

ALCOHOL WIPES (75% ALCOHOL)- alcohol cloth
NanTong Guarder Medical Technology 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.

Alcohol wipes (75% Alcohol)

Active Ingredient(s)

Alcohol 75% v/v. Purpose: Antiseptic

Purpose

Antiseptic

Use

Alcohol wet wipe to help reduce bacteria that potentially can cause disease. For use when soap and water are not available.

Warnings

For external use only. Flammable. Keep away from heat or flame

Do not use

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

When using this product keep out of eyes, ears, and mouth. In case of contact with eyes, rinse eyes thoroughly with water.

Stop use and ask a doctor if irritation or rash occurs. These may be signs of a serious condition.

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

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Directions

Wet hands thoroughly with product and allow to dry.

Discard wipes in trash receptacle after use. Do not flush.

Children under 6 years of age should be supervised when using this product.

Other information

- Store between 15-30C (59-86F)
- Avoid freezing and excessive heat above 40C (104F)

Inactive ingredients

Aqua

Package Label - Principal Display Panel



800wipes in onebox NDC: 79795-003-09

ALCOHOL WIPES (75% ALCOHOL)

alcohol cloth

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:79795-003
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958 V90M) (ALCOHOL - UNII:3K9958 V90M)	ALCOHOL	0.75 g

Inactive Ingredients

Ingredient Name			Strength	
WATER (UNII: 059QF0KO0R)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:79795-003-01	3 in 1 BAG; Type 2: Prefilled Drug Delivery Device/System (syringe, patch, etc.)	11/06/2020	
2	NDC:79795-003-02	5 in 1 BAG; Type 0: Not a Combination Product	11/06/2020	
3	NDC:79795-003-03	10 in 1 BAG; Type 0: Not a Combination Product	11/06/2020	
4	NDC:79795-003-04	20 in 1 BAG; Type 0: Not a Combination Product	11/06/2020	
5	NDC:79795-003-05	50 in 1 BAG; Type 0: Not a Combination Product	11/06/2020	
6	NDC:79795-003-06	80 in 1 BAG; Type 0: Not a Combination Product	11/06/2020	
7	NDC:79795-003-07	75 in 1 BOX; Type 0: Not a Combination Product	11/06/2020	
8	NDC:79795-003-08	80 in 1 BOX; Type 0: Not a Combination Product	11/06/2020	
9	NDC:79795-003-09	100 in 1 BOX; Type 0: Not a Combination Product	11/06/2020	
10	NDC:79795-003-10	150 in 1 BOX; Type 0: Not a Combination Product	11/06/2020	
11	NDC:79795-003-11	160 in 1 BOX; Type 0: Not a Combination Product	11/06/2020	
12	NDC:79795-003-12	200 in 1 BOX; Type 0: Not a Combination Product	11/06/2020	
13	NDC:79795-003-13	600 in 1 BOX; Type 0: Not a Combination Product	11/06/2020	
14	NDC:79795-003-14	800 in 1 BOX; Type 0: Not a Combination Product	11/06/2020	
15	NDC:79795-003-15	1000 in 1 BOX; Type 0: Not a Combination Product	11/06/2020	
16	NDC:79795-003-16	1200 in 1 BOX; Type 0: Not a Combination Product	11/06/2020	
Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC mono graph not final	part333A	11/06/2020		

Labeler - NanTong Guarder Medical Technology Co.,Ltd. (554537961)

Registrant - NanTong Guarder Medical Technology Co.,Ltd. (554537961)

Establishment

Name	Address	ID/FEI	Business Operations
NanTong Guarder Medical Technology Co.,Ltd.		554537961	manufacture(79795-003)

Revised: 11/2020

NanTong Guarder Medical Technology Co.,Ltd.