

UP (AND) UP GREEN APPLE SCENT- chloroxyleneol 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

Chloroxyleneol 0.30%

Use

for handwashing to decrease bacteria on the skin

Warnings

For external use only

In case of accidental ingestion, drink a glass of water to dilute. If eye contact occurs, rinse thoroughly with water.

Directions wet hands and forearms. Apply 5 ml or palmful to hands and forearms. Scrub thoroughly for 30 sec. and rinse.

Purpose

Antibacterial hand soap

Inactive ingredients

Water, Sodium Laureth Sulfate, Sodium Dodecylbenzenesulfonate, Sodium Chloride, Alcohol Denat., Sodium Xylenesulfonate, Sodium Lauryl Sulfate, Tetrasodium EDTA, Benzisothiazolinone, Methylisothiazolinone, Methylchlorisothiazolinone, Fragrance, FD&C Yello No. 5, FD&C Blue No. 1

Questions? 1-800-910-6874

Target 60oz GAAB LDU 690004755_690005736-page-001.jpg
Target 24oz GAAB LDU 4900008630_4900008631-page-001.jpg

Compare to
Dawn® Ultra Antibacterial Hand Soap*
green apple
dish soap
and
antibacterial hand soap



24 FL OZ (1.5 PT) 709 mL

CAUTION: KEEP OUT OF REACH OF CHILDREN.
See caution on back label.

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FOR DISHES, POTS AND PANS
One firm squeeze of Up & Up dish soap and antibacterial hand soap into sink, turn on hot water, wash, rinse and drain.

Drug Facts

Active ingredient Purpose
Chloroxylenol 0.30%.....Antibacterial hand soap

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Do not use in automatic dishwashers.
Do not add bleach.

*Dawn® is a registered trademark of The Procter & Gamble Company. This product is not manufactured or distributed by The Procter & Gamble Company.

U We welcome any questions you may have at Target.com/commitments or 1-800-910-6874

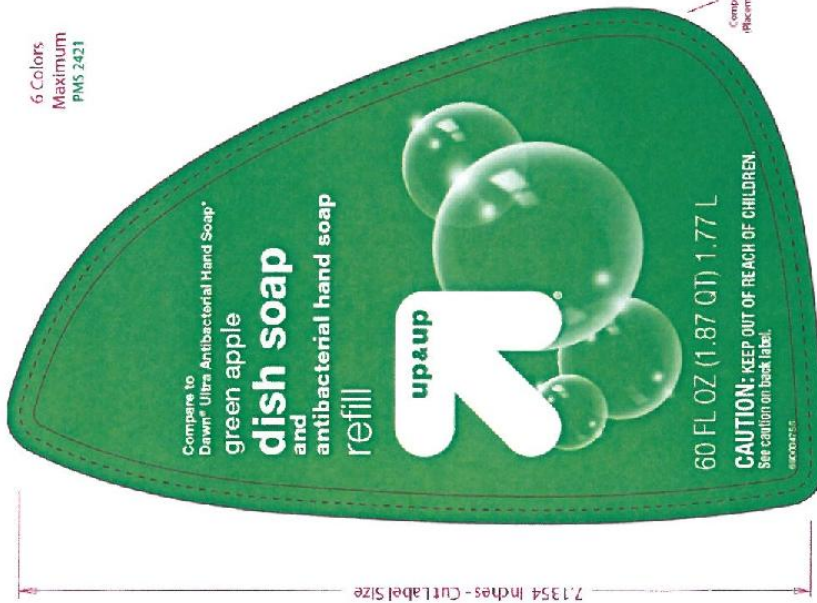
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Amy Finacker
7/14/15

UP (AND) UP GREEN APPLE SCENT



chloroxylenol soap

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:63691-029
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
CHLOROXYLENOL (UNII: 0F32U78V2Q) (CHLOROXYLENOL - UNII:0F32U78V2Q)	CHLOROXYLENOL	0.3 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
SODIUM LAURETH SULFATE (UNII: BPV390UAP0)	
SODIUM XYLENESULFONATE (UNII: G4LZF950UR)	
EDETATE SODIUM (UNII: MP1J8420LU)	
BENZISOTHIAZOLINONE (UNII: HRA0F1A4R3)	
WATER (UNII: 059QF0K00R)	
SODIUM DODECYLBENZENESULFONATE (UNII: 554127163Y)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
ALCOHOL (UNII: 3K9958V90M)	
SODIUM LAURYL SULFATE (UNII: 368GB5141J)	
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	
METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	
FD&C BLUE NO. 1 (UNII: HBR47K3TBD)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:63691-029-30	1774 mL in 1 BOTTLE; Type 0: Not a Combination Product	12/01/2015	
2	NDC:63691-029-10	709 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	04/30/2016	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	11/30/2015	

Labeler - Sun Products Corporation (070931480)