COLABORATORY HAND SANITIZER- ethyl alcohol gel
Doehler USA Inc.

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.

CoLaboratory Hand Sanitizer

Drug Facts

Active ingredient

Ethyl Alcohol v/v 80%

Purpose

Antiseptic

USE

• Hand sanitizer to help reduce bacteria that could potentially cause disease. For use when soap and water are not readily available.

Warnings

Flammable. Keep away from fire or flame.

For external use only

When using this product
• Avoid contact with eyes. If contact occurs, flush eyes with water.
• Avoid contact with broken skin.

Stop using and consult doctor
if irritation or redness develop and last.

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

Directions.

• Wet hands with product & allow to dry without wiping
• For children under 6, use only under adult supervision.
• Not recommended for infants

Other information

• Store below 80°F (26.6°C)
• May discolor certain fabrics or surfaces

Inactive ingredients:

Fragrance, Glycerin, Hydrogen peroxide, Hydroxypropyl methylcellulose, Water

Alcohol antiseptic 80%

topical solution

non-sterile solution

Doehler North America
400 High Point Road SE
Cartersville, GA

for consumer complaints please call 770 387 0451
select the prompt to customer service

Packaging

COLABORATORY HAND SANITIZER

ethyl alcohol gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:77278-002
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	80 mL in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**
GLYCERIN (UNII: PDC6A3C0OX)	
HYDROGEN PEROXIDE (UNII: BBX060AN9V)	
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
WATER (UNII: 059QF0KO0R)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:77278-002-01	50 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/30/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	06/30/2020	

Labeler - Doehler USA Inc. (040455356)

Revised: 8/2020 Doehler USA Inc.