

ALCOHOL WIPES- alcohol wipes patch
Fujian Yifa Healthcare Products 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.

Yifa, Alcohol Wipes, 3 AIs, *50

Active ingredient

Benzalkonium Chloride, 0.15% w/w;

Didecyldimonium Chloride, 0.2% w/w;

Ethanol Alcohol, 15% w/w.

Inactive ingredient

Phenoxyethanol, Propylene Glycol, Glycerin, Water.

Purpose

Sterilization.

When using

Skin's cleaning care, Article surface's cleaning and maintenance.

Do not use

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

Stop use

Discontinue use if allergic reactions occur.

Keep out of reach of children

Keep out of reach of children.

Indications & usage

Skin's cleaning care, Article surface's cleaning and maintenance.

Remove seal, pull out sheet and wipe skin or articles.

(For external use only)

Dosage & administration

Use it as needed, after following the usage instructions.

For external use only.

Warnings

This product is disposable and should not be reused.

Check whether the package is in good condition before use. If the package is found to be untidy, it is not suitable for use.

Package label. Principal display panel



整体效果图



正面效果图



内包效果图

ALCOHOL WIPES

alcohol wipes patch

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:75543-101
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.15 U in 100 U
DIDECYLDIMONIUM CHLORIDE (UNII: JXN40O9Y9B) (DIDECYLDIMONIUM - UNII:Z7F472XQPA)	DIDECYLDIMONIUM CHLORIDE	0.2 U in 100 U
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	15 U in 100 U

Inactive Ingredients

Ingredient Name	Strength
GLYCERIN (UNII: PDC6A3C0OX)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
WATER (UNII: 059QF0K00R)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	

Product Characteristics

Color		Score	
Shape	FREEFORM	Size	
Flavor		Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:75543-101-15	50 in 1 PACKAGE	05/05/2020	
1	NDC:75543-101-01	1 U in 1 PATCH; Type 0: Not a Combination Product		

Marketing Information

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

Labeler - Fujian Yifa Healthcare Products Co., Ltd. (420510793)

Registrant - A03 Lab of BTS (548009541)

Establishment

Name	Address	ID/FEI	Business Operations
Fujian Yifa Healthcare Products Co., Ltd.		420510793	label(75543-101) , manufacture(75543-101)

Revised: 5/2020

Fujian Yifa Healthcare Products Co., Ltd.