
KEEP 4MM MIN. QUIET ZONE

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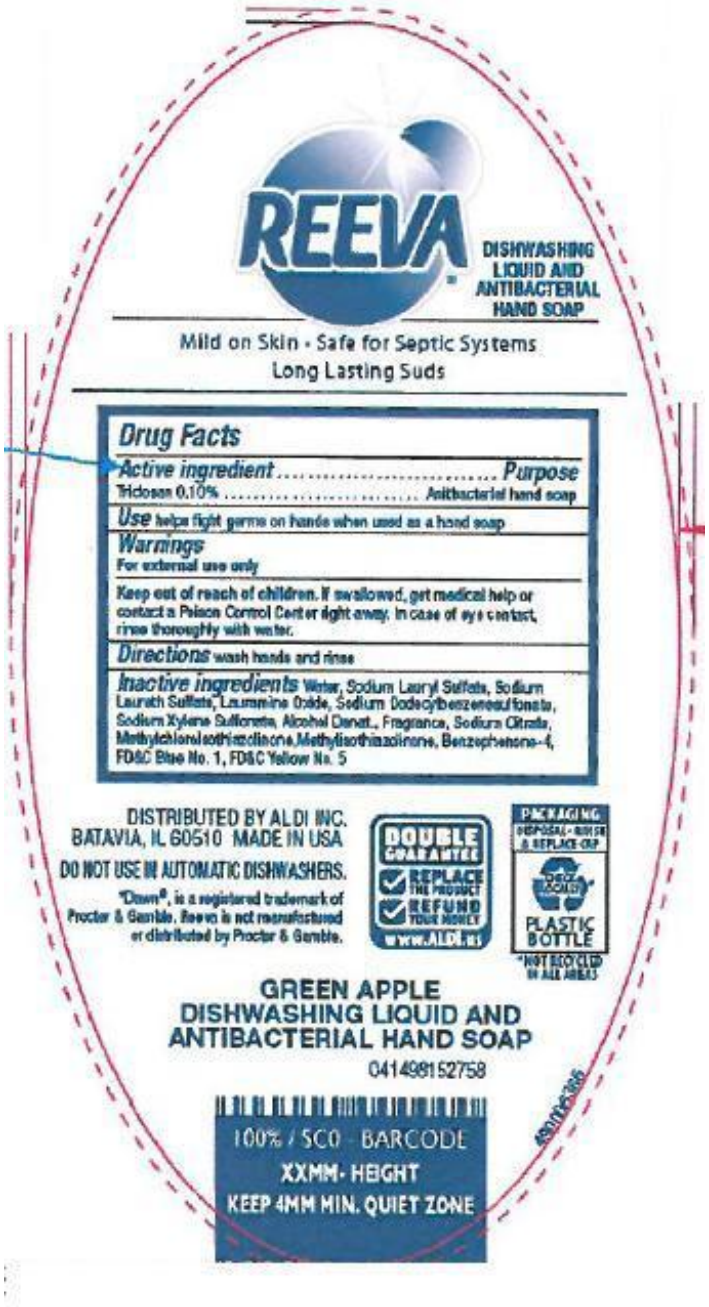

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REEVA ANTIBACTERIAL GREEN APPLE

triclosan soap

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:63691-020
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 LAURYL SULFATE (UNII: 368GB5141J)	
SODIUM LAURETH SULFATE (UNII: BPV390UAP0)	
ALCOHOL (UNII: 3K9958V90M)	
LAURAMINE OXIDE (UNII: 4F6FC4MI8W)	
SODIUM XYLENESULFONATE (UNII: G4LZF950UR)	
SODIUM CITRATE (UNII: 1Q73Q2JULR)	
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:63691-020-10	709 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/01/2014	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	01/01/2014	

Labeler - Sun Products Corporation (070931480)

Revised: 12/2015 Sun Products Corporation