75% ALCOHOL WIPES- 75% alcohol wipes cloth
Guangzhou Daieme Cosmetic 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.

Active Ingredient(s)

Alcohol 75% v/v. Purpose: Antiseptic

Purpose

Antiseptic, Hand Sanitizer Wipes

Use

Hand Sanitize Wipesr 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

- Place enough product on hands to cover all surfaces. Rub hands together until dry.
- Supervise children under 6 years of age when using this product to avoid swallowing.

Other information

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

Inactive ingredients

glycerin, water,polyurethane,aloe extract

Package Label - Principal Display Panel

80 WIPES NDC: 54237-500-01

75% ALCOHOL WIPES

75% alcohol wipes cloth

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:54237-500
Route of Administration	EXTRACORPOREAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	0.75 U in 1 U

Inactive Ingredients	
Ingredient Name	**Strength**
GLYCERIN (UNII: PDC6A3C0OX)	
POLYURETHANE-35 (NOT MORE THAN 500 MPA.S AT 40%) (UNII: Q2LKX89BE0)	
WATER (UNII: 059QF0KO0R)	
ALOE (UNII: V5VD430YW9)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:54237-500-01	80 U in 1 PACKAGE; Type 0: Not a Combination Product	03/30/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	03/30/2020	

Labeler - Guangzhou Daieme Cosmetic co,.Ltd (542359812)

Registrant - SDI INTERNATIONAL CORP (069936261)

Establishment

Name	Address	ID/FEI	Business Operations
Guangzhou Daieme Cosmetic Co,.Ltd		542359812	manufacture(54237-500)

Revised: 5/2020 Guangzhou Daieme Cosmetic co,.Ltd