FIRST SAFETY HAND SANITIZING TISSUE- ethyl alcohol cloth
BonitoLab Co.

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

Ethy Alcohol 66% v/v

INACTIVE INGREDIENTS

Water, Glycerin, Citric Acid

PURPOSE

Antiseptic

WARNINGS

For external use only. Flammable. Keep away from heat or flame

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Do not use

■ on infants
■ on open skin wounds

When using this product

■ avoid getting into the eyes
■ In case of eye contact immediately flush eyes thoroughly with water

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Stop use and ask a doctor

■ if irritation or redness develops
■ conditions persist for more than 72 hours
■ redness is present

KEEP OUT OF REACH OF CHILDREN

■ in case of accidental ingestion, contact a doctor or Poison Center immediately

Uses

■ hand sanitizing wipes to help reduce bacteria on the skin

Directions

■ Tear open packet, remove wipe
■ thoroughly wipes hands with cloth
■ Supervise children under 6 years of age when using this product

Other Information

■ Storage in a cool, dry place, Avoid freezing and excessive heat above 40 □ (104 □)
■ may discolor certain fabrics or surfaces

PACKAGE LABEL.PRINCIPAL DISPLAY PANEL

FIRST SAFETY HAND SANITIZING TISSUE

ethyl alcohol cloth

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:79984-010
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
Alcohol (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	Alcohol	66 in 100

Inactive Ingredients	
Ingredient Name	**Strength**
Water (UNII: 059QF0KO0R)	
Glycerin (UNII: PDC6A3C0OX)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**

1	NDC:79984-010-01	30 in 1 CARTON; Type 0: Not a Combination Product	07/01/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	07/01/2020	

Labeler - BonitoLab Co. (695505454)

Registrant - BonitoLab Co. (695505454)

Establishment

Name	Address	ID/FEI	Business Operations
HANUL CO.,LTD		557814370	manufacture(79984-010)

Revised: 8/2020 BonitoLab Co.