

INSTANT HAND SANITIZER- instant hand sanitizer gel

Accessory Myxx LLC

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.

Hand Sanitizer

Instant Hand Sanitizer

Warnings

For external use only: hands

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

Stop use and ask a doctor if irritation and redness develop condition persists for more than 72 hours

avoid contact with broken skin do not inhale or ingest

General Warnings

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Ethyl alcohol 70%-75%

Use to decrease bacteria on the skin. Use on hands only.

Water, Aloe Barbadosensis Leaf Juice, Maltodextrin, Glycerin, Propylene Glycol, Acrylates/C10-30 Alkyl Acrylate Crosspolymer, Triethanoamine, Fragrance, Lactose,

Microcrystalline Cellulose, Sucrose, Zea Mays (Corn) starch, Ultramarine blue CI 77007, Tocopheryl Acetate, Hydroxypropyl Methyl Cellulose, D&C Red NO.33, FD&C Blue No.1

To clean hands and fight bacteria on that skin

Squeeze the bottle on your hands so the liquid comes out. Rub hands together. Allow hands to dry.

Not intended for children under the age of 12. Please keep out of reach of children.







**ANTI-BACTERIAL
HAND SANITIZER**



Kills Up To
99.9%
Of Germs

1oz / 3 ML





Drug Facts

Active ingredient

Ethyl Alcohol 70-75%.....

Purpose

Antiseptic

Use for hand washing to decrease bacteria on the skin.

Warnings

warnings

For external use only.

Flammable. Keep away from fire or flames.

When using this product keep out of eyes. In case of contact with eyes, rinse with water.

Stop use and ask doctor if irritation and redness develops and persists.

Keep out of reach of children

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

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

Other information Store under 105° F.

Inactive Ingredients Purified Water, Glycerin, Isopropyl, Myristate, Carbomer, Aloe Vera, Fragrance.

Drug Facts

Active ingredient

Ethyl Alcohol 70%

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

Other information Store under 105° F.

Inactive ingredient

Water(aqua), Glycerin, Carbomer, Aminomethyl propanol, Fragrance(parfum), Aloe barbadensis leaf extract, FD & C Blue No.1 (CI 42090).

Distributed by: Accessory Myxx LLC New York, NY 10018

Expiry Date: 08/2022

Batch No: HS2020218

Made in China

www.shopmyxx.com



Drug Facts

Active ingredient

Ethyl Alcohol 70%

Purpose Antiseptic

Use

For hand washing to decrease bacteria on the skin.

Warnings

For external use only.

Flammable. Keep away from fire or flame.

When using this product keep out of eyes. In case of contact with eyes, rinse with water.

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Distributed by: Accessory Myxx LLC New York, NY 10018
Expiry Date: 07/2022 Batch No: HS2020301

Made in China



www.shopmyxx.com

Style No: 8940CM



Drug Facts (continued)

Warnings

Keep out of reach of children.
If swallowed, get medical help or contact a Poison Control Center right away.

Directions

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

Other information

Store under 105° F.

Inactive ingredients

Water(aqua), Glycerin, Carbomer, Aminomethyl propanol, Fragrance(parfum), Aloe barbadensis leaf extract, FD & C Blue No.1 (CI 42090).

Distributed by: Accessory Myxx LLC New York, NY 10018
Expiry Date: 07/2022 Batch No: HS2020301

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Style No: 8940CM



INSTANT HAND SANITIZER

instant hand sanitizer gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:726 19-2020
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	75 mL in 100 mL

Inactive Ingredients

Ingredient Name	Strength
AMINOMETHYL PROPANEDIOL (UNII: CZ7BU4QZJZ)	0.18 mL in 100 mL

WATER (UNII: 059QF0KO0R)	28.42 mL in 100 mL
GLYCERIN (UNII: PDC6A3C0OX)	1 mL in 100 mL
CARBOMER 980 (UNII: 4Q93RCW27E)	0.35 mL in 100 mL
FD&C BLUE NO. 1 (UNII: HBR47K3TBD)	0.0001 mL in 100 mL
FRAGRANCE CLEAN ORC0600327 (UNII: 329LCV5BTF)	0.04 mL in 100 mL
ALOE VERA LEAF (UNII: ZY81Z83H0X)	0.35 mL in 100 mL

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72619-2020-1	56 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	05/18/2020	
2	NDC:72619-2020-2	500 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	05/18/2020	
3	NDC:72619-2020-4	3 mL in 1 PACKET; Type 0: Not a Combination Product	05/18/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC mono graph not final	part333A	05/18/2020	

INSTANT HAND SANITIZER

instant hand sanitizer gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:72619-1219
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	62 mL in 100 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
FD&C BLUE NO. 1 (UNII: HBR47K3TBD)	
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)	
SUCROSE (UNII: C151H8M554)	
ZEA MAYS POLLEN (UNII: 74PD8J616H)	
ULTRAMARINE BLUE (UNII: I39WR998B1)	
3-HEXYLOXYPROPYLENE GLYCOL (UNII: 3485P35DA4)	
(C10-C30)ALKYL METHACRYLATE ESTER (UNII: XH2FQZ38D8)	

GLYCERIN (UNII: PDC6A3C0OX)				
MALTODEXTRIN (UNII: 7CVR7L4A2D)				
D&C RED NO. 33 (UNII: 9DBA0SBB0L)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72619-1219-3	2.9 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	10/10/2018	
2	NDC:72619-1219-2	56 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	10/10/2018	
Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph not final	part333A	10/10/2018		

INSTANT HAND SANITIZER				
instant hand sanitizer gel				
Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:72619-1001	
Route of Administration	TOPICAL			
Active Ingredient/Active Moiety				
Ingredient Name	Basis of Strength	Strength		
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	62 mL in 100 mL		
Inactive Ingredients				
Ingredient Name	Strength			
WATER (UNII: 059QF0KO0R)				
CARBOMER 940 (UNII: 4Q93RCW27E)				
AMINOMETHYL PROPANEDIOL (UNII: CZ7BU4QZJZ)				
GLYCERIN (UNII: PDC6A3C0OX)				
ALOE VERA LEAF (UNII: ZY81Z83H0X)				
FD&C BLUE NO. 1 (UNII: HBR47K3TBD)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72619-1001-3	2.9 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	10/10/2018	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	10/10/2018	

Labeler - Accessory Myxx LLC (080553023)

Establishment

Name	Address	ID/FEI	Business Operations
Leeds (China) Biotechnology Co., Ltd		416037073	manufacture(72619-2020)

Establishment

Name	Address	ID/FEI	Business Operations
Nantong Health & Beyond Hygienic Products Inc.		421280161	manufacture(72619-1219)

Establishment

Name	Address	ID/FEI	Business Operations
NINGBO OCEANSTAR CHEMICAL PRODUCTS CO LTD		544493972	manufacture(72619-1001)

Revised: 5/2020

Accessory Myxx LLC