80% ALCOHOL GEL- alcohol solution Phoenix Products Company

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 80% v/v. Purpose: Antiseptic

Purpose

Antiseptic, Hand Rub

Use

Health care personnel hand rub to help reduce bacteria that potentially can cause disease.

Warnings

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

Do not use

- in children less than 2 months of age
- on open skin wounds

When using this product keep out of eyes, ears, and mouth. In case of contact with eyes, rinse eyes thoroughly with water.

Stop use and ask a doctor if irritation or rash occurs. These may be signs of a serious condition.

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

Directions

- Place enough product on hands to cover all surfaces. Rub hands together until dry.
- Supervise children under 6 years of age when using this product to avoid swallowing.

Other information

- Store between 15-30C (59-86F)
- Avoid freezing and excessive heat above 40C (104F)

Inactive ingredients

purified water, glycerin USP, hydrogen peroxide, carbomer, TEA.

Package Label - Principal Display Panel

[Divina 16 fl oz]

NDC: 73956-294-16

MADE IN USA Divina Professional Products Terryville, CT 06786 www.divinaproducts.com

HAND SANITIZER Alcohol Antiseptic 80% Topical Solution Antiseptic Hand Rub Non-Sterile Solution 16 fl oz. | 473 mL

Since 1969 Divina has been manufacturing premium quality salon-use solutions that professionals rely on every day to provide superior results and happy clients. Product integrity and value are the cornerstones of our business.

Professional Salon Solutions

OVER 50 YEARS

Members of the Professional Beauty Association

DIRECTIONS FOR USE:

- 1. If possible, first thoroughly wash hands with soap and water for at least 20 seconds.
- 2. Apply enough of the product to the palm of your hand to wet your hands completely.
- 3. Rub your hands and fingers together to completely cover all areas until dry. Keep away from EYES or any mucous membranes.

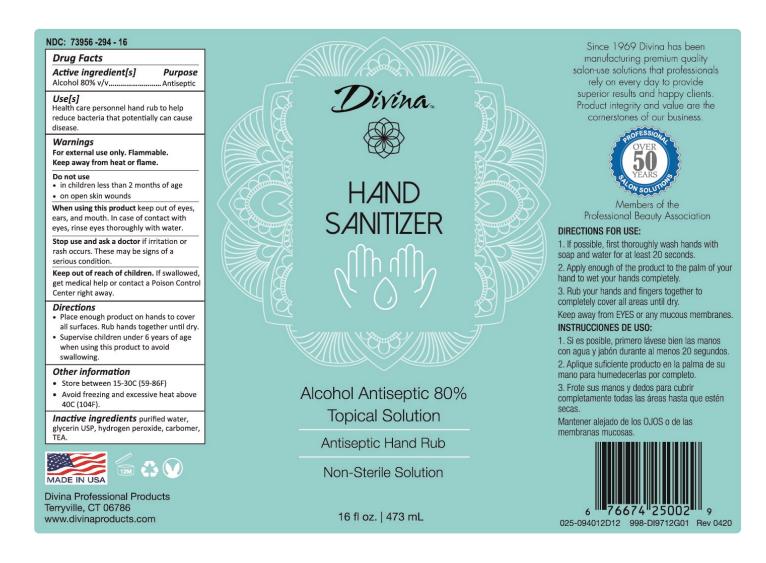
INSTRUCCIONES DE USO:

- 1. Si es posible, primero lávese bien las manos con agua y jabón durante al menos 20 segundos.
- 2. Aplique suficiente producto en la palma de su mano para humedecerlas por completo.
- 3. Frote sus manos y dedos para cubrir completamente todas las áreas hasta que estén secas.

Mantener alejado de los OJOS o de las membranas mucosas.

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025-094012D12 998-DI9712G01 Rev 0420



Package Label - Principal Display Panel

[Divina 32 fl oz]

NDC: 73956-294-32

MADE IN USA Divina Professional Products Terryville, CT 06786 www.divinaproducts.com

HAND SANITIZER Alcohol Antiseptic 80% Topical Solution Antiseptic Hand Rub Non-Sterile Solution 32 fl oz. | 946 mL

Since 1969 Divina has been manufacturing premium quality salon-use solutions that professionals rely on every day to provide superior results and happy clients. Product integrity and value are the cornerstones of our business.

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DIRECTIONS FOR USE:

- 1. If possible, first thoroughly wash hands with soap and water for at least 20 seconds.
- 2. Apply enough of the product to the palm of your hand to wet your hands completely.
- 3. Rub your hands and fingers together to completely cover all areas until dry. Keep away from EYES or any mucous membranes.

INSTRUCCIONES DE USO:

- 1. Si es posible, primero lávese bien las manos con agua y jabón durante al menos 20 segundos.
- 2. Aplique suficiente producto en la palma de su mano para humedecerlas por completo.
- 3. Frote sus manos y dedos para cubrir completamente todas las áreas hasta que estén secas.

Mantener alejado de los OJOS o de las membranas mucosas.

