## BQ PHARMA HAND SANITIZER- alcohol gel BQ PHARMA HAND SANITIZER FRESH SCENT- alcohol gel Frank por Laboratories, S.A. de C.V.

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.

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#### **BQ Pharma Hand Sanitizer 70 Percent**

#### DRUG FACTS

## Active Ingredient(s)

Alcohol 70% v/v

## **Purpose**

Antiseptic

## Use(s)

Hand Sanitizer to help reduce bacteria that potentially can cause disease. For use when soap and water are not available.

## Warnings

For external use only. Flammable. Keep away from jeat 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. I 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 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

- Do not store above 105°F (40°C)
- May discolor certain wood surfaces.

## **Inactive Ingredients**

Camphor, Carbomer, Glycerin, Menthol, Triethanolamine, Water.

## Questions or comments?

#-###-###

\*Mix of mostiurizers that will keep your hands moisturized for up to 24 hours.

Made in Mexico by Frank Por Laboratories S.A. de C.V.

Distributed by XXXXX

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## **Inactive Ingredients**

Carbomer, C.I. 42090, Fragrance, Glycerin, Menthol, Triethanolamine, Water.

#### Questions or comments?

#-###-###

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#### PRINCIPAL DISPLAY PANEL

BQ Pharma Hand Sanitizer 70 % Ethyl Alcohol 3.04 fl oz.



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# Drug Facts (continued)

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#### PRINCIPAL DISPLAY PANEL

BQ PHARMA HAND SANITIZER 70 % ETHYL ALCOHOL 13.52 fl Oz.





# **BQ PHARMA HAND SANITIZER**

alcohol gel

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:78817-001
Route of Administration	TOPICAL		

l	Active Ingredient/Active Moiety		
l	Ingredient Name	Basis of Strength	Strength
l	ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	.70 mL in 1 mL

Inactive Ingredients				
Ingredient Name	Strength			
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0 A5MM307FC)				
CAMPHOR (SYNTHETIC) (UNII: 5TJD82A1ET)				
GLYCERIN (UNII: PDC6A3C0OX)				
MENTHOL (UNII: L7T10EIP3A)				
TROLAMINE (UNII: 9O3K93S3TK)				
WATER (UNII: 059QF0KO0R)				

F	Packaging				
#	Item Code	Package Description	<b>Marketing Start Date</b>	Marketing End Date	
1	NDC:78817-001-01	90 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/08/2020		
2	NDC:78817-001-02	180 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/08/2020		
3	NDC:78817-001-03	1000 mL in 1 BOTTLE; Type 0: Not a Combination Product	09/15/2020		
4	NDC:78817-001-04	400 mL in 1 BOTTLE; Type 0: Not a Combination Product	09/15/2020		
5	NDC:78817-001-05	400 mL in 1 BOTTLE; Type 0: Not a Combination Product	09/15/2020		
6	NDC:78817-001-06	500 mL in 1 BOTTLE; Type 0: Not a Combination Product	09/15/2020		

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC MONOGRAPH NOT FINAL	part333E	06/08/2020		

# BQ PHARMA HAND SANITIZER FRESH SCENT

alcohol gel

<b>Product Information</b>			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:78817-002
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	.70 mL in 1 mL	

Inactive Ingredients				
Ingredient Name	Strength			
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0 A5MM307FC)				
CAMPHOR (SYNTHETIC) (UNII: 5TJD82A1ET)				
GLYCERIN (UNII: PDC6A3C0OX)				
MENTHOL (UNII: L7T10EIP3A)				
TROLAMINE (UNII: 9O3K93S3TK)				
WATER (UNII: 059QF0KO0R)				

]	Packaging			
#	# Item Code	Package Description	<b>Marketing Start Date</b>	Marketing End Date
1	NDC:78817-002-03	1000 mL in 1 BOTTLE; Type 0: Not a Combination Product	09/15/2020	
2	NDC:78817-002-04	400 mL in 1 BOTTLE; Type 0: Not a Combination Product	09/15/2020	
3	NDC:78817-002-05	400 mL in 1 BOTTLE; Type 0: Not a Combination Product	09/15/2020	
4	NDC:78817-002-06	500 mL in 1 BOTTLE; Type 0: Not a Combination Product	09/15/2020	

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC MONOGRAPH NOT FINAL	part333E	09/15/2020		

# Labeler - Frank por Laboratories, S.A. de C.V. (812756987)

Revised: 8/2020 Frank por Laboratories, S.A. de C.V.