

## **ANTIMICROBIAL HAND WASH- chloroxylenol soap**

**Henry Schein, Inc.**

*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.*

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### **Drug Facts**

#### **Active Ingredient**

Chloroxylenol (PCMX) 0.6%

#### **Purpose**

Antiseptic

#### **Uses**

- for handwashing to decrease bacteria on skin. Recommended for repeated use.

#### **Warnings**

**For external use only.**

**When using this product** do not get into eyes. If contact occurs, rinse eyes thoroughly with water

#### **Stop use and ask doctor if**

- irritation and redness develops and persists for more than 72 hours

**Keep out of reach of children.** If swallowed, get medical help or contact a Poison Control Center right away

#### **Directions**

- wet hands and forearms
- apply 5 milliliters (teaspoonful) or palmful to hands and forearms
- scrub thoroughly for 30 seconds
- supervise children under 6 years of age

#### **Inactive Ingredients**

aloe vera, cocamide DEA, cocamidopropyl betaine, D&C Red #33, fragrance, germaben II, propylene glycol, purified water, sodium chloride, sodium laureth sulfate, tetrasodium EDTA, vitamin E acetate

Distributed by **HENRY SCHEIN, INC.** Melville, NY 11747 USA

Made in U.S.A.

### **PRINCIPAL DISPLAY PANEL – 27.05 fl. oz. Box Label**

**Henry Schein**

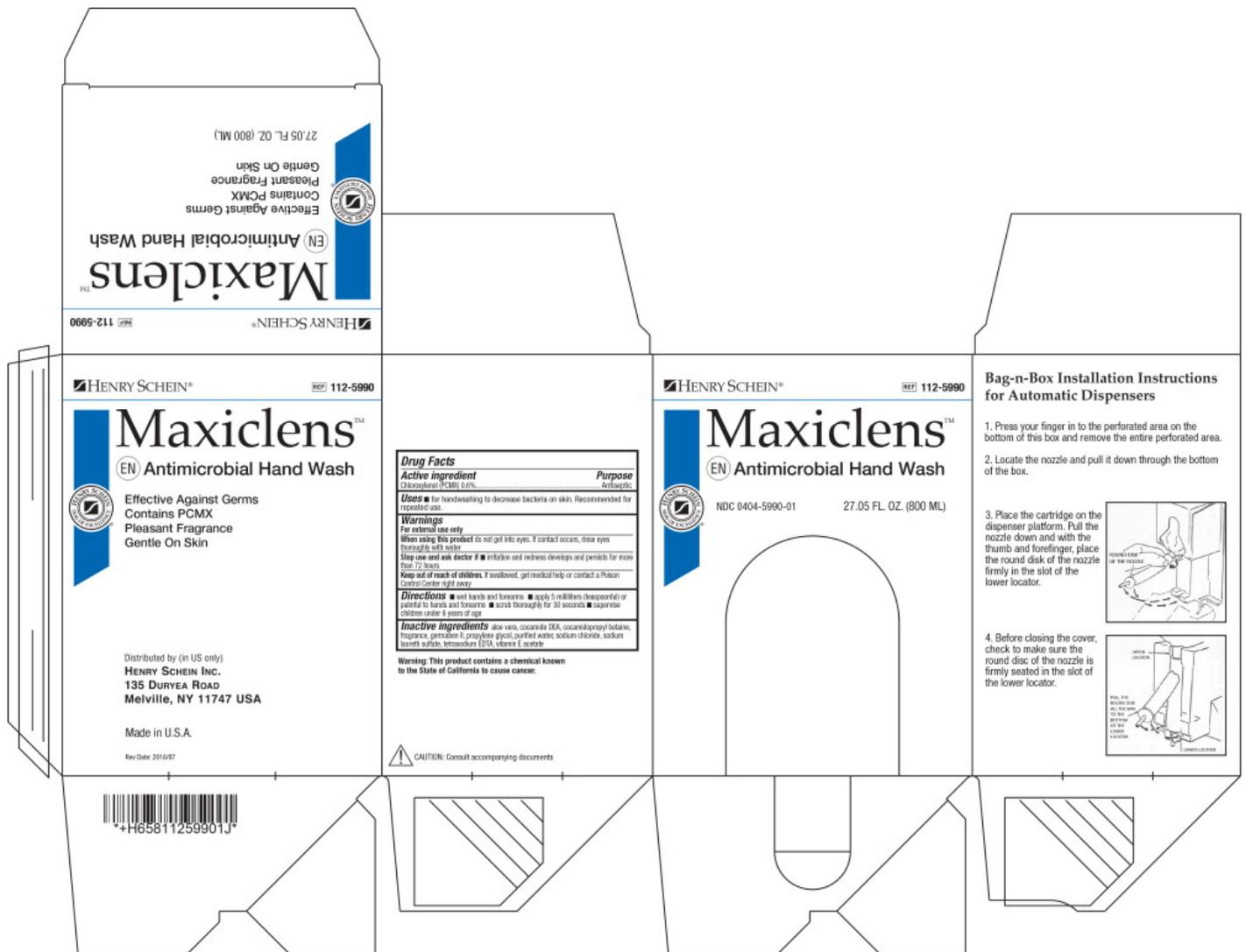
112-5990

**Maxiclens**

**Antimicrobial Hand Wash**

NDC 0404-5990-01

27.05 FL. OZ. (800 ML)



## ANTIMICROBIAL HAND WASH

chloroxylenol soap

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:0404-5990
<b>Route of Administration</b>	TOPICAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
chloroxylenol (UNII: 0F32U78V2Q) (chloroxylenol - UNII:0F32U78V2Q)	chloroxylenol	6.0 mg in 1000 mL

### Inactive Ingredients

Ingredient Name	Strength
aloe vera leaf (UNII: ZY81Z83H0X)	
coco diethanolamide (UNII: 92005F972D)	
cocamidopropyl betaine (UNII: 5OCF3O11KX)	
diazolidinyl urea (UNII: H5RIZ3MPW4)	
propylene glycol (UNII: 6DC9Q167V3)	
water (UNII: 059QF0KO0R)	
sodium chloride (UNII: 451W47IQ8X)	
sodium laureth sulfate (UNII: BPV390UAP0)	
edetate sodium (UNII: MPLJ8420LU)	
alpha-tocopherol acetate (UNII: 9E8X80D2L0)	
propylparaben (UNII: Z8IX2SC1OH)	
methylparaben (UNII: A2I8C7HI9T)	

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0404-5990-01	800 mL in 1 BOX; Type 0: Not a Combination Product	03/01/2012	

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	03/01/2012	

**Labeler** - Henry Schein, Inc. (012430880)

**Registrant** - Safetec of America, Inc. (874965262)

### Establishment

Name	Address	ID/FEI	Business Operations
Safetec of America, Inc.		874965262	MANUFACTURE(0404-5990)

Revised: 7/2016

Henry Schein, Inc.