PREFERENCE HAND SANITIZER- alcohol spray
Whip Mix Corporation

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.

Preference Hand Sanitizer

Active Ingredient

Ethanol 72%

Purpose

Liquid Hand Sanitizer

Use

Used by Health Care Professionals for preventing the spread of both pathogenic microorganisms as well as resident bacterial flora and reducing healthcare-associated infections.

Warnings

Flammable, Keep away from fire or flame.

For External Use Only

Do not get in the eyes. If splashed in eyes, rinse promptly and thoroughy with water. Discontinue use if irritation and redness develop. If irritation persists for more that 5 days, consult a physician.

Keep out of reach of children

In case of alcohol ingestion, seek professional assistance or contact a Poison Control Center immediately.

Directions

Apply product liberally on hands and rub together until dry.

Inactive Ingredients

Cyclomethicone, Sodium Lactate, Isopropyl Myristace, Cyclomethicone (and) Trimethyisiloxyl Silicane.

Principal Display Panel

Preference

Liquid Hand Sanitizer

Favored 3:1

over the leanding liquid sanitizer

- Kills 99.9% of most common germs
- Dries in less than 10 seconds
- Moisturizes the skin

- Not harmful to beneficial micro-organisms

PREFERENCE HAND SANITIZER

alcohol spray

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:62223-001
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	340.56 mL in 470 mL

Inactive Ingredients	
Ingredient Name	**Strength**
CYCLOMETHICONE (UNII: NMQ347994Z)	
SODIUM LACTATE (UNII: TU7HW0W0QT)	
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:62223-001-16	470 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/23/2020	

2	NDC:62223-001-32	950 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/23/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	03/23/2020	

Labeler - Whip Mix Corporation (006385173)

Establishment

Name	Address	ID/FEI	Business Operations
Whip Mix Corporation		006385173	repack(62223-001) , manufacture(62223-001) , relabel(62223-001)

Revised: 3/2020 Whip Mix Corporation