

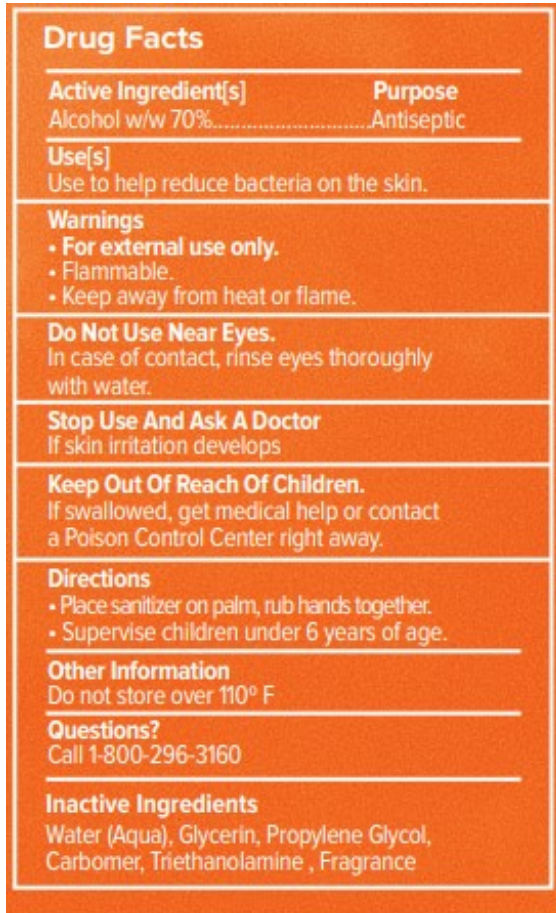
MOXE- citrus hand sanitizer 4oz gel
Nutrix International, LLC.

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.

Active Ingredient[s]

Alcohol 70%

Use[s]



Drug Facts	
Active Ingredient[s]	Purpose
Alcohol w/w 70%	Antiseptic
Use[s]	
Use to help reduce bacteria on the skin.	
Warnings	
• For external use only.	
• Flammable.	
• Keep away from heat or flame.	
Do Not Use Near Eyes.	
In case of contact, rinse eyes thoroughly with water.	
Stop Use And Ask A Doctor	
If skin irritation develops	
Keep Out Of Reach Of Children.	
If swallowed, get medical help or contact a Poison Control Center right away.	
Directions	
• Place sanitizer on palm, rub hands together.	
• Supervise children under 6 years of age.	
Other Information	
Do not store over 110° F	
Questions?	
Call 1-800-296-3160	
Inactive Ingredients	
Water (Aqua), Glycerin, Propylene Glycol, Carbomer, Triethanolamine, Fragrance	

Use to help reduce bacteria on the skin.

Warnings

- **For external use only.**
- Flammable.
- Keep away from heat or flame.

Do not use

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Stop Use And Ask

A Doctor If skin irritation develops

Keep out of reach of children

If swallowed, get medical help or contact a Poison Control Center right away

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Other Information

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Purpose

Antiseptic

Artwork

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Label

MOXE			
citrus hand sanitizer 4oz gel			
Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:73333-011
Route of Administration	TOPICAL		
Active Ingredient/Active Moiety			
	Ingredient Name	Basis of Strength	Strength
	ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	0.7 g in 118 g
Inactive Ingredients			
	Ingredient Name	Strength	
	TANGERINE (UNII: KH3E3096OO)	0.005 g in 118 g	
	TROLAMINE (UNII: 9O3K93S3TK)	0.0012 g in 118 g	
	CARBOMER 940 (UNII: 4Q93RCW27E)	0.004 g in 118 g	
	PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	0.015 g in 118 g	
	WATER (UNII: 059QF0KO0R)	0.2598 g in 118 g	
	GLYCERIN (UNII: PDC6A3C0OX)	0.015 g in 118 g	
Packaging			

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:73333-011-01	118 g in 1 BOTTLE; Type 0: Not a Combination Product	05/11/2020	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	05/11/2020	

Labeler - Nutrix International, LLC. (117341868)

Establishment			
Name	Address	ID/FEI	Business Operations
Nutrix International, LLC.		117341868	manufacture(73333-011) , pack(73333-011) , label(73333-011)

Revised: 5/2020

Nutrix International, LLC.