

EXIHUAN HAND SANITIZER- alcohol liquid
Yunnan Thousand Sheng Pharmaceutical Group 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.

EXIHUAN Hand Sanitizer

This is a hand sanitizer manufactured according to the Temporary Policy for Preparation of Certain Alcohol-Based Hand Sanitizer Products During the Public Health Emergency (CoViD-19); Guidance for Industry.

The hand sanitizer is manufactured using only the following United States Pharmacopoeia (USP) grade ingredients in the preparation of the product (percentage in final product formulation) consistent with World Health Organization (WHO) recommendations:

- a. Alcohol (ethanol) (USP or Food Chemical Codex (FCC) grade) (75%, volume/volume (v/v)) in an aqueous solution denatured according to Alcohol and Tobacco Tax and Trade Bureau regulations in 27 CFR part 20.
- b. Sodium hyaluronate
- c. Butanediol
- d. Triethanolamine
- e. collagen
- f. Carbomer
- g. Sterile distilled water or boiled cold water.

The firm does not add other active or inactive ingredients. Different or additional ingredients may impact the quality and potency of the product.

Active Ingredient(s)

Alcohol 75% v/v. Purpose: Antiseptic

Purpose

Antiseptic, Hand Sanitizer

Use

Hand Sanitizer to help reduce bacteria that potentially can cause disease. For use when soap and water are not available.

Warnings

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

Do not use

- in children less than 2 months of age
- on open skin wounds

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

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

WATER (UNII: 059QF0KO0R)					
BUTANEDIOL (MIXED ISOMERS) (UNII: TMS4MGA0H4)					
TROLAMINE (UNII: 9O3K93S3TK)					
COLLAGENASE CLOSTRIDIUM HISTOLYTICUM (UNII: 9X7O8V25IT)					
Packaging					
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:74431-208-01	10 L in 1 BOTTLE; Type 0: Not a Combination Product	03/30/2020		
Marketing Information					
Marketing Category		Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph not final		part333A	03/30/2020		

Labeler - Yunnan Thousand Sheng Pharmaceutical Group Co. Ltd. (554528835)

Establishment				
Name		Address	ID/FEI	Business Operations
Yunnan Thousand Sheng Pharmaceutical Group Co. Ltd.			554528835	manufacture(74431-208)

Revised: 4/2020

Yunnan Thousand Sheng Pharmaceutical Group Co. Ltd.