SANITIZING HAND MIST UNSCENTED- ethyl alcohol spray
SANITIZING HAND MIST LEMONGRASS- ethyl alcohol spray
SANITIZING HAND GEL UNSCENTED- ethyl alcohol gel
SANITIZING HAND GEL PEPPERMINT EUCALYPTUS- ethyl alcohol gel
SANITIZING HAND MIST PEPPERMINT EUCALYPTUS- ethyl alcohol spray
SANITIZING HAND GEL LEMONGRASS- ethyl alcohol gel
ALO New York LLC

ALO Brands Hand Sanitzer

Active ingredients

Ethyl Alcoahol

Purpose

Antiseptic

Uses

For handwashing to decrease bacteria on the skin

Warnings

Flammable. Keep away from fire or flame. For external use only.

When using this product

Do not use in eyes. In case of contact with eyes, rinse with water.

Stop use and ask doctor if

irritation and redness develop and persist.

Keep out of reach of chidren.

If swallowed, get medical help promptly.

Directions

Wet hands thoroughly with product and allow to dry without wiping.

Other information

Store under 105°F

Inactive Ingredients

NDC 80588-010:
Water, Glycerine, Aloe Barbadensis Leaf
NDC 80588-011:
Water, Glycerine, Alce Barbadensis Leaf, Eucalyptus Radiata Flower/leaf/Stem Oil, Piper Mentha Piperita (Peopermint) Oil

NDC 80588-012:

Water, Glycerine, Aloe Barbadensis Leaf, Cymbopogon Schoenanthus (Lemongrass) Oil

NDC 80588-020:
Water, Glyperine, Rapidgel EZ. Aloe barbadensis Leaf
NDC 80588-011:
Water. Glycerine, Rapidgel EZ. Aloe Barbadensis Leaf. Eucalyptus Radiate Flower/Leaf/Stem Oil, Piper Mentha Piperita (Peppermint) Oil

NDC 80588-012:
Water, Glycerine, Rapidgel EZ. Aloe Barbadensis Leaf, Cymbopogon Schoenanthus (Lemongrass) Oil

Sanitizing Hand Mist - Lemongrass +Aloe NDC 82355-012-02

Sanitizing Hand Gel ReCharge - Peppermint Euclyptus +Aloe NDC 82355-021-32

Sanitizing Hand Gel ReCharge Lemongrass + Aloe NDC 82355-022-32

Sanitizing Hand Gel ReCharge - Unscented + Aloe NDC 82355-020-32

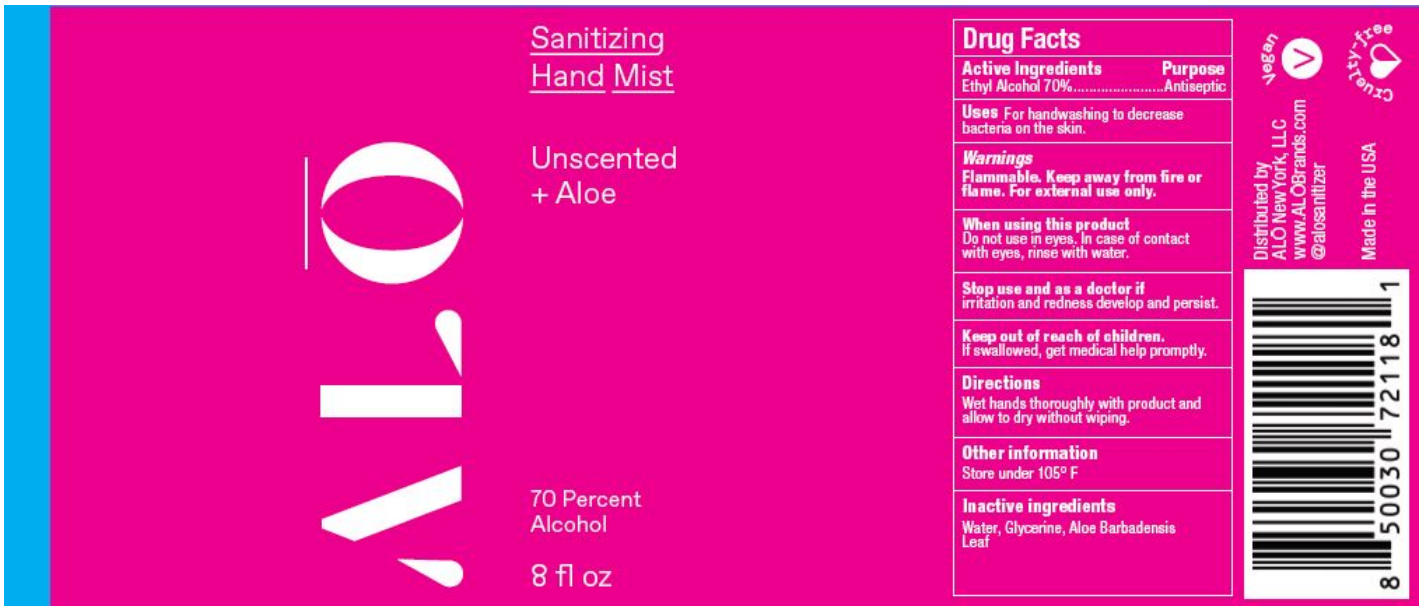
Sanitizing Hand Mist - Peppermint Eucalyptus + Aloe NDC 82355-011-02

