DIAL PROFESSIONAL ANTIBACTERIAL HAND SANITIZER - ethyl alcohol liquid
The Dial Corporation

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.

Drug Facts

Active ingredient

Ethyl Alcohol 72%

Purpose

Antibacterial

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 and redness develops

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

Directions

pump foam into dry hands, wet thoroughly 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, n-propanol, bis-PEG-12 dimethicone, behentrimonium chloride, PEG-200 hydrogenated glyceryl palmate, PEG-7 glyceryl cocoate, coco-glucoside, glyceryl linoleammonium chloride

Questions?

1-877-777-3277

Dial Professional

Antibacterial Hand Sanitizer Fragrance Free Foam

Kills 99.99% of germs instantly

Made in Canada

2010 Distributed by The Dial Corporation/A Henkel Company

Scottsdale, AZ 85255

www.dialprofessional.com

Install with label toward wall

Instalar con etiqueta hacia la pared

US Patent Numbers

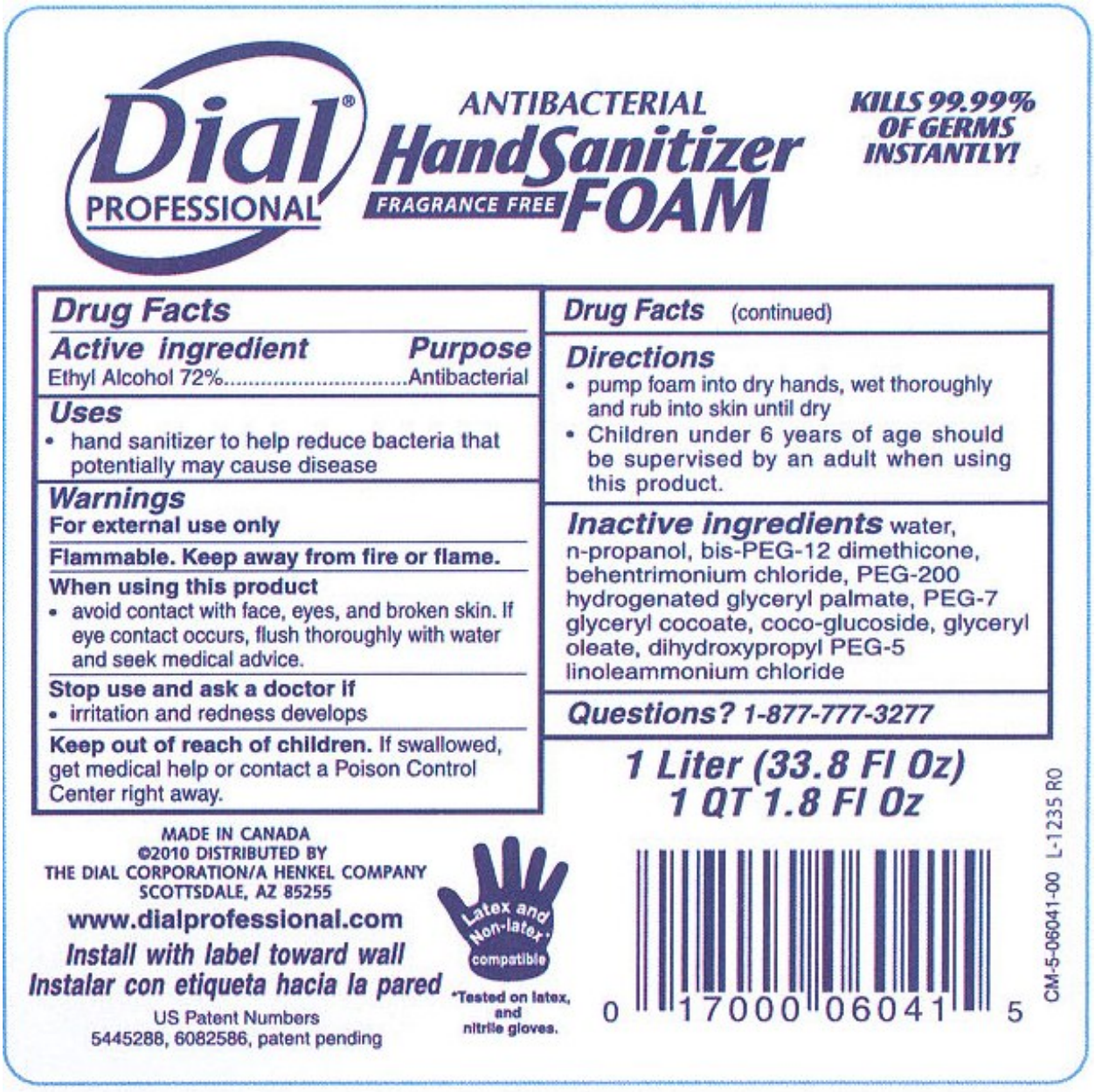
5445288, 6082586, patent pending

1 Liter (33.8 Fl Oz)

1 QT 1.8 Fl Oz

Latex and Non-latex compatible

Tested on latex and nitrile gloves.



DIAL PROFESSIONAL ANTIBACTERIAL HAND SANITIZER

ethyl alcohol liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:54340-107
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
Alcohol (UNII: 3K9958V90M) (Alcohol - UNII:3K9958V90M)	Alcohol	72 mL in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
DIHYDROXYPROPYL PEG-5 LINOLEAMMONIUM CHLORIDE (UNII: 0Y0NQR2GH1)	
PROPYL ALCOHOL (UNII: 96F264O9SV)	
BEHENTRIMONIUM CHLORIDE (UNII: X7GNG3S47T)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:54340-107-27	1000 mL in 1 BOTTLE, PLASTIC		
2	NDC:54340-107-12	1000 mL in 1 BOTTLE, PLASTIC		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	12/01/2010	

Labeler - The Dial Corporation (070252531)

Establishment

Name	Address	ID/FEI	Business Operations
Deb Worldwide Healthcare Inc.		205662831	manufacture

Revised: 3/2011 The Dial Corporation