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.

69821-003 70% Alcohol Wet Wipes

Active ingredient

Ethyl Alcohol 70%

Purpose

Antiseptic

Uses

●For hand sanitizing to decrease bacteria on the skin
●Recommended for repeated use

Warnings

. FOR EXTERNAL USE ONLY
. FLAMMABL E, KEEP AWAY FROM FIRE OR FL AME
●When using this product avoid contact with eyes. In case of eye contact, flush eyes with water and seek medical help.
●If swallowed drink large amounts of water and seek medical attention.
●Avoid contact with face and broken skin. If contact occurs, flush with water.
●Stop use and ask a doctor if rritation or redness develops

Note Do not store above00°F. Non-stainingMay discolor some fabrics.

Keep out of reach of children

Keep out of reach of children.

Directions

●Take wipe and rub thoroughly over all surfaces of both hands
●Allow to dry without wiping
●Dispose of wipe in trash after use. Do not flush.

Purfied Water, Anionic Surfactant,
Propylene Glycol, Glycerin, Aloe Barbadensis Leaf Juice.

WET WIPES

ethyl alcohol swab

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:69821-003
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	70 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
GLYCERIN (UNII: PDC6A3C0OX)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
DECYL GLUCOSIDE (UNII: Z17H97EA6Y)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69821-003-01	10 in 1 PACKAGE	04/20/2020	
1		2.5 g in 1 POUCH; Type 0: Not a Combination Product		
2	NDC:69821-003-02	20 in 1 PACKAGE	04/20/2020	
2		2.5 g in 1 POUCH; Type 0: Not a Combination Product		
3	NDC:69821-003-03	30 in 1 PACKAGE	04/20/2020	
3		2.5 g in 1 POUCH; Type 0: Not a Combination Product		
4	NDC:69821-003-04	50 in 1 PACKAGE	04/20/2020	
4		2.5 g in 1 POUCH; Type 0: Not a Combination Product		
5	NDC:69821-003-05	80 in 1 PACKAGE	04/20/2020	
5		2.5 g in 1 POUCH; Type 0: Not a Combination Product		
6	NDC:69821-003-06	100 in 1 BAG	04/20/2020	
6		2.5 g in 1 POUCH; Type 0: Not a Combination Product		
7	NDC:69821-003-07	160 in 1 BAG	04/20/2020	
7		0.48 g in 1 POUCH; Type 0: Not a Combination Product		
8	NDC:69821-003-09	135 in 1 PAIL	04/20/2020	
8		0.48 g in 1 POUCH; Type 0: Not a Combination Product		
9	NDC:69821-003-08	1200 in 1 BAG	04/20/2020	
9		0.47 g in 1 POUCH; Type 0: Not a Combination Product		
10	NDC:69821-003-10	250 in 1 PAIL	04/20/2020	
10		0.48 g in 1 POUCH; Type 0: Not a Combination Product		
11	NDC:69821-003-11	1500 in 1 PAIL	04/20/2020	
11		0.48 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)

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

Establishment

Name	Address	ID/FEI	Business Operations
Zhejiang Qimei Commodity Co.,Ltd.		544331136	manufacture(69821-003)

Revised: 11/2022 Zhejiang Qimei Commodity Co.,Ltd.