HAND SANITIZER WIPES- ethyl alcohol cloth 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.

69821-006 70% Ethyl Alcohol Hand Sanitizer Wipes

Active ingredient

Ethyl Alcohol 70%

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 with product and allow to dry without wiping.

Aloe Barbadensis Leaf Juice, Carbormer, Diazolidinyl Urea, Glycerin, Methylparaben, Polysorbate 20, Propylene Glycol, Propylparaben, Tocopheryl Acetate, Trietha nolamine, Water



HAND SANITIZER WIPES

ethyl alcohol cloth

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:69821-006

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name

Basis of Strength

ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)

ALCOHOL

ALCOHOL

70 mL in 100 g

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 0590F0KO0R)	

ALOE VERA LEAF (UNII: ZY81Z83H0X)

CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)

DIAZOLIDINYL UREA (UNII: H5RIZ3MPW4)

GLYCERIN (UNII: PDC6A3C0OX)

PROPYLENE GLYCOL (UNII: 6DC9Q167V3)
PROPYLPARABEN (UNII: Z8IX2SC10H)

TROLAMINE (UNII: 903K93S3TK)

POLYSORBATE 20 (UNII: 7T1F30V5YH)

ALPHA-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)

METHYLPARABEN (UNII: A2I8C7HI9T)

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#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69821-006- 01	135 in 1 PAIL	11/01/2020	

1		570 g in 1 POUCH; Type 0: Not a Combination Product		
2	NDC:69821-006- 02	20 in 1 BAG	11/01/2020	
2		50 g in 1 POUCH; Type 0: Not a Combination Product		
3	NDC:69821-006- 03	30 in 1 BAG	11/01/2020	
3		75 g in 1 POUCH; Type 0: Not a Combination Product		
4	NDC:69821-006- 04	50 in 1 BAG	11/01/2020	
4		125 g in 1 POUCH; Type 0: Not a Combination Product		
5	NDC:69821-006- 05	80 in 1 BAG	11/01/2020	
5		200 g in 1 POUCH; Type 0: Not a Combination Product		
6	NDC:69821-006- 06	100 in 1 BAG	11/01/2020	
6		250 g in 1 POUCH; Type 0: Not a Combination Product		
7	NDC:69821-006- 07	160 in 1 BAG	11/01/2020	
7		77 g in 1 POUCH; Type 0: Not a Combination Product		
8	NDC:69821-006- 09	160 in 1 PAIL	11/01/2020	
8		675 g in 1 POUCH; Type 0: Not a Combination Product		
9	NDC:69821-006- 08	250 in 1 BAG	11/01/2020	
9		120 g in 1 POUCH; Type 0: Not a Combination Product		
10	NDC:69821-006- 10	250 in 1 PAIL	11/01/2020	
10		120 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	11/01/2020		

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

Establishment				
Name	Address	ID/FEI	Business Operations	
Zhejiang Qimei Commodity Co.,Ltd.		544331136	manufacture(69821-006)	