ANTIBACTERIAL FOAMING HAND SANITIZER- benzalkonium chloride solution
Jets, Sets, & Elephants Beauty Corp.

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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bath+body+etc® antibacterial foaming hand sanitizer - Warm Vanilla Sugar, Sweet Cherry Blossom, Fresh Ocean Breeze, Cool Cucumber Melon, Irresistible Green Apple, Alluring Red Pomegranate, Sparkling White Cranberry, Crisp White Citrus, Sheer Cotton Whisper

Active ingredient

Benzalkonium chloride 0.11%

Purpose

Antibacterial

Uses

- to decrease bacteria on the skin and clean hands
- recommended for repeated use

Warnings

For external use only. Flammable, keep away from fire or flame.

Keep out of reach of children.

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

Do not get into eyes. If contact occurs, rinse thoroughly with water.

Discontinue use if irritation or redness develop. If irritation persists for more than 72 hours, consult a doctor.

Directions

- apply to hands until thoroughly wet
- rub vigorously until dry
- supervise children in the use of this product

Inactive Ingredients

Warm Vanilla Sugar

Aqua, Alcohol, Decyl Glucoside, Parfum, Aloe Barbadensis, Methylisothiazolinone, Methylchloroisothiazolinone, Disodium EDTA, Benzophenone-4, Citric Acid, Tocopheryl Acetate, Retinyl Palmitate, Red 33, Blue 1, Yellow 5

Sweet Cherry Blossom

Aqua, Alcohol, Decyl Glucoside, Parfum, Aloe Barbadensis, Methylisothiazolinone, Methylchloroisothiazolinone, Disodium EDTA, Benzophenone-4, Citric Acid, Tocopheryl Acetate, Retinyl Palmitate, Red 33, Yellow 5

Fresh Ocean Breeze

Aqua, Alcohol, Decyl Glucoside, Parfum, Aloe Barbadensis, Methylisothiazolinone, Methylchloroisothiazolinone, Disodium EDTA, Benzophenone-4, Citric Acid, Tocopheryl Acetate, Retinyl Palmitate, Red 33, Blue 1, Yellow 5

Cool Cucumber Melon

Aqua, Alcohol, Decyl Glucoside, Parfum, Aloe Barbadensis, Methylisothiazolinone, Methylchloroisothiazolinone, Disodium EDTA, Benzophenone-4, Citric Acid, Tocopheryl Acetate, Retinyl Palmitate, Red 33, Yellow 5

Irresistible Green Apple

Aqua, Alcohol, Decyl Glucoside, Parfum, Aloe Barbadensis, Methylisothiazolinone, Methylchloroisothiazolinone, Disodium EDTA, Benzophenone-4, Citric Acid, Tocopheryl Acetate, Retinyl Palmitate, Blue 1, Yellow 5

Alluring Red Pomegranate

Aqua, Alcohol, Decyl Glucoside, Parfum, Aloe Barbadensis, Methylisothiazolinone, Methylchloroisothiazolinone, Disodium EDTA, Benzophenone-4, Citric Acid, Tocopheryl Acetate, Retinyl Palmitate, Red 40, Red 33, Blue 1

Sparkling White Cranberry

Aqua, Alcohol, Decyl Glucoside, Parfum, Aloe Barbadensis, Methylisothiazolinone, Methylchloroisothiazolinone, Disodium EDTA, Benzophenone-4, Citric Acid, Tocopheryl Acetate, Retinyl Palmitate, Red 33, Yellow 5

Crisp White Citrus

Aqua, Alcohol, Decyl Glucoside, Parfum, Aloe Barbadensis, Methylisothiazolinone, Methylchloroisothiazolinone, Disodium EDTA, Benzophenone-4, Citric Acid, Tocopheryl Acetate, Retinyl Palmitate, Red 33, Yellow 5

Sheer Cotton Whisper

Aqua, Alcohol, Decyl Glucoside, Parfum, Aloe Barbadensis, Methylisothiazolinone, Methylchloroisothiazolinone, Disodium EDTA, Benzophenone-4, Citric Acid, Tocopheryl Acetate, Retinyl Palmitate, Blue 1, Red 33, Yellow 5

Questions?

