

**PROFORMANCE SERIES FOAM NON-ALCOHOL SANITIZER - benzalkonium
chloride 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

Benzalkonium Chloride, 0.13%

Purpose

Antibacterial

Uses

For hand sanitizing to reduce bacteria on the skin

Warnings

For external use only

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 to dry hands, rub into skin

No rinsing required

Water, Propylene Glycol, Aloe Barbadosis Leaf Juice, Cocamidopropyl Betaine, Lauramine Oxide, Tetrasodium EDTA, Magnesium Nitrate, Methylchloroisothiazolinone, Magnesium Chloride, Methylisothiazolinone.

ProFormance Series

Foam Non-Alcohol Sanitizer

Pro-Link

MSN1000

Distributed exclusively by

Pro-Link, Inc.

1 Liter

33.8 Fluid Ounces



Foam Non-Alcohol Sanitizer

Drug Facts

Active ingredient	Purpose
Benzalkonium Chloride, 0.13%	Antibacterial

Uses ■ For hand sanitizing to reduce bacteria on the skin

Warnings

For external use only

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 to dry hands, rub into skin
- No rinsing required

Inactive ingredients

Water, Propylene Glycol, Aloe Barbadosensis Leaf Juice, Cocamidopropyl Betaine, Lauramine Oxide, Tetrasodium EDTA, Magnesium Nitrate, Methylchloroisothiazolinone, Magnesium Chloride, Methylisothiazolinone.

MSN1000

Distributed exclusively by:
Pro-Link, Inc.
Canton, MA 02021
Ph.: 800-74-LINKS
www.prolinkhq.com



Rev. 12-10

MSN100-01-116



7 23622 13302 3

1 Liter • 33.8 Fluid Ounces

PROFORMANCE SERIES FOAM NON-ALCOHOL SANITIZER

benzalkonium chloride liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:66908-106
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
Benzalkonium Chloride (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	Benzalkonium Chloride	0.13 mL in 100 mL

Inactive Ingredients

Ingredient Name	Strength
-----------------	----------

WATER (UNII: 059QF0KO0R)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
COCAMIDOPROPYL BETAINE (UNII: 5OCF3O11KX)	
LAURAMINE OXIDE (UNII: 4F6FC4M18W)	
EDETATE SODIUM (UNII: MP1J8420LU)	
MAGNESIUM NITRATE (UNII: 77CBG3UN78)	
METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)	
MAGNESIUM CHLORIDE (UNII: 02F3473H9O)	
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:66908-106-27	1000 mL in 1 BOTTLE, PLASTIC		
2	NDC:66908-106-12	1200 mL in 1 BOTTLE, PLASTIC		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	01/01/2011	

Labeler - Pro-Link, Inc. (144650637)

Registrant - Deb USA, Inc. (607378015)

Establishment

Name	Address	ID/FEI	Business Operations
Deb USA, Inc.		607378015	manufacture