

ADVANCED- ethyl alcohol gel
Perrigo Direct, Inc

Good Sense 370.001/370AE Rev 2
Advanced Hand Sanitizer

Active ingredient

Ethyl Alcohol 70%

Purpose

Antiseptic

Uses

- to decrease bacteria on the skin that could cause disease
- recommended for repeated use

Warnings

For external use only: hands

Flammable, keep away from fire or 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 and redness develop
- 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
- for children under 6, use only under adult supervision
- not recommended for infants

Other information

- do not store above 105°F
- may discolor some fabrics
- harmful to wood finishes and plastics

Inactive ingredients

water, glyceryl caprylate/caprinate, glycerin, isopropyl myristate, tocopheryl acetate, carbomer or acrylates/C10-30 alkyl acrylate crosspolymer, fragrance, benzophenone-4

Claims

*Effective at eliminating more than 99.99% of many common harmful germs and bacteria in as little as 15 seconds.

Disclaimer

**This product is not manufactured or distributed by GOJO Industries, Inc., distributor of Purell Advanced Hand Sanitizer.

ADVERSE REACTION

Distributed By: Perrigo Direct, Inc.

Peachtree City, GA 30269

www.PerrigoDirect.com

1-888-593-0593

GoodSense is a registered trademark of Perrigo Pharma International DAC.

Principal display panel

GOODSENSE®

Advanced

Hand Sanitizer

Fast & Effective

Kills more than 99.99% of germs*

Compare to active ingredient of Purell® Advanced**

100% SATISFACTION GUARANTEED

8 FL OZ (236 mL)

GOODSENSE.

Advanced Hand Sanitizer

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8 FL OZ (236 mL)

L0012668FC



Drug Facts

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Pat. 9,161,982
L0012668FC



46036-00068



ADVANCED

ethyl alcohol gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:75981-370
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
WATER (UNII: 059QF0KO0R)	
GLYCERYL CAPRYLATE/CAPRATE (UNII: G7515SW10N)	
GLYCERIN (UNII: PDC6A3COOX)	
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)	
CARBOMER COPOLYMER TYPE A (ALLYL PENTAERYTHRITOL CROSSLINKED) (UNII: 71DD5V995L)	
SULISOBENZONE (UNII: 1W6L629B4K)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date

1	NDC:75981-370-34	236 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	09/20/2012	
2	NDC:75981-370-16	59 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	09/20/2012	
3	NDC:75981-370-45	946 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	09/20/2012	06/01/2023

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	09/20/2012	

Labeler - Perrigo Direct, Inc (076059836)

Registrant - Nice-Pak Products, LLC (119091520)

Establishment

Name	Address	ID/FEI	Business Operations
Nice-Pak Products, LLC		119091514	manufacture(75981-370)

Establishment

Name	Address	ID/FEI	Business Operations
Nice-Pak Products, LLC		119091520	manufacture(75981-370)

Revised: 2/2026

Perrigo Direct, Inc