DIAL PROFESSIONAL ANTIBACTERIAL HAND SANITIZER FOAM- dial professional antibacterial hand sanitizer foam gel Henkel Corporation

Dial Professional Antibacterial Gel/Foaming Hand Sanitizer

Dial Professional Antibacterial Hand Sanitizer Gel

Dial Professional Foaming Hand Sanitizer - 4/15.2oz Pump

Dial Professional Foaming Hand Sanitizer FIT Universal Touch Free - Refill 3/1L

Dial Professional Hand Sanitizer Foam FIT X1 Touch Free - 3/1L

Dial Professional Foaming Hand Sanitizer FIT X2 Touch Free - Refill 3/1L

Dial Professional Foam Hand Santizer FIT Universal Manual - Refill 3/1.2L

Dial Professional Hand Santizer Foam FIT X1 Manual - Refill 3/1.2L

Dial Professional Foam Hand Santizer FIT X2 Manual - Refill 3/1.2L

Dial Professional Foam Hand Sanitizer Dial 1700 Manual Refill - 3/1.2L

Dial Hand Sanitzer (Gel/Foaming)

Dial Professional Gel Hand Sanitizer

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

Inactive Ingredients: Aqua (Water, Eau) · Propylene Glycol · Glycerin · Carbomer ·

Aminomethyl Propanol · Isopropyl Myristate · T-Butyl Alcohol

Dial Professional Foaming Hand Sanitizer

Active Ingredient: Ethyl Alcohol 65% v/v Purpose: Antiseptic

Inactive Ingredients: Agua (Water, Eau) · PEG-10 Dimethicone · Glycerin ·

Cocamidopropyl PG-Dimonium Chloride · Isopropyl Myristate · PEG-12 Allyl Ether · T-

Butyl Alcohol · Diazolidinyl Urea · Methylparaben

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 develops ORStop use and ask a doctor if irritation 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 that potentially may cause disease

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Inactive Ingredients:

Aqua

PEG-10 Dimethicone

Glycerin

Cocamidopropyl PG-Dimonium Chloride

Isopropyl Myristate

PEG-12 Allyl Ether

T-Butyl Alcohol

Diazolidinyl Urea

Methylparaben

Dial Gel/Foaming Hand Sanitizer



ANTIBACTERIAL

FRAGRANCE FREE

Drug Facts

Active ingredient Purpose Ethyl Alcohol 62.0%......Antiseptic

Uses

· hand sanitizer to help reduce bacteria that potentially may cause disease

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.

Stop use and ask a doctor if

irritation or redness develops I

Drug Facts (continued)

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

Directions

- · wet hands thoroughly with product and rub into skin until dry
- · Children under 6 years of age should be supervised by an adult when using this product.

Inactive ingredients

water, glycerin, isopropyl myristate, propylene glycol, carbomer, aminomethyl propanol

Questions? 1-877-777-3277

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www.dialprofessional.com

800 mL (27 FI Oz) 1 Pt 11 FI Oz



KILLS 99.999% OF BACTERIA!



latex, vinyl, and



CM-5-95862-04/1716833









DIAL PROFESSIONAL ANTIBACTERIAL HAND SANITIZER FOAM

dial professional antibacterial hand sanitizer foam gel

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:54340-130	
Route of Administration	TOPICAL			

Active Ingredient/Active Moiety				
Ingredient Name	Basis of Strength	Strength		
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	62.75 mL in 100 mL		

Inactive Ingredients				
Ingredient Name	Strength			
WATER (UNII: 059QF0KO0R)	33 mL in 100 mL			

P	Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:54340- 130-04	3 in 1 CARTON	06/21/2022		
1	NDC:54340- 130-03	1200 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product			
2	NDC:54340- 130-02	3 in 1 CARTON	06/21/2022		
2	NDC:54340- 130-01	1000 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product			
3	NDC:54340- 130-06	4 in 1 CARTON	06/21/2022		
3	NDC:54340- 130-05	444 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: 12/2024 Henkel Corporation