

**INSTANTFOAM NON-ALCOHOL - benzalkonium chloride liquid**  
**PARAGON AERO INSTANT - benzalkonium chloride liquid**  
**HY5 SOAPOPOPULAR AERO - benzalkonium chloride liquid**  
**Deb USA, 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.*

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**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

Inactive ingredients

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

deb

InstantFOAM

non-alcohol-dye and fragrance free

hand sanitizer

refreshing

no water required

use anywhere, anytime

use everyday

Kills 99.99% of common germs

deb foam technology

NSF

Nonfood Compounds Program Listed E-3 140059

55854-01-116

1 Liter - 33.8 Fluid Ounces



# INSTANTFOAM NON-ALCOHOL

benzalkonium chloride liquid

## Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:11084-058
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: 4F6FC4MI8W)	
EDETATE SODIUM (UNII: MPIJ8420LU)	
METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)	
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	
MAGNESIUM NITRATE (UNII: 77CBG3UN78)	
MAGNESIUM CHLORIDE (UNII: 02F3473H9O)	

## Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11084-058-01	47 mL in 1 BOTTLE, PUMP		
2	NDC:11084-058-05	3780 mL in 1 BOTTLE, PLASTIC		
3	NDC:11084-058-12	1200 mL in 1 BOTTLE, PLASTIC		
4	NDC:11084-058-20	2000 mL in 1 BOTTLE, PLASTIC		
5	NDC:11084-058-27	1000 mL in 1 BOTTLE, PLASTIC		
6	NDC:11084-058-40	400 mL in 1 BOTTLE, PUMP		

## Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	01/01/2010	

# PARAGON AERO INSTANT

benzalkonium chloride liquid

**Product Information**

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:11084-137
<b>Route of Administration</b>	TOPICAL		

**Active Ingredient/Active Moiety**

<b>Ingredient Name</b>	<b>Basis of Strength</b>	<b>Strength</b>
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.13 mL in 100 mL

**Inactive Ingredients**

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

**Packaging**

#	<b>Item Code</b>	<b>Package Description</b>	<b>Marketing Start Date</b>	<b>Marketing End Date</b>
1	NDC:11084-137-27	1000 mL in 1 BOTTLE, PLASTIC		

**Marketing Information**

<b>Marketing Category</b>	<b>Application Number or Monograph Citation</b>	<b>Marketing Start Date</b>	<b>Marketing End Date</b>
OTC mono graph not final	part333A	01/01/2010	

**HY5 SOAPOPULAR AERO**

benzalkonium chloride liquid

**Product Information**

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:11084-136
<b>Route of Administration</b>	TOPICAL		

**Active Ingredient/Active Moiety**

<b>Ingredient Name</b>	<b>Basis of Strength</b>	<b>Strength</b>
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM -	BENZALKONIUM	0.13 mL

UNII:7N6JUD5X6Y)	CHLORIDE	in 100 mL		
<b>Inactive Ingredients</b>				
<b>Ingredient Name</b>		<b>Strength</b>		
WATER (UNII: 059QF0KO0R)				
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)				
ALOE VERA LEAF (UNII: ZY81Z83H0X)				
COCAMIDO PROPYL BETAINE (UNII: 5OCF3O11KX)				
LAURAMINE OXIDE (UNII: 4F6FC4M18W)				
EDETATE SODIUM (UNII: MP1J8420LU)				
METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)				
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)				
MAGNESIUM NITRATE (UNII: 77CBG3UN78)				
MAGNESIUM CHLORIDE (UNII: 02F3473H9O)				
<b>Packaging</b>				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11084-136-27	1000 mL in 1 BOTTLE, PLASTIC		
<b>Marketing Information</b>				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph not final	part333A	01/01/2010		

**Labeler** - Deb USA, Inc. (607378015)

### Establishment

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

Revised: 9/2010

Deb USA, Inc.