

HUMAN GUARD HAND SANITIZER WIPES- alcohol swab
DERMACOS KOREA

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 70% v/v. Purpose: Antimicrobial

Purpose

Antimicrobial, Hand Sanitizer

Use

To help reduce bacteria on the skin.

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

- Open package and apply wipe thoroughly to hands.
- Allow to dry without wiping.
- Children under 6 years of age should be supervised when using this product.
- Not recommended for infants.

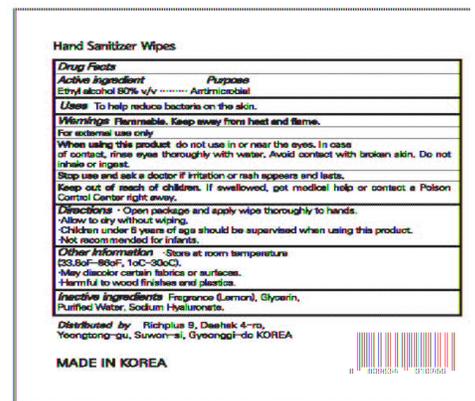
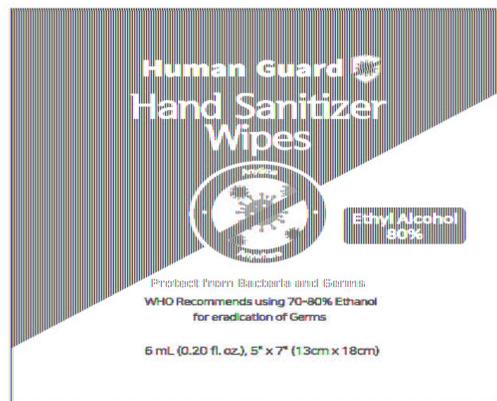
Other information

- Store at room temperature (33.8oF-86oF, 1oC-30oC) and avoid cold or dry place.
- May discolor certain fabrics or surfaces.
- Harmful to wood finishes and plastics.

Inactive ingredients

Fragrance (Lemon), Glycerin, Purified Water, Sodium Hyaluronate.

Package Label - Principal Display Panel



HUMAN GUARD HAND SANITIZER WIPES

alcohol swab

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:77637-200
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
GLYCERIN (UNII: PDC6A3C0OX)	
WATER (UNII: 059QF0KO0R)	

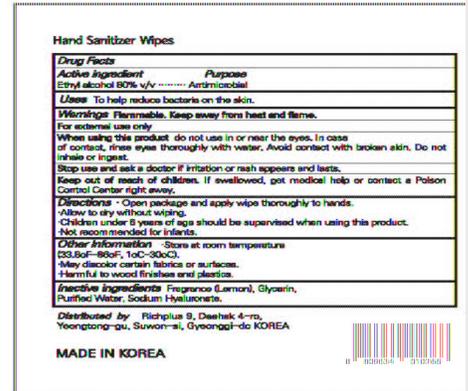
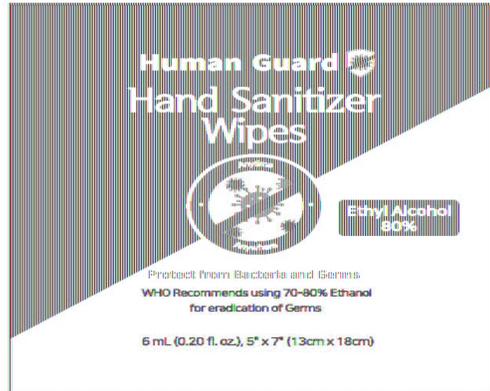
HYALURONATE SODIUM (UNII: YSE9PPT4TH)

Product Characteristics

Color		Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:77637-200-01	20 in 1 BOX	05/20/2020	
1		6 mL in 1 PACKET; 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	05/13/2020	

Labeler - DERMACOS KOREA (695625857)

Registrant - DERMACOS KOREA (695625857)

Establishment

Name	Address	ID/FEI	Business Operations
DERMACOS KOREA		695625857	manufacture(77637-200)

Revised: 5/2020

DERMACOS KOREA