1-800-FDA-1088

SATISFACTION GUARANTEED BY REFUND OR EXCHANGE

Made in China

Distributed by Jets, Sets & Elephants Beauty Corp.

For Shopko Stores Operating Co., LLC

Green Bay, WI 54307, U.S.A.

Package/Label Principal Display Panel - Bottle Label

bath+body+etc®

Warm

VANILLA SUGAR

antibacterial

foaming

hand sanitizer

PARABEN FREE

contains aloe vera and

vitamins A & E

1.7 fl oz

Warm Vanilla Sugar Bottle Label

Package/Label Principal Display Panel - Bottle Label

bath+body+etc®

Sweet

CHERRY BLOSSOM

antibacterial

foaming

hand sanitizer

PARABEN FREE

contains aloe vera and

vitamins A & E

1.7 fl oz

Sweet Cherry Blossom Bottle Label

Package/Label Principal Display Panel - Bottle Label

bath+body+etc®

Fresh

OCEAN BREEZE

antibacterial

foaming

hand sanitizer

PARABEN FREE

contains aloe vera and

vitamins A & E

1.7 fl oz

Fresh Ocean Breeze Bottle Label

Package/Label Principal Display Panel - Bottle Label

bath+body+etc®

Cool

CUCUMBER MELON

antibacterial

foaming

hand sanitizer

PARABEN FREE

contains aloe vera and

vitamins A & E

1.7 fl oz

Cool Cucumber Melon Bottle Label

Package/Label Principal Display Panel - Bottle Label

bath+body+etc®

Irresistible

GREEN APPLE

antibacterial

foaming

hand sanitizer

PARABEN FREE

contains aloe vera and

vitamins A & E

1.7 fl oz

Irresistible Green Apple Bottle Label

Package/Label Principal Display Panel - Bottle Label

bath+body+etc®

Alluring

RED POMEGRANATE

antibacterial

foaming

hand sanitizer

PARABEN FREE

contains aloe vera and

vitamins A & E

1.7 fl oz

Alluring Red Pomegranate Bottle Label

Package/Label Principal Display Panel - Bottle Label

bath+body+etc®

Sparkling

WHITE CRANBERRY

antibacterial

foaming

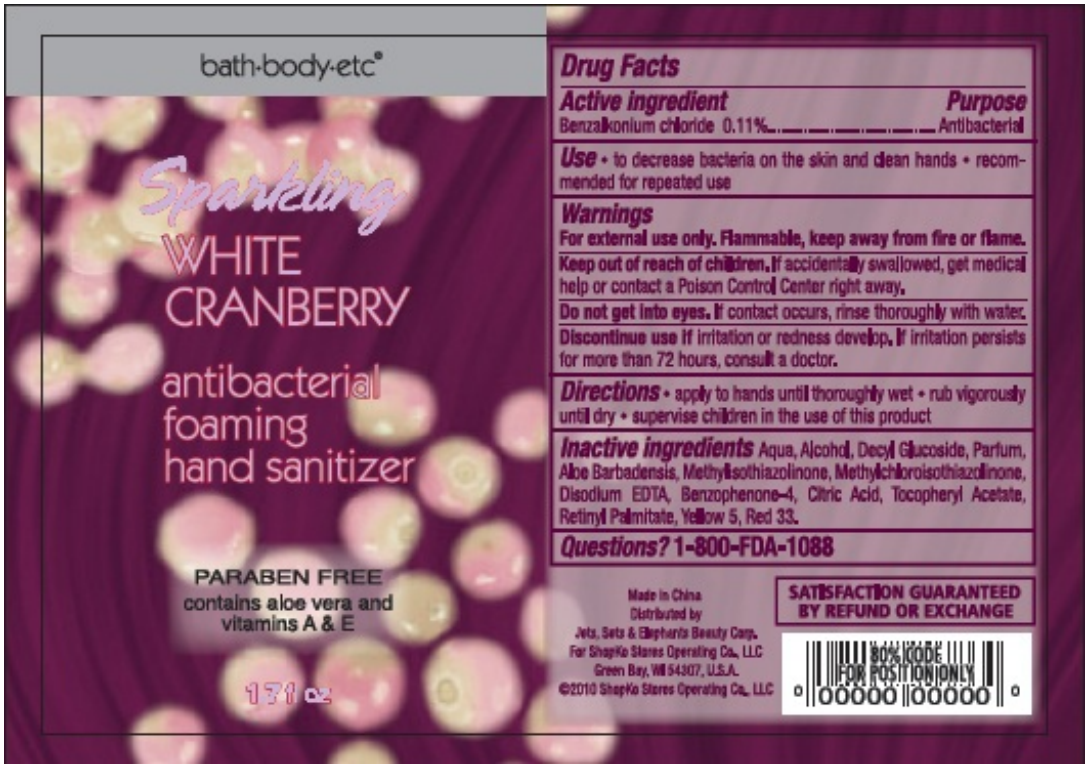
hand sanitizer

PARABEN FREE

contains aloe vera and

vitamins A & E

1.7 fl oz



Sparkling White Cranberry Bottle Label

Package/Label Principal Display Panel - Bottle Label

bath+body+etc®

Crisp

WHITE CITRUS

antibacterial

foaming

hand sanitizer

PARABEN FREE

contains aloe vera and

vitamins A & E

1.7 fl oz

Crisp White Citrus Bottle Label

Package/Label Principal Display Panel - Bottle Label

bath+body+etc®

Sheer

COTTON WHISPER

antibacterial

foaming

hand sanitizer

PARABEN FREE

contains aloe vera and

vitamins A & E

1.7 fl oz

Sheer Cotton Whisper Bottle Label

ANTIBACTERIAL FOAMING HAND SANITIZER

benzalkonium chloride solution

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:50988-190
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.055 mL in 50.275 mL

