WET WIPES- ethyl alcohol swab
Zhejiang Qimei Commodity 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.

69821-002 75% Ethyl Alcohol Wet Wipes

Active ingredient

Ethyl Alcohol 75%

Purpose

Antiseptic Handwash

Uses

■ For hand washing to dec rease bacteria on the skin. Recommended for repeated use.

Warnings

For external use only
Flammable, Keep away from fire or flame
When using this product avoid contact with eyes.
In case of eye contact, rins e with water to remove.
Stop use and ask a doctor if irritation and redness develop

Keep out of reach of children

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

Directions

■Wet hands thoroughly wi th product and allow to dry without wiping.

Aqua

WET WIPES

ethyl alcohol swab

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:69821-002
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	75 g in 100 g

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:69821-002-01	1 in 1 PACKAGE	04/20/2020	
1		3 g in 1 POUCH; Type 0: Not a Combination Product		
2	NDC:69821-002-02	10 in 1 PACKAGE	04/20/2020	
2		3 g in 1 POUCH; Type 0: Not a Combination Product		
3	NDC:69821-002-03	15 in 1 PACKAGE	04/20/2020	
3		3 g in 1 POUCH; Type 0: Not a Combination Product		
4	NDC:69821-002-04	20 in 1 PACKAGE	04/20/2020	
4		3 g in 1 POUCH; Type 0: Not a Combination Product		
5	NDC:69821-002-05	50 in 1 PACKAGE	04/20/2020	
5		3 g in 1 POUCH; Type 0: Not a Combination Product		
6	NDC:69821-002-06	60 in 1 PACKAGE	04/20/2020	
6		3 g in 1 POUCH; Type 0: Not a Combination Product		
7	NDC:69821-002-07	80 in 1 PACKAGE	04/20/2020	

7		3 g in 1 POUCH; Type 0: Not a Combination Product		
8	NDC:69821-002-09	50 in 1 PAIL	04/20/2020	
8		1.7 g in 1 POUCH; Type 0: Not a Combination Product		
9	NDC:69821-002-08	600 in 1 PACKAGE	04/20/2020	
9		3 g in 1 POUCH; Type 0: Not a Combination Product		
10	NDC:69821-002-10	160 in 1 PAIL	04/20/2020	
10		1.7 g in 1 POUCH; Type 0: Not a Combination Product		
11	NDC:69821-002-11	250 in 1 PAIL	04/20/2020	
11		1.7 g in 1 POUCH; Type 0: Not a Combination Product		
12	NDC:69821-002-12	600 in 1 PAIL	04/20/2020	
12		1.7 g in 1 POUCH; 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	04/20/2020	

Labeler - Zhejiang Qimei Commodity Co.,Ltd. (544331136)

Establishment

Name	Address	ID/FEI	Business Operations
Zhejiang Qimei Commodity Co.,Ltd.		544331136	manufacture(69821-002)

Revised: 5/2020 Zhejiang Qimei Commodity Co.,Ltd.