

AMOVEO UNSCENTED- alcohol gel
Protair-X Health Solutions 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.

Amoveo Unscented

Drug Facts

Active Ingredient

Alcohol 70%

Purpose

Antiseptic

Uses

- For handwashing 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
- may discolor certain fabrics
- harmful to wood finishes and plastics

Inactive Ingredients

Carbomer, Glycerin, Triethanolamine, Water

Questions?

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

Package Labeling:



amoveo™

UNSCENTED

**ANTISEPTIC HAND SANITIZER
ALCOHOL 70%**

**Fast-acting
Hypoallergenic Gel**

**KILLS BACTERIA AND GERMS
NON-STICKY GEL**

33.8 fl oz (1.06 qt) (1 L)

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Supervise children in use of this product.
Use without water.

Made in Canada

AMOVEO is a trademark of Protair-X Health Solutions Inc. 

Protair-X Health Solutions Inc.,
Boucherville, QC, J4B 8P1, Canada.
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NVA_400_01 Product Code 1281-100



AMOVEO UNSCENTED

alcohol gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:49955-702
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	700 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
CARBOXYPOLYMETHYLENE (UNII: 0A5MM307FC)	
GLYCERIN (UNII: PDC6A3C0OX)	
TROLAMINE (UNII: 9O3K93S3TK)	
WATER (UNII: 059QF0K00R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:49955-702-50	500 mL in 1 BOTTLE; Type 0: Not a Combination Product	02/24/2017	
2	NDC:49955-702-01	1000 mL in 1 BOTTLE; Type 0: Not a Combination Product	02/24/2017	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	02/24/2017	

Labeler - Protair-X Health Solutions Inc. (205614519)

Revised: 2/2017

Protair-X Health Solutions Inc.