**DIAL PROFESSIONAL GEL SANITIZER- dial professional gel hand sanitizer gel**
**Henkel Corporation**

---

**Dial Professional Gel Hand Sanitizer**
**Dial Professional Hand Sanitizer Gel 11 Fl Oz (12)**
**Dial Professional Antibacterial Hand Sanitizer Gel**

**Dial Hand Sanitzer (Gel/Foaming)**

**Dial Professional Gel Hand Sanitizer**

Active Ingredient: Ethyl Alcohol 62.0% v/v Purpose: Antiseptic

## Warnings

Warnings For external use only
Flammable. Keep away from fire or flame.

When using this product•Avoid contact with face, eyes, and broken skin. •If eye contact occurs, flush thoroughly with water and seek medical advice.ORWhen using this productavoid contact with face, eyes, and broken skin.If eye contact occurs, flush thoroughly with water and seek medical advice.
Stop use and ask a doctor if•Irritation or redness developsORStop use and ask a doctor ifirritation or redness develops
Keep out of reach of children. •If swallowed, get medical help or contact a Poison Control Center right away

Use(s)Hand sanitizer to help reduce bacteria thatpotentially may cause disease

Use(s)Hand sanitizer to help reduce bacteria that potentially may cause disease

Inactive ingredients Aqua (Water, Eau) · Propylene Glycol ·

Glycerin . Carbomer . Aminomethyl Propanol . Isopropyl Myristate .
T-Butyl Alcohol

**Dial Gel/Foaming Hand Sanitizer**


# DIAL PROFESSIONAL GEL SANITIZER

dial professional gel hand sanitizer gel

| Product Information | | | |
|---|---|---|---|
| **Product Type** | HUMAN OTC DRUG | **Item Code (Source)** | NDC:54340-136 |
| **Route of Administration** | TOPICAL | | |

| Active Ingredient/Active Moiety | | |
|---|---|---|
| **Ingredient Name** | **Basis of Strength** | **Strength** |
| **ALCOHOL** (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M) | ALCOHOL | 60.14 mL in 100 mL |

| Inactive Ingredients | |
|---|---|
| **Ingredient Name** | **Strength** |

| | |
|---|---|
| **WATER** (UNII: 059QF0KO0R) | 38.19 mL in 100 mL |

## Packaging

| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
|---|---|---|---|---|
| **1** | NDC:54340-136-02 | 3 in 1 CARTON | 06/21/2022 | 01/01/2024 |
| **1** | NDC:54340-136-01 | 1200 mL in 1 BOTTLE, DISPENSING; Type 0: Not a Combination Product | | |
| **2** | NDC:54340-136-04 | 12 in 1 CARTON | 06/21/2022 | 01/01/2024 |
| **2** | NDC:54340-136-03 | 221 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product | | |
| **3** | NDC:54340-136-06 | 6 in 1 CARTON | 06/21/2022 | 01/01/2024 |
| **3** | NDC:54340-136-05 | 443 mL in 1 POUCH; Type 0: Not a Combination Product | | |
| **4** | NDC:54340-136-08 | 3 in 1 CARTON | 06/21/2022 | 01/01/2024 |
| **4** | NDC:54340-136-07 | 1000 mL in 1 BOTTLE, DISPENSING; Type 0: Not a Combination Product | | |
| **5** | NDC:54340-136-10 | 144 in 1 CARTON | 06/21/2022 | 01/01/2024 |
| **5** | NDC:54340-136-09 | 59 mL in 1 BOTTLE, DISPENSING; Type 0: Not a Combination Product | | |
| **6** | NDC:54340-136-11 | 12 in 1 CARTON | 01/01/2024 | |
| **6** | | 325 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product | | |

## Marketing Information

| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|---|---|---|---|
| OTC Monograph Drug | 505G(a)(3) | 06/21/2022 | |

## Labeler - Henkel Corporation (080887708)

Revised: 12/2025 Henkel Corporation