

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

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

Ethyl alcohol 62%

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,

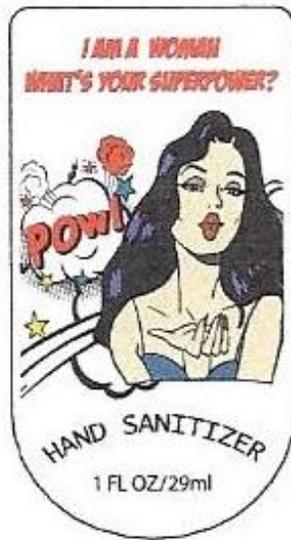
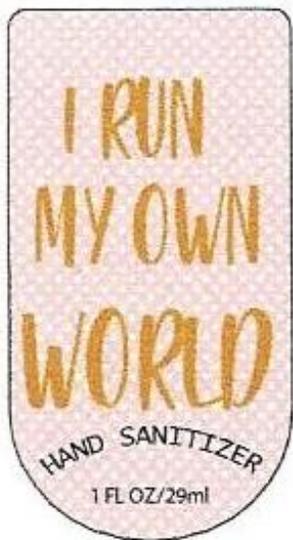
Microcrystalline Cellulose, Sucrose,
Zea Mays (Corn) starch, Ultramarine
blue CI 77007, Tocopheryl Acetate,
Hydroxypropyl Methyl Cellulose,
D&C Red NO.33, FD&C Blue No.1

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.





Factor II S top use and ask a



INSTANT HAND SANITIZER

instant hand sanitizer gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:726 19-1219
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: H3R47K3TBD)	
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:726 19-1219-1	2.9 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	10/10/20 18	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	10/10/20 18	

INSTANT HAND SANITIZER

instant hand sanitizer gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:726 19-100 1
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: 059QF0K00R)	
CARBOMER 940 (UNII: 4Q93RCW27E)	
TROLAMINE (UNII: 9O3K93S3TK)	
GLYCERIN (UNII: PDC6A3C0OX)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:726 19-100 1-1	2.9 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	10/10/20 18	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	10/10/20 18	

Labeler - Accessory Myxx LLC (080553023)

Establishment			
Name	Address	ID/FEI	Business Operations
Nantong Health & Beyond Hygienic Products Inc.		421280 16 1	manufacture(726 19-1219)

Establishment			
Name	Address	ID/FEI	Business Operations
NINGBO OCEANSTAR CHEMICAL PRODUCTS CO LTD		54449 39 72	manufacture(726 19-100 1)