

**MEDISHIELD PRO ORIGINAL ANTISEPTIC HAND SANITIZER- alcohol gel
7327641 Canada, 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.

MediShield Pro Original Antiseptic Hand Sanitizer

Drug Facts

Active ingredient

Ethyl alcohol 70%

Purpose

Antiseptic

Uses

- For hand washing to decrease bacteria on the skin
- Recommended for repeated use

Warnings

For external use only

Flammable, keep away from fire or flame

Do not use

in the eyes

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

Other information

- Avoid temperatures over 110°F (43°C)

- May discolor certain fabrics
- Harmful to wood finishes and plastics

Inactive ingredients

Water, glycerin, carbomer, triethanolamine, fragrance, tocopheryl acetate, aloe vera powder

Questions?

1-877-874-0606, Weekdays, 9AM – 5PM EST

Package Labeling:

MEDISHIELD 

ORIGINAL

ANTISEPTIC HAND SANITIZER
70% ETHYL ALCOHOL

KILLS BACTERIA AND GERMS
NON-STICKY GEL

16.9 fl oz (1.06 pt) (500 mL)

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Questions? 1-877-874-0606, Weekdays, 9AM – 5PM EST	

Use without water.
Made in Canada

Manufactured by
7327641 Canada, Inc.
DBA Protair-X Solutions Group Supply Services
149-1 J.-A Bombardier
Boucherville, QC J4B 8P1
Canada

Medishield is a trademark of
Protair-X Solutions Group Supply Services

Protair-X Solutions Group Supply Services,
Boucherville, QC, J4B 8P1, Canada.
www.protairxgrp.com


NVA_410_01 Product Code: 1380-500

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MEDISHIELD PRO ORIGINAL ANTISEPTIC HAND SANITIZER

alcohol gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:81818-001
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
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ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)		ALCOHOL	70 mL in 100 mL	
Inactive Ingredients				
Ingredient Name				Strength
WATER (UNII: 059QF0K00R)				
GLYCERIN (UNII: PDC6A3C00X)				
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)				
TROLAMINE (UNII: 9O3K93S3TK)				
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)				
ALOE VERA LEAF (UNII: ZY81Z83H0X)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:81818-001-00	500 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/15/2021	
Marketing Information				
Marketing Category	Application Number or Monograph Citation		Marketing Start Date	Marketing End Date
OTC monograph not final	part333E		04/15/2021	

Labeler - 7327641 Canada, Inc. (204236330)

Revised: 4/2021

7327641 Canada, Inc.