PROFORMANCE SERIES FOAM ALCOHOL HAND SANITIZER - ethyl alcohol liquid
Pro-Link, 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.

Drug Facts

Active ingredient

Ethyl Alcohol 72%

Purpose

Antibacterial

Uses

For hand sanitizing to reduce bacteria on skin

Warnings

For external use only

Flammable: Keep away from fire or flame.

When using this product avoid contact with eyes.

In case of eye contact, flush with water.

Keep out of reach of children.

If swallowed, get medical help or contact a Poison Control Center right away.

Directions

Apply one shot of foaming sanitizer to dry hands

Rub into skin

No rinsing required

Inactive ingredients

Water, n-Propanol, Bis-PEG-12 Dimethicone, Behentrimonium Chloride, PEG-200 Hydrogenated Glyceryl Palmate, PEG-7 Glyceryl Cocoate, Coco-Glucoside, Glyceryl Oleate, Dihydroxypropyl PEG-5 Linoleammonim Chloride

ProFormance Series

Foam Alcohol Sanitizer

Pro-Link Green

Certified EcoLogo

Certified Instant Hand Antiseptic CCD-170

MSA1000

Distributed exclusively by:

Pro-Link, Inc.

1 Liter

33.8 Fluid Ounces

PROFORMANCE SERIES FOAM ALCOHOL HAND SANITIZER

ethyl alcohol liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:66908-107
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:66908-107-27	1000 mL in 1 BOTTLE, PLASTIC		
2	NDC:66908-107-12	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	01/01/2011	

Labeler - Pro-Link, Inc. (144650637)

Registrant - Deb USA, Inc. (607378015)

Establishment

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

Revised: 1/2011 Pro-Link, Inc.