APOLLO MEDICAL PRODUCTS DISINFECTING WIPES- alcohol liquid
KKJ & Associates LLC

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.

Apollo Disinfecting Wipes

Active Ingredient(s)

Ethyl Alcohol 70% v/v.

Purpose

Antiseptic

Use

For first aid and for cleansing areas to reduce bacteria and viruses.

Warnings

For external use only. Flammable. Keep away from heat or flame. Store at room temperature 59F-86F (15C-30C)

Do not use

- on 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. POISON CONTROL (800)-222-1222

Directions

Clean the affected area by using wipe to apply and clean skin or surface. Let air dry.

Other information

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

Inactive ingredients

Water.

Dispose of wipe in trash after use. Do not flush. Tightly close lids between uses to retain moisture. Nonrefillable container. Do not reuse or refill this empty container. Offer empty containers for

recycling. If recycling is not available, discard container in trash.

Package Label - Principal Display Panel

APOLLO MEDICAL PRODUCTS DISINFECTING WIPES

alcohol liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:79642-003
Route of Administration	TOPICAL		

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

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

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:79642-003-01	1 in 1 CONTAINER	09/18/2020	
1		500 mL in 1 CONTAINER; 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	09/18/2020	

Labeler - KKJ & Associates LLC (057060784)

Establishment

Name	Address	ID/FEI	Business Operations
KKJ & Associates LLC		057060784	manufacture(79642-003)

Revised: 9/2020 KKJ & Associates LLC