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
ALCOHOL (UNII: 3K9958V90M)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	
METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)	
EDETATE CALCIUM DISODIUM ANHYDROUS (UNII: 8U5D034955)	
SULISOBENZONE (UNII: 1W6L629B4K)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
.ALPHA.-TOCOPHEROL ACETATE, D- (UNII: A7E6112E4N)	
VITAMIN A PALMITATE (UNII: 1D1K0N0VVC)	
D&C RED NO. 33 (UNII: 9DBA0SBB0L)	

FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:50988-190-00	50.275 mL in 1 BOTTLE, PUMP		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part333E	08/03/2010	

ANTIBACTERIAL FOAMING HAND SANITIZER

benzalkonium chloride solution

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:50988-191
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.055 mL in 50.275 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
ALCOHOL (UNII: 3K9958V90M)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	
METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)	
EDETATE CALCIUM DISODIUM ANHYDROUS (UNII: 8U5D034955)	
SULISOBENZONE (UNII: 1W6L629B4K)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
.ALPHA.-TOCOPHEROL ACETATE, D- (UNII: A7E6112E4N)	
VITAMIN A PALMITATE (UNII: 1D1K0N0VVC)	
D&C RED NO. 33 (UNII: 9DBA0SBB0L)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date

1	NDC:50988-191-00	50.275 mL in 1 BOTTLE, PUMP		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part333E	08/03/2010	

ANTIBACTERIAL FOAMING HAND SANITIZER

benzalkonium chloride solution

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:50988-192
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.055 mL in 50.275 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
ALCOHOL (UNII: 3K9958V90M)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	
METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)	
EDETATE CALCIUM DISODIUM ANHYDROUS (UNII: 8U5D034955)	
SULISOBENZONE (UNII: 1W6L629B4K)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
.ALPHA.-TOCOPHEROL ACETATE, D- (UNII: A7E6112E4N)	
VITAMIN A PALMITATE (UNII: 1D1K0N0VVC)	
D&C RED NO. 33 (UNII: 9DBA0SBB0L)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:50988-192-00	50.275 mL in 1 BOTTLE, PUMP		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date

OTC monograph final	part333E	08/03/2010	

ANTIBACTERIAL FOAMING HAND SANITIZER

benzalkonium chloride solution

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:50988-193
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.055 mL in 50.275 mL

