PURE CLEAN PREMIUM HAND SANITIZER- hand sanitizer liquid
WOMANSWORLD Co., Ltd.

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.

Active Ingredient

Alcohol 75% v/v

Purpose

Antiseptic

Use

- Hand sanitizer to help reduce bacteria that potentially can cause disease.

Warnings

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

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

Stop use and ask a doctor if irritation or rash occurs. These may be signs of serious condition.

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

Directions

- Place enough product on your palm to thoroughly cover your hands.
- Rub hands together briskly until dry.
- Supervise children when using this product to avoid swallowing.

Other Information

- Store between 15-30 °C (59-86°F).
- Avoid freezing and excessive heat above 104°F(40°C).
- May discolor certain fabrics or surface.

Inactive Ingredients

Water, Butylene Glycol, Glycerin, Aloe Vera Leaf, Sodium Hyaluronate, 1,2-Hexanediol

Package Label - Principal Display Panel

PURE CLEAN PREMIUM HAND SANITIZER

hand sanitizer liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:77708-001
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**
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	
HYALURONATE SODIUM (UNII: YSE9PPT4TH)	
1,2-HEXANEDIOL (UNII: TR046Y3K1G)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:77708-001-10	50 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	05/20/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	05/20/2020	

Labeler - WOMANSWORLD Co., Ltd. (688320057)

Registrant - WOMANSWORLD Co., Ltd. (688320057)

Establishment

Name	Address	ID/FEI	Business Operations
ICOMAX Co., Ltd.		694834031	manufacture(77708-001)

Revised: 6/2020 WOMANSWORLD Co., Ltd.