

HAND SANITIZER FRAGRANCE FREE- ethyl alcohol solution
Office Max

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.

70% Alcohol Hand Sanitizer
701.000/701AA

Active ingredient

Ethyl Alcohol 70% v/v

□ Purpose

Antiseptic

□ Uses

To decrease bacteria on the skin that could cause disease

Recommended for repeated use

□ Warnings

FLAMMABLE, keep away from fire and flame

For external use only

When using this product

do not use in or near eyes

Stop use and ask a doctor

if irritation or redness develops. If condition persists for more than 72 hours consult a doctor.

Keep out of reach of children.

If swallowed, seek immediate medical attention or call a poison control center.

□ Directions

Wet hands thoroughly with product and allow to dry without wiping.

□ *Inactive ingredients*

Water, Isopropyl Alcohol, PEG-12 Dimethicone, Glycerin, Caprylic/Capric Triglyceride, Isopropyl Myristate

Additional rear label text for manual dispensers

HighMark Manual Dispenser Foam Hand Sanitizer Refill

Distributed by: Office Depot, LLC

6600 N. Military Trail, Boca Raton, FL 33496 USA

Highmark is a trademark or registered trademark of OMX, Inc.

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Made in USA with foreign and domestic materials

Manufactured for Georgia-Pacific Consumer Products LP, Atlanta, GA 30303

Questions? Call 1-877-674-0688

Alcohol permit SDS-MO-15036 DSP-MO-28 DSP-MO-34

33.9 FL OZ (1000 mL)

Additional rear label text for automatic dispensers

HighMark Automatic Dispenser Foam Hand Sanitizer Refill

Distributed by: Office Depot, LLC

6600 N. Military Trail, Boca Raton, FL 33496 USA

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33.9 FL OZ (1000 mL)

principal display panel

MANUAL DISPENSER REFILL

FOAM HAND SANITIZER

FRAGRANCE FREE

33.9 fl oz (1000 mL)



principal display panel

AUTOMATED DISPENSER REFILL

FOAM HAND SANITIZER

FRAGRANCE FREE

33.9 FL OZ (1000 mL)



HAND SANITIZER FRAGRANCE FREE

ethyl alcohol solution

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:61139-701
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	70 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
ISOPROPYL ALCOHOL (UNII: ND2M416302)	
PEG-12 DIMETHICONE (UNII: ZEL54N6W95)	
GLYCERIN (UNII: PDC6A3C0OX)	

GLYCERYL CAPRYLATE/CAPRATE (UNII: G7515SW10N)

ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:61139-701-86	1000 mL in 1 BOTTLE, DISPENSING; Type 0: Not a Combination Product	10/06/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	06/02/2016	

Labeler - Office Max (009073099)

Registrant - Vi-Jon, LLC. (790752542)

Establishment

Name	Address	ID/FEI	Business Operations
Vi-Jon, LLC.		790752542	manufacture(61139-701)

Revised: 11/2022

Office Max