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
ALCOHOL (UNII: 3K9958V90M)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	
METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)	
EDETATE CALCIUM DISODIUM ANHYDROUS (UNII: 8U5D034955)	
SULISOBENZONE (UNII: 1W6L629B4K)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
.ALPHA.-TOCOPHEROL ACETATE, D- (UNII: A7E6112E4N)	
VITAMIN A PALMITATE (UNII: 1D1K0N0VVC)	
D&C RED NO. 33 (UNII: 9DBA0SBB0L)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:50988-193-00	50.275 mL in 1 BOTTLE, PUMP		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part333E	08/03/2010	

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Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.055 mL in 50.275 mL

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
ALCOHOL (UNII: 3K9958V90M)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	
METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)	
EDETATE CALCIUM DISODIUM ANHYDROUS (UNII: 8U5D034955)	
SULISOBENZONE (UNII: 1W6L629B4K)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
.ALPHA.-TOCOPHEROL ACETATE, D- (UNII: A7E6112E4N)	
VITAMIN A PALMITATE (UNII: 1D1K0N0VVC)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:50988-194-00	50.275 mL in 1 BOTTLE, PUMP		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part333E	08/03/2010	

ANTIBACTERIAL FOAMING HAND SANITIZER

benzalkonium chloride solution

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:50988-195
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.055 mL in 50.275 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
ALCOHOL (UNII: 3K9958V90M)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	
METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)	
EDETATE CALCIUM DISODIUM ANHYDROUS (UNII: 8U5D034955)	
SULISOBENZONE (UNII: 1W6L629B4K)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
.ALPHA.-TOCOPHEROL ACETATE, D- (UNII: A7E6112E4N)	
VITAMIN A PALMITATE (UNII: 1D1K0N0VVC)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
D&C RED NO. 33 (UNII: 9DBA0SBB0L)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:50988-195-00	50.275 mL in 1 BOTTLE, PUMP		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part333E	08/03/2010	

ANTIBACTERIAL FOAMING HAND SANITIZER

benzalkonium chloride solution

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:50988-196
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.055 mL in 50.275 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
ALCOHOL (UNII: 3K9958V90M)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	
METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)	
EDETATE CALCIUM DISODIUM ANHYDROUS (UNII: 8U5D034955)	
SULISOBENZONE (UNII: 1W6L629B4K)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
.ALPHA.-TOCOPHEROL ACETATE, D- (UNII: A7E6112E4N)	
VITAMIN A PALMITATE (UNII: 1D1K0N0VVC)	
D&C RED NO. 33 (UNII: 9DBA0SBB0L)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:50988-196-00	50.275 mL in 1 BOTTLE, PUMP		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part333E	08/03/2010	

ANTIBACTERIAL FOAMING HAND SANITIZER

benzalkonium chloride solution

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:50988-197
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.055 mL in 50.275 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
ALCOHOL (UNII: 3K9958V90M)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	
METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)	
EDETATE CALCIUM DISODIUM ANHYDROUS (UNII: 8U5D034955)	

SULISOBENZONE (UNII: 1W6L629B4K)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
.ALPHA.-TOCOPHEROL ACETATE, D- (UNII: A7E6112E4N)	
VITAMIN A PALMITATE (UNII: 1D1K0N0VVC)	
D&C RED NO. 33 (UNII: 9DBA0SBB0L)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:50988-197-00	50.275 mL in 1 BOTTLE, PUMP		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part333E	12/01/2010	

ANTIBACTERIAL FOAMING HAND SANITIZER

benzalkonium chloride solution

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:50988-198
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.055 mL in 50.275 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
ALCOHOL (UNII: 3K9958V90M)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	
METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)	
EDETATE CALCIUM DISODIUM ANHYDROUS (UNII: 8U5D034955)	
SULISOBENZONE (UNII: 1W6L629B4K)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
.ALPHA.-TOCOPHEROL ACETATE, D- (UNII: A7E6112E4N)	
VITAMIN A PALMITATE (UNII: 1D1K0N0VVC)	
D&C RED NO. 33 (UNII: 9DBA0SBB0L)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:50988-198-00	50.275 mL in 1 BOTTLE, PUMP		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part333E	12/01/2010	

Labeler - Jets, Sets, & Elephants Beauty Corp. (243254039)

Establishment

Name	Address	ID/FEI	Business Operations
Gold Orient International Limited		679905914	MANUFACTURE

Revised: 11/2010 Jets, Sets, & Elephants Beauty Corp.