

**SIMPLIFY ANTIBACTERIAL HAND ORANGE SCENT- chloroxylenol soap**  
**Zhejiang Qimei Cosmetics Co., Ltd.**

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**Drug Facts**

**Active Ingredient**

Chloroxylenol 0.30%

**Purpose**

Antibacterial hand soap

**Use**

For handwashing to decrease bacteria on the skin

**Warnings**

**For external use only**

KEEP OUT OF REACH OF CHILDREN.

May cause moderate eye irritation. Avoid contact with eyes and skin. If in eyes, flush eyes gently with water for 15 minutes. If swallowed, drink water to dilute. If irritation persists, contact a Physician or Poison Control Center.

**Directions**

- Wet hands.
- Apply palmful to hands.
- Scrub thoroughly and rinse with water.

**Inactive Ingredients**

Carbamide, citric acid, cocamidopropyl betaine, DMDM hydroton, dodecylbenzene sulfonic acid, ethyl alcohol, fragrance, lauramine oxide, sodium chloride, sodium hydroxide, sodium lauryl ether sulfate, sodium xylene sulfonate, water

**19.4 FL OZ (1.21 PT) 573 mL**

Compare to

Dawn<sup>®</sup> Ultra Antibacterial

Simplify<sup>®</sup>

Antimicrobial

Hand Soap & Ultra Dish

Detergent

- Contains no

phosphates

- Cuts Grease

ORANGE

SCENT

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**Simplify®**

**Antibacterial  
Hand Soap & Ultra Dish  
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the simplify promise:



Ammonia Free



Responsibly Sourced



Phthalate Free

peace of mind  
guarantee

If you are not satisfied,  
we'll happily refund  
your money.

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200 NEWBERRY COMMONS  
ETTERS, PA 17319  
[www.riteaid.com](http://www.riteaid.com)

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or distributed by Procter & Gamble®,  
owner of the registered trademark DAWN®.



MADE IN CHINA

**SIMPLIFY ANTIBACTERIAL HAND ORANGE SCENT**

chloroxylenol soap

Product Information				
<b>Product Type</b>		HUMAN OTC DRUG	<b>Item Code (Source)</b> NDC:81773-014	
<b>Route of Administration</b>		TOPICAL		
Active Ingredient/Active Moiety				
<b>Ingredient Name</b>			<b>Basis of Strength</b>	<b>Strength</b>
CHLOROXYLENOL (UNII: 0F32U78V2Q) (CHLOROXYLENOL - UNII:0F32U78V2Q)			CHLOROXYLENOL	0.3 g in 100 mL
Inactive Ingredients				
<b>Ingredient Name</b>				<b>Strength</b>
DODECYLBENZENESULFONIC ACID (UNII: 60NSK897G9)				
WATER (UNII: 059QF0KO0R)				
UREA (UNII: 8W8T17847W)				
SODIUM C12-14 ALKETH-3 SULFATE (UNII: 5G80BO01PI)				
ALCOHOL (UNII: 3K9958V90M)				
COCAMIDOPROPYL BETAINE (UNII: 5OCF3O11KX)				
SODIUM XYLENESULFONATE (UNII: G4LZF950UR)				
DMDM HYDANTOIN (UNII: BYR0546TOW)				
LAURAMINE OXIDE (UNII: 4F6FC4MI8W)				
SODIUM HYDROXIDE (UNII: 55X04QC32I)				
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)				
SODIUM CHLORIDE (UNII: 451W47IQ8X)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:81773-014-01	573 mL in 1 BOTTLE; Type 0: Not a Combination Product	08/09/2023	
Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC Monograph Drug	505G(a)(3)	08/09/2023		

**Labeler** - Zhejiang Qimei Cosmetics Co., Ltd. (709887693)

Revised: 12/2023

Zhejiang Qimei Cosmetics Co., Ltd.