HAND SANITIZER WIPES- alcohol cloth
HANUL CO.,LTD

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hand sanitizer wipes Hanul 75%

Active Ingredient(s)

Ethyl Alcohol 75% Purpose: Antiseptic

Purpose

Antiseptic, Hand Sanitizer Wipes

Use

To decrease bacteria on the skin that could cause disease

Warnings

For external use only: hands

Flammable. Keep away from fire and flame.

When using this product

keep out of eyes. In case of contact with eyes, flush thoroughly with water.

avoid contact with broken skin

do not inhale or ingest

Stop Use and ask a doctor if

irritation or redness develops

condition persists for more than 72 hours

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

Directions

- Wet hands thoroughly with product and allow to dry without wiping
- Not recommended for infants

Other information

do not store above 105ºF/40°C

may discolor some fabrics

harmful to wood finishes and plastics

Inactive ingredients

water, glycerin, hydrogen peroxide

Package Label - Principal Display Panel

rico

hand sanitizing wipes

non-sterile solution

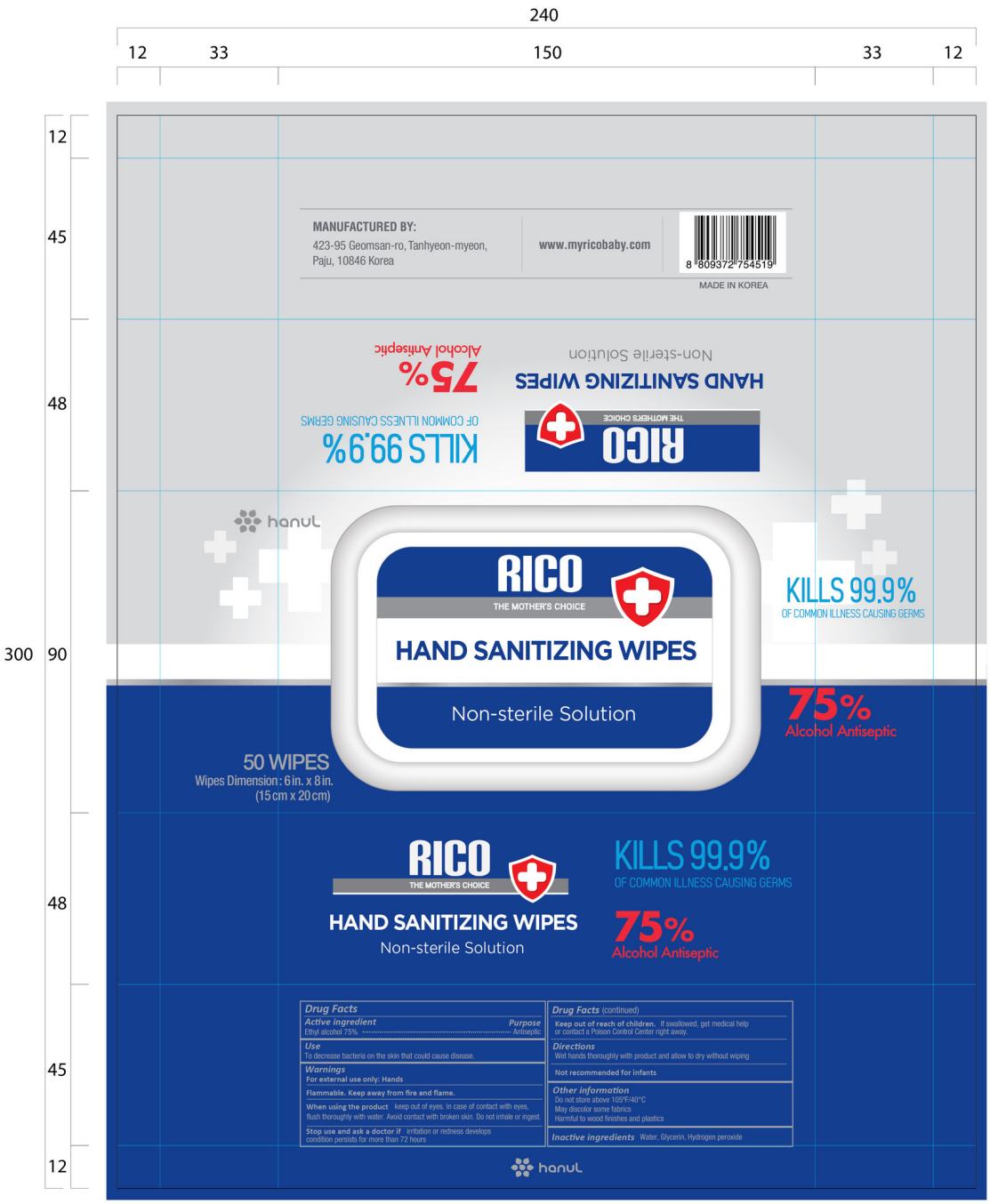
kills 99.9 of common illness-causing germs

75% alcohol antiseptic

RICO HAND SANITIZING WIPES (80), 75% Export

RICO HAND SANITIZING WIPES (10), 75% Export

RICO HAND SANITIZING WIPES (50), 75% Export



HAND SANITIZER WIPES

alcohol cloth

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:75214-503
Route of Administration	TOPICAL		

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)	1.45 mL in 100 mL
HYDROGEN PEROXIDE (UNII: BBX060AN9V)	0.125 mL in 100 mL
WATER (UNII: 059QF0KO0R)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:75214-503-50	334 mL in 1 PACKAGE; Type 0: Not a Combination Product	03/30/2020	
2	NDC:75214-503-10	134 mL in 1 PACKAGE; Type 0: Not a Combination Product	03/30/2020	
3	NDC:75214-503-80	534 mL 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 Drug	M016	03/30/2020	

Labeler - HANUL CO.,LTD (557814370)

Establishment

Name	Address	ID/FEI	Business Operations
HANUL CO.,LTD		557814370	manufacture(75214-503)

Revised: 1/2025 HANUL CO.,LTD