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025-094013D06 998-DI9713G01 Rev 0420



[Phoenix Gallon]

NDC: 73956-194-28

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Phoenix Problem Solving Products

HAND

SANITIZER

Alcohol Antiseptic 80% Topical Solution Antiseptic Hand Rub - Non-Sterile Solution

3.79 Liters (1 Gallon)

DIRECTIONS FOR USE:

- 1. If possible, first thoroughly wash hands with soap and water for at least 20 seconds.
- 2. Apply enough of the product to the palm of your hand to wet your hands completely.
- 3. Rub your hands and fingers together to completely cover all areas until dry. Keep away from EYES or any mucous membranes.

INSTRUCCIONES DE USO:

- 1. Si es posible, primero lávese bien las manos con agua y jabón durante al menos 20 segundos.
- 2. Aplique suficiente producto en la palma de su mano para humedecerlas por completo.
- 3. Frote sus manos y dedos para cubrir completamente todas las áreas hasta que estén secas.

Mantener alejado de los OJOS o de las membranas mucosas.

Distributed by:

Phoenix Products Company

55 Container Drive, Terryville, CT 06786

Rev. No 0320

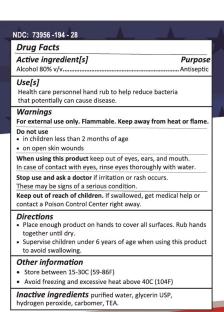
Item# 001-094015P04

998-PHHS15G01

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www.phoenixproductsco.com





Alcohol Antiseptic 80% **Topical Solution**

Antiseptic hand rub - Non-sterile solution

DIRECTIONS FOR USE:

- 1. If possible first thoroughly wash hands with soap and water for at least 20 seconds.
- 2. Apply enough of the product to the palm of your hand to wet your hands completely.
- 3. Rub your hands and fingers together to completely cover

Keep away from EYES or any mucous membranes.

INSTRUCCIONES DE USO:

- 1. Si es posible, primero lávese bien las manos con agua y jabón durante al menos 20 segundos.
- 2. Aplique suficiente producto en la palma de su mano para humedecerlas por completo.
- 3. Frote sus manos y dedos para cubrir completamente todas las áreas hasta que estén secas.

Mantener alejado de los OJOS o de las membranas

For further information on the chemicals used in this formula. review the MSDS.

Distributed by:
Phoenix Products Company
55 Container, Drive Terryville, CT 06786
Rev. No. 03.20
Item # 001-094015P04
998-PHST 5601
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www.phoenixproductsco.com



3.79 Liters (1 Gallon)

80% ALCOHOL GEL

alcohol solution

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:73956-294	
Route of Administration	TOPICAL			

l	Active Ingredient/Active Moiety				
l	Ingredient Name	Basis of Strength	Strength		
ı	ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	80 mL in 100 mL		

Inactive Ingredients				
Ingredient Name	Strength			
GLYCERIN (UNII: PDC6A3C0OX)	.5 mL in 100 mL			
HYDROGEN PERO XIDE (UNII: BBX060AN9V)	0.25 mL in 100 mL			
WATER (UNII: 059QF0KO0R)	18.35 mL in 100 mL			
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0 A5MM307FC)	.5 mL in 100 mL			
TRIETHANOLAMINE LAURYL SULFATE (UNII: E8458C1KAA)	.3 mL in 100 mL			

P	Packaging					
#	Item Code	Package Description	Marketing Start Date	Marketing End Date		
1	NDC:73956-294-16	473 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/01/2020			
2	NDC:73956-294-32	946 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/01/2020			
3	NDC:73956-294-28	3785 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/01/2020			

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	06/01/2020	

80% ALCOHOL GEL

alcohol solution

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:73956-194	
Route of Administration	TOPICAL			

Active Ingredient/Active Moiety		
Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	80 mL in 100 mL

Inactive Ingredients			
Ingredient Name	Strength		
GLYCERIN (UNII: PDC6A3C0OX)	.5 mL in 100 mL		
HYDRO GEN PERO XIDE (UNII: BBX060AN9V)	0.25 mL in 100 mL		
WATER (UNII: 059QF0KO0R)	17.6 mL in 100 mL		
TRIETHANO LAMINE LAURYL SULFATE (UNII: E8458 C1KAA) .3 mL in 100 mL			
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0 A5MM307FC)	.5 mL in 100 mL		

P	Packaging					
#	Item Code	Package Description	Marketing Start Date	Marketing End Date		
1	NDC:73956-194-04	118 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/01/2020			
2	NDC:73956-194-16	473 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/01/2020			
3	NDC:73956-194-32	946 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/01/2020			
4	NDC:73956-194-28	3785 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/01/2020			

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph not final	part333A	06/01/2020		

Labeler - Phoenix Products Company (101332237)

Establishment			
Name	Address	ID/FEI	Business Operations

Phoenix Products Company	10	1332237	manufacture(73956-194, 73956-294)

Revised: 8/2020 Phoenix Products Company