# ALCOHOL WIPES- alcohol cloth Guangzhou Baihua Co., Ltd.

Disclaimer: This drug has not been found by FDA to be safe and effective, and this labeling has not been approved by FDA. For further information about unapproved drugs, click here.

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### **Alcohol Wipes**

#### **Purpose**

PURPOSE: ANTIMICROBIAL

#### **Active Ingredient(s)**

**ACTIVE INGREDIENT: ETHYL ALCOHOL 75%** 

#### Use

USES: HAND SANITIZER TO HELP REDUCE BACTERIA ON THE SKIN.

### Warnings

WARNINGS: FLAMMABLE, KEEP AWAY FROM FIRE OR FLAME

- FOR EXTERNAL USE ONLY.
- DO NOT USE IN OR NEAR THE EYES.

IN CASE OF CONTACT, RINSE EYES THOROUGHLY WITH WATER.

- STOP USE AND CONSULT WITH A PHYSICIAN IF IRRITATION DEVELOPS AND LASTS FOR MORE THAN 72 HOURS.
- KEEP OUT OF REACH OF CHILDREN.
- IF SWALLOWED, GET MEDICAL HELP OR CONTACT A POISON CONTROL CENTER IMMEDIATELY.

#### Do not use

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#### **Directions**

- DISPENSING. LIFT THE FRONT LID. OPEN PROTECTIVE SEAL. PULL OUT WIPE, RESEAL, AND CLOSE THE LID.
- USE. WIPE HANDS THOROUGHLY.
- DISPOSAL. DO NOT FLUSH.

#### Directions of Use:

- 1. Open the lid and slowly peel back label. 2. Pull out wipes as needed.
- 3. Reseal label to retain moisture.

#### Other information

STORAGE. STORE AT ROOM TEMPERATURE.

#### **Inactive ingredients**

Glycerol, Hydrogen Peroxide, RO pure water

# **Package Label - Principal Display Panel**

Alcohol Wipes(80pcs) 77665-104-01







Alcohol Wipes (20pcs) 77665-104-02





## **ALCOHOL WIPES**

alcohol cloth

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:77665-104	

**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		
GLYCERIN (UNII: PDC6A3C0OX)			
HYDROGEN PEROXIDE (UNII: BBX060AN9V)			
WATER (UNII: 059QF0KO0R)			

Packaging			
# Item Code	Package Description	Marketing Start Date	Marketing End Date
NDC:77665-104-	80 mL in 1 PACKAGE; Type 0: Not a Combination Product	07/18/2023	
NDC:77665-104- 02	20 mL in 1 PACKAGE; Type 0: Not a Combination Product	07/18/2023	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		07/18/2023	
		07/18/2023	

# Labeler - Guangzhou Baihua Co., Ltd. (545015588)

# Registrant - Guangzhou Baihua Co., Ltd. (545015588)

Establishment				
Name	Address	ID/FEI	<b>Business Operations</b>	
Guangzhou Baihua Co., Ltd		545015588	manufacture(77665-104)	

Revised: 1/2024 Guangzhou Baihua Co., Ltd.