Sanitizing Hand Mist - Unscented + Aloe NDC 82355-010-02

Sanitizing Hand Mist - Lemongrass +Aloe NDC 82355-012-08

Sanitizing Hand Mist - Peppermint Eucalypyus + Aloe NDC 82355-011-08

Sanitizing Hand Mist - Unscented + Aloe NDC 82355-010-08



Sanitizing Hand Gel - Lemongrass + Aloe NDC 82355-022-08

Sanitizing Hand Gel - Peppermint Eucalyptus + Aloe NDC 82355-021-08

Sanitizing Hand Gel - Unscented + Aloe NDC 82355-020-08

SANITIZING HAND MIST UNSCENTED

ethyl alcohol spray

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:82355-010
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	70 mL in 100 mL

Inactive Ingredients

Ingredient Name	Strength
ALOE (UNII: V5VD430YW9)	
GLYCERIN (UNII: PDC6A3C0OX)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:82355-010-02	59 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	06/09/2022	
2	NDC:82355-010-08	236 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	06/09/2022	

Marketing Information

Marketing	Application Number or Monograph	Marketing Start	Marketing End

Category	Citation	Date	Date
OTC Monograph Drug	M003	06/09/2022	

SANITIZING HAND MIST LEMONGRASS

ethyl alcohol spray

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:82355-012
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	70 mL in 100 mL

Inactive Ingredients

Ingredient Name	Strength
ALOE (UNII: V5VD430YW9)	
EAST INDIAN LEMONGRASS OIL (UNII: UP0M8M3VZW)	
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:82355-012-02	59 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	06/06/2022	
2	NDC:82355-012-08	238 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	06/06/2022	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M003	06/06/2022	

SANITIZING HAND GEL UNSCENTED

ethyl alcohol gel

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:82355-020
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	70 mL in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**
GLYCERIN (UNII: PDC6A3C0OX)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WQ0SDW1A)	
ACRYLATES/VINYL ISODECANOATE CROSSPOLYMER (10000 MPA.S NEUTRALIZED AT 0.5%) (UNII: 2N8MDB79NA)	
WATER (UNII: 059QF0KO0R)	
ALOE (UNII: V5VD430YW9)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:82355-020-08	238 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	06/09/2022	
2	NDC:82355-020-32	900 mL in 1 BOTTLE, DISPENSING; Type 0: Not a Combination Product	06/09/2022	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M003	06/09/2022	

SANITIZING HAND GEL PEPPERMINT EUCALYPTUS

ethyl alcohol gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:82355-021
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	70 mL in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**

EUCALYPTUS OIL (UNII: 2R04ONI662)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WQ0SDW1A)	
ACRYLATES/VINYL ISODECANOATE CROSSPOLYMER (10000 MPA.S NEUTRALIZED AT 0.5%) (UNII: 2N8MDB79NA)	
ALOE (UNII: V5VD430YW9)	
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
PEPPERMINT OIL (UNII: AV092KU4JH)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:82355-021-08	238 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	06/06/2022	
2	NDC:82355-021-32	900 mL in 1 BOTTLE, DISPENSING; Type 0: Not a Combination Product	06/06/2022	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M003	06/06/2022	

SANITIZING HAND MIST PEPPERMINT EUCALYPTUS

ethyl alcohol spray

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:82355-011
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	70 mL in 100 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
ALOE (UNII: V5VD430YW9)	
PEPPERMINT OIL (UNII: AV092KU4JH)	
EUCALYPTUS OIL (UNII: 2R04ONI662)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:82355-011-02	59 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	06/06/2022	
2	NDC:82355-011-08	238 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	06/06/2022	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M003	06/06/2022	

SANITIZING HAND GEL LEMONGRASS

ethyl alcohol gel

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:82355-022
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	70 mL in 100 mL

Inactive Ingredients

Ingredient Name	Strength
GLYCERIN (UNII: PDC6A3C0OX)	
WATER (UNII: 059QF0KO0R)	
ALOE (UNII: V5VD430YW9)	
EAST INDIAN LEMONGRASS OIL (UNII: UP0M8M3VZW)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WQ0SDW1A)	
ACRYLATES/VINYL ISODECANOATE CROSSPOLYMER (10000 MPA.S NEUTRALIZED AT 0.5%) (UNII: 2N8MDB79NA)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:82355-022-08	238 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	06/06/2022	
2	NDC:82355-022-32	900 mL in 1 BOTTLE, DISPENSING; Type 0: Not a Combination Product	06/06/2022	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M003	06/06/2022	

Labeler - ALO New York LLC (110122374)

Registrant - Alkaline Corporation (790098318)

Establishment

Name	Address	ID/FEI	Business Operations
Alkaline Corporation		790098318	manufacture(82355-010, 82355-011, 82355-012, 82355-020, 82355-021, 82355-022) , pack(82355-010, 82355-011, 82355-012, 82355-020, 82355-021, 82355-022) , label(82355-010, 82355-011, 82355-012, 82355-020, 82355-021, 82355-022)

Revised: 2/2024 ALO New York LLC