**HANDI-KLEEN ANTIBACTERIAL FOAMING HAND WASH- chloroxylenol liquid**
**Diamond Chemical Co. 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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**Handi-Kleen Antibacterial Foaming Hand Wash**

***Drug Facts***

***Active Ingredient***

Cloroxylenol 0.3% w/w

***Purpose***

Antiseptic

***Uses***

- Handwash to help reduce bacteria on the skin that potentially can cause disease.
- Recommended for repeated use.

***Warnings***

**For external use only.**

**When using this product** keep out of eyes. In case of eye contact, flush eyes with water.

**Stop use and ask a doctor if** irritation or redness develop or if condition 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 with water and dispense sufficient amount of product into cupped palm of hand.
- Wash both hands thoroughly for 15 minutes.
- Rinse under running water and dry thoroughly.

***Inactive Ingredients*** Water, Sodium Laureth Sulfate, Cocamide DEA, DMDM Hydantoin, Ethyl Alcohol, Phenoxyethanol, Isopropyl Alcohol, Fragrance, Citric Acid, FD&C Red 4

***Handi-Kleen™***

**Antibacterial Foaming Hand Wash**

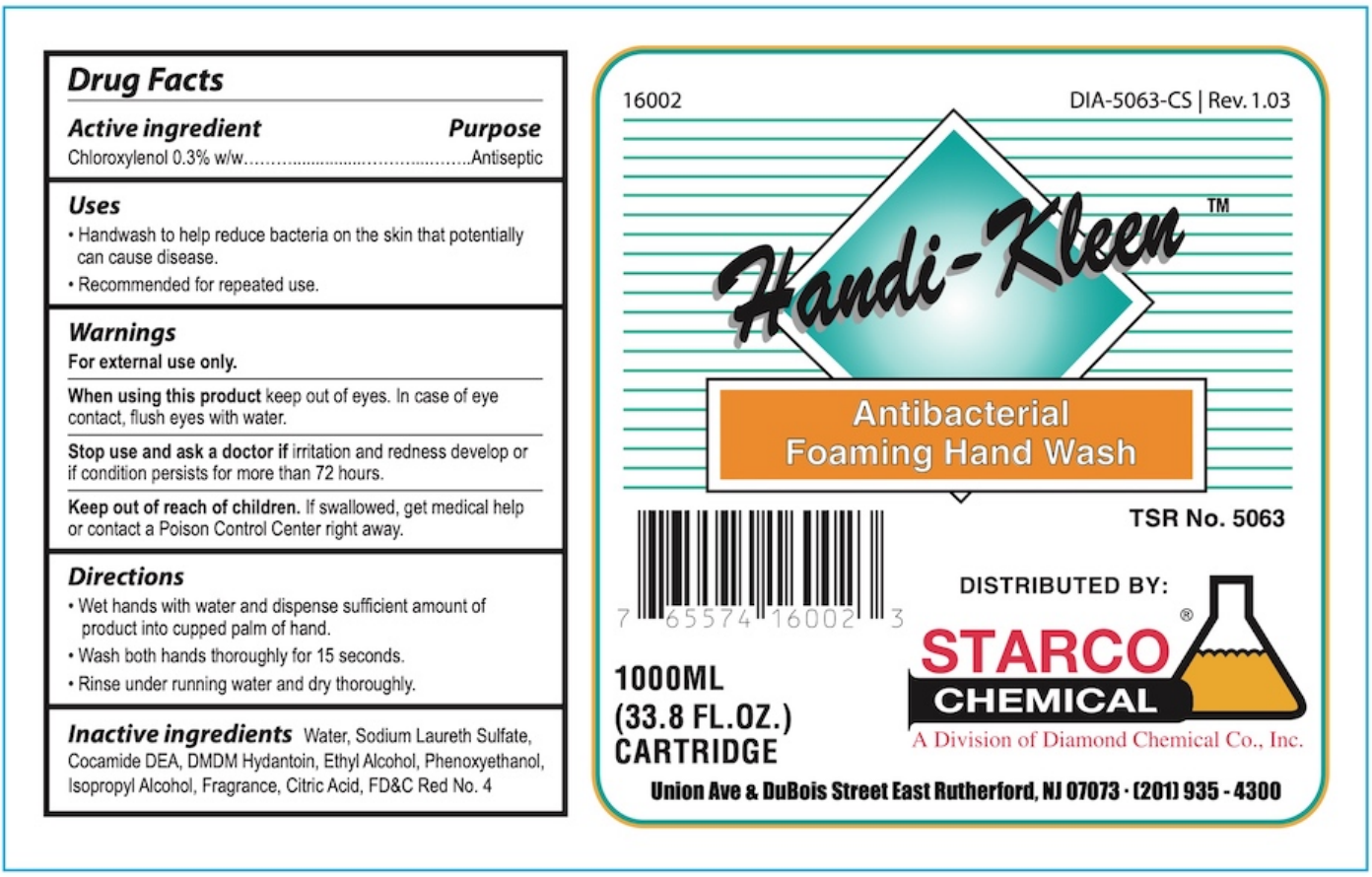
**1000 ML (33.8 FL.OZ.) CARTRIDGE**

DISTRIBUTED BY:

**STARCO CHEMICAL**

A Division of Diamond Chemical Co., Inc.

**Union Ave & Dubois Street East Rutherford, NJ 07073 • (201) 935-4300**




# HANDI-KLEEN ANTIBACTERIAL FOAMING HAND WASH

chloroxylenol liquid

| Product Information | | | |
|---|---|---|---|
| **Product Type** | HUMAN OTC DRUG | **Item Code (Source)** | NDC:72177-180 |
| **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** |
| **CITRIC ACID MONOHYDRATE** (UNII: 2968PHW8QP) | |
| **COCAMIDOPROPYL DIMETHYLAMINE** (UNII: L36BM7DG2T) | |
| **DMDM HYDANTOIN** (UNII: BYR0546TOW) | |
| **ALCOHOL** (UNII: 3K9958V90M) | |
| **ISOPROPYL ALCOHOL** (UNII: ND2M416302) | |
| **PHENOXYETHANOL** (UNII: HIE492ZZ3T) | |

| | |
|---|---|
| **FD&C RED NO. 4** (UNII: X3W0AM1JLX) | |
| **SODIUM LAURETH SULFATE** (UNII: BPV390UAP0) | |
| **WATER** (UNII: 059QF0KO0R) | |

## Packaging

| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
|---|---|---|---|---|
| **1** | NDC:72177-180-10 | 1000 mL in 1 CARTRIDGE; Type 0: Not a Combination Product | 02/03/2021 | 07/26/2024 |

## Marketing Information

| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|---|---|---|---|
| OTC monograph not final | part333E | 02/03/2021 | 07/26/2024 |

## Labeler - Diamond Chemical Co. Inc. (001381482)

Revised: 9/2022 Diamond Chemical Co. Inc.