

HAND HERO HAND SANITIZER- ethyl alcohol gel
Natural Technology, LLC

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.

Ethyl Alcohol 70%. Purpose: Antimicrobial

3.79 L Label NDC: 79458-000-02

DRUG FACTS	
Active ingredient Ethyl Alcohol 70% v/v.....	Purpose Antibacterial
Use Hand sanitizer to help reduce bacteria on the skin	
Warnings Flammable Keep away from fire or flame Only for external use	
When using this product Avoid contact with the face and eyes. If you experience eye irritation, wash thoroughly with water and seek medical advice. Stop use and consult a doctor if irritation or redness keeps you out of reach of others. If swallowed, get medical attention and contact poison control right away.	
Directions - Wet hands and rub hands together until dry - Children under 6 years of age should be supervised by an adult when using this product	
Other information Store below 119°F (49°C)	
Inactive ingredients Purified water, Acrylates/Amlyl cresol/trimethylammonium chloride copolymer	
HAND HERO LLC PO BOX 78378 MIAMI, FL 33178	WWW.HANDHERO.COM 817264 MADE IN USA



**ANTIBACTERIAL
HAND
SANITIZER**
FRAGRANCE FREE

**KILLS 99.99%
OF GERMS***

128 FL OZ (3.79 L)



Purpose: Antimicrobial

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HAND HERO LLC NDC 79458-000-02 Rev. 04/19/2019	WWW.HANDHERO.COM H17004 MADE IN USA



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Avoid contact with the face and eyes.

If eye contact occurs, flush thoroughly with water and seek medical advice.

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HAND HERO LLC PO BOX 79174 DALLAS, TX 75279	www.usmhc.com 817264 MADE IN USA



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128 FL OZ (3.79 L)



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3.79L Label NDC: 79458-000-02

Purified Water, Acrylates/vinyl crosspolymer

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HAND HERO HAND SANITIZER

ethyl alcohol gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:79458-000
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	72 mL in 100 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	23 mL in 100 mL
RAPIDGEL EZ1 (UNII: 33JH4A7R2K)	5 mL in 100 mL

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:79458-000-02	3790 mL in 1 BOTTLE; Type 0: Not a Combination Product	07/14/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	07/14/2020	

Labeler - Natural Technology, LLC (618561906)

Registrant - Natural Technology, LLC (618561906)

Establishment

Name	Address	ID/FEI	Business Operations
Natural Technology, LLC		618561906	manufacture(79458-000)