AQ PURE INSTANT HAND SANITIZER- alcohol gel DAY GREAT 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.

AQ pure Instant Hand Sanitizer

Drug Facts

Active Ingredients

Alcohol 80%

Purpose

Antiseptic

Uses:

- wet hands bacteria on the skin that could disease
- recommended for repeated uses

Directions:

- wet hands thoroughly with product and allow to dry without wiping
- not recommende for infants

Warnings:

- For external use only
- Flammable
- Keep away from heat and flame
- Do not store above 40 degrees Celcius (15 degrees Farenheit)

Do not use

• in mouth, ears or eyes

When using this product,

- avoid contact with eyes
- In case of contact, flush eyes with water

Stop use and seek a doctor

- if redness or irritation develops and persist for more than 72 hours
- Children must be supervise in use of this product
- Pump as needed into your palms and thoroughly spread on both hands
- Rub in to skin until dry

Keep out of reach of children

• children must be supervised in the use of this product

Other Information:

- may discolor some fabrics
- harmful to wood finishes and plastics

Inactive Ingredients:

- water
- triethanolamine
- acrylates/c 10-30 alkyl crosspolymer
- glycerin

Package Labeling:290ml



9.8 FL OZ (290ml) NDC: xxxxx-xxx-xx

Drug Facts

l lene

• to decrease bacteria on the skin that could cause disease • recommended for repeated uses

Directions

• wet hands thoroughly with product and allow to dry without wiping • not recommended for infants

Warnings:

• For external use only • Flammable • Keep away from heat and flame • Do not store above 40 degrees Celcius (105 degrees Farenheit) • Do not use in mouth, ears or eyes • When using this product, avoid contact with eyes • In case of contact, flush eyes with water • Stop use and seek a doctor if redness or irritation develops and persist for more than 72 hours • Children must be supervised in use of this product • Pump as needed into your palms and thoroughly spread on both hands • Rub into skin until dry • children must be supervised in the use of this product

Other Information:

• may discolor some fabrics • harmful to wood finishes and plastics

Inactive Ingredients:

• water • triethanolamine • acrylates/c10-30 alkyl acrylate crosspolymer • glycerin

Distributed by: Day Great LLC, Miami, FL 33181











Package Labeling:480ml

er Information:

may discolor some fabrics • harmful to wood finishes and plastics

Inactive Ingredients:

• water • triethanolamine • acrylates/c10-30 alkyl acrylate crosspolymer • glycerin

Distributed by: Day Great LLC , Miami, FL 33181





Z (480ml)

Drug Facts

• to de

to decrease bacteria on the skin that could cause disease * recommended for repeated uses

Directions:

 \bullet wet hands thoroughly with product and allow to dry without wiping \bullet not recommended for infants

Warnings:

• For external use only • Tammable • Keep away from heat and flame • Do not store above 40 degrees Celcius (105 degrees Farenheit) • Do not use in mouth, each greys • When using this product, avoid contact with eyes • In case of contact, flush eyes with water • Stop use and seek a doctor if redness or irritation develops and persist for more than 72 hours • Children must be supervised in use of this product • Pump as needed into your palms and thoroughly spread on both hands • Rub into skin until dry • Keep out of reach of children

AQ PURE INSTANT HAND SANITIZER

alcohol gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:79333-001
Route of Administration	TOPICAL		

l	Active Ingredient/Active Moiety		
l	Ingredient Name	Basis of Strength	Strength
ı	ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	0.8 mL in 1 mL

Inactive Ingredients	
Ingredient Name	Strength
CARBOMER INTERPOLYMER TYPE A (ALLYL SUCROSE CROSSLINKED) (UNII: 59TL3WG5CO)	
WATER (UNII: 059QF0KO0R)	
TROLAMINE (UNII: 9O3K93S3TK)	
GLYCERIN (UNII: PDC6 A3C0 OX)	

l	Packaging			
l	# Item Code	Package Description	Marketing Start Date	Marketing End Date
l	1 NDC:79333-001-07	290 mL in 1 BOTTLE; Type 0: Not a Combination Product	07/13/2020	
l	2 NDC:79333-001-08	480 mL in 1 BOTTLE; Type 0: Not a Combination Product	07/13/2020	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	07/13/2020	

Labeler - DAY GREAT LLC (122472915)

Revised: 7/2020 DAY GREAT LLC