SWEET CAREFOR ALCOHOL WIPES- alcohol cloth Imperial Palace Commodity(shenzhen)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.

DRUG FACTS

Active Ingredient(s)

• Alcohol 75% v/v.

Purpose

• Cleaning and disinfection

Uses

Packs

- Peel back lid and then label.
- Remove wipes as required.
- Replace lid and label to keep wipes moist.

Buckets

- Open the bucket lid from the top.
- Locate tear cut in the inner pouch, tear straight to open. Do not remove wipesroll from pouch.
- Pull first wipes from center of roll up through opening.
- Thread first wipe through dispensing nozzle in top of roll.
- Close the bucket lid and dispense the wipes as required.

Warnings

- For external use only.
- Store in cool and dry place.keep away from heat and direct sunlight.
- Avoid touching eyes. In case of eye contact, flush thoroughly with water.
- Avoid contact with broken skin.
- Don't flush into toilet.
- Stop use and ask a doctor if skin irritation develops.
- In case of accidental ingestion, seekprofessional assistance or contact a Poison Control Center immediately.

Directions

- Thoroughly moisten the skin or solid surface with the product and let it dry without wiping
- For children under 6, use only under adult supervision.
- Not recommended for infants.

Inactive ingredients

• Water, Glycerin

Package Label - Principal Display Panel







1020 mL NDC: 52489-001-03



3025 mL NDC: 52489-001-04



SWEET CAREFOR ALCOHOL WIPES alcohol cloth **Product Information Product Type** HUMAN OTC DRUG Item Code (Source) NDC:52489-001 TOPICAL **Route of Administration Active Ingredient/Active Moiety Ingredient Name Basis of Strength** Strength ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII: 3K9958V90M) ALCOHOL 75 mL in 100 mL **Inactive Ingredients** Ingredient Name Strength GLYCERIN (UNII: PDC6A3C0OX)

Packaging						
# Item Code	Package Description	Marketing Start Date	Marketing End Date			
1 NDC:52489-001-01	53 mL in 1 PACKAGE; Type 0: Not a Combination Product	04/30/2020				
2 NDC:52489-001-02	285 mL in 1 PACKAGE; Type 0: Not a Combination Product	04/30/2020				
3 NDC:52489-001-03	1020 mL in 1 PAIL; Type 0: Not a Combination Product	04/30/2020				
4 NDC:52489-001-04	3025 mL in 1 PAIL; Type 0: Not a Combination Product	04/30/2020				
Marketing Information						
	y Application Number or Monograph Citation	Marketing Start Date	Marketing End Date			
Marketing Catego						

Labeler - Imperial Palace Commodity(shenzhen)CO.,LTD (527796368)

Registrant - Imperial Palace Commodity(Dongguan)CO.,LTD (544367643)

Establishment

Name	Address	ID/FEI	Business Operations
Imperial Palace Commodity(Dongguan)CO.,LTD		544367643	manufacture(52489-001)

Revised: 4/2020

Imperial Palace Commodity(shenzhen)CO.,LTD