DRPURX HAND SANITIZER- alcohol gel
Loom USA

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.

DrPuRx HAND SANITIZER

Drug Facts

Active Ingredients

Ethyl Alcohol 70%

Purpose

Antimicrobial

Uses

- to decrease bacteria on the skin that could cause disease
- recommended for repeated use

Warnings

For external use only: hands

Flammable, keep away from fire or flame

When using this product

- keep out of eyes. In case of contact with eyes, flush thoroughly with water.
- avoid contact with broken skin
- do not inhale or ingest.

Stop use and ask a doctor if

- irritation and redness develop
- condition persists for more than 72 hours

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
- for children under 6, use only under adult supervision
- not recommended for infants

Other information

- do not store above 85°
- may discolor some fabrics
- harmful to wood finished and plastics

Inactive ingredients

water, acrylates/C10-30 alkyl acrylate crosspolymer, sodium PCA, hyaluronic acid

Distributed by:
LoomUSA LLC
3 Graphics Dr
Ewing, NJ 08628

Manufactured by:
LoomUSA LLC
FDA Registered Facility

PRINCIPAL DISPLAY PANEL - 499.79 mL Bottle Label

Made in
the USA

DrPuRX

HAND SANITIZER

Kills more than 99% of infectious agents*

Moisturizers leavehands smooth

16.9 FL OZ (499.79mL)

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*Effective at eliminating more than 99.98% of many harmful germs and bacteria in as little as 15 seconds.

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DUNS 11-751-4119

DRPURX HAND SANITIZER

alcohol gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:80607-921

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)	
CARBOMER INTERPOLYMER TYPE A (ALLYL SUCROSE CROSSLINKED) (UNII: 59TL3WG5CO)	
SODIUM PYRROLIDONE CARBOXYLATE (UNII: 469OTG57A2)	
HYALURONIC ACID (UNII: S270N0TRQY)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:80607-921-01	250 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/01/2020	
2	NDC:80607-921-02	500 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/01/2020	
3	NDC:80607-921-03	50 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/01/2020	
4	NDC:80607-921-04	118.2 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/01/2020	
5	NDC:80607-921-05	3785.4 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/01/2020	
6	NDC:80607-921-06	208197.65 mL in 1 DRUM; Type 0: Not a Combination Product	03/25/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC MONOGRAPH NOT FINAL	part333A	03/01/2020	

Labeler - Loom USA (117514119)

Establishment

Name	Address	ID/FEI	Business Operations
LoomUSA LLC		117514119	MANUFACTURE(80607-921)