BUFFERIN HAND SANITIZER- alcohol gel Genomma Lab USA

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.

Bufferin® Hand Sanitizer

Drug Facts

Active Ingredient

Ethyl alcohol 70% v/v

Purpose

Antiseptic

Uses

- To decrease bacteria on the skin when water, soap & towel are not available
- Recommended for repeated use.

Warnings

For external use only.

Flammable. Keep away from fire or flame.

When using this product

- Keep out of eyes. In case of contact with eyes, rinse thoroughly with water
- Do not use on broken or irritated skin.

Stop use and ask a doctor if irritation or redness develop and last more than 72 hours.

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

Directions

• Apply enough product to wet hands. Rub hands together until dry. Supervise children in use of this product.

Other information

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

Inactive ingredients

water, carbomer, triethanolamine, fragrance, propylene glycol, glycerin, aloe barbadensis extract, tocopheryl acetate.

Questions?

Call toll free 1-877-994-3666 Monday to Friday, 8am to 6pm, Central time

Distributed by: Genomma Lab USA

Inc., Houston, TX, 77027

PRINCIPAL DISPLAY PANEL - 236 ml Bottle Label

The Science of Healthy Hands

70% ETHYL ALCOHOL SOURCED IN THE USA

BUFFERIN® HAND SANITIZER

Antibacterial Gel

KILLS 99.99% OF BACTERIA

enriched with Vitamin E & Aloe

8 fl oz (236 ml)

2000009663 S-20N092 The Science of Healthy Hands



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Made in Mexico.
Distributed by: Genomma Lab USA
Inc., Houston, TX, 77027
www.genommalab.us

Lot number and expiration date printed on bottle.

BUFFERIN HAND SANITIZER

alcohol gel

Product Information Product Type HUMAN OTC DRUG Item Code (Source) NDC:50066-608 Route of Administration TOPICAL

Active Ingredient/Active Moiety	e Ingredient/Active Moiety				
Ingredient Name	Basis of Strength	Strength			
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	70 mL in 100 mL			

Inactive Ingredients

Strength				
CARBOMER HO MO PO LYMER, UNSPECIFIED TYPE (UNII: 0 A5MM307FC)				

ı	P	Packaging			
	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
		NDC:50066-608- 08	236 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	08/12/2020	
		NDC:50066-608- 02	70 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	09/16/2020	

Marketing Informat	rketing Information					
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date			
OTC MONOGRAPH NOT FINAL	part333E	08/12/2020				

Labeler - Genomma Lab USA (832323534)

Revised: 9/2020 Genomma Lab USA