DR GEORGES VIRUBLAST- ethyl alcohol gel
M&M Innovations Inc

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.

Dr Georges Virublast

Active ingredient

Ethyl Alcohol 75% v/v

Purpose

Handwash, antiseptic

Use

For handwashing to decrease bacteria on the skin.

Warnings

Flammable. Keep away from fire or flame.

For external use only.

Do not use

Do not use in or near the eyes, in case of contact rinse eyes thoroughly with water

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

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

Directions

- wet hands thoroughly with product
- briskly rub hands together until dry
- supervise children in the use of this product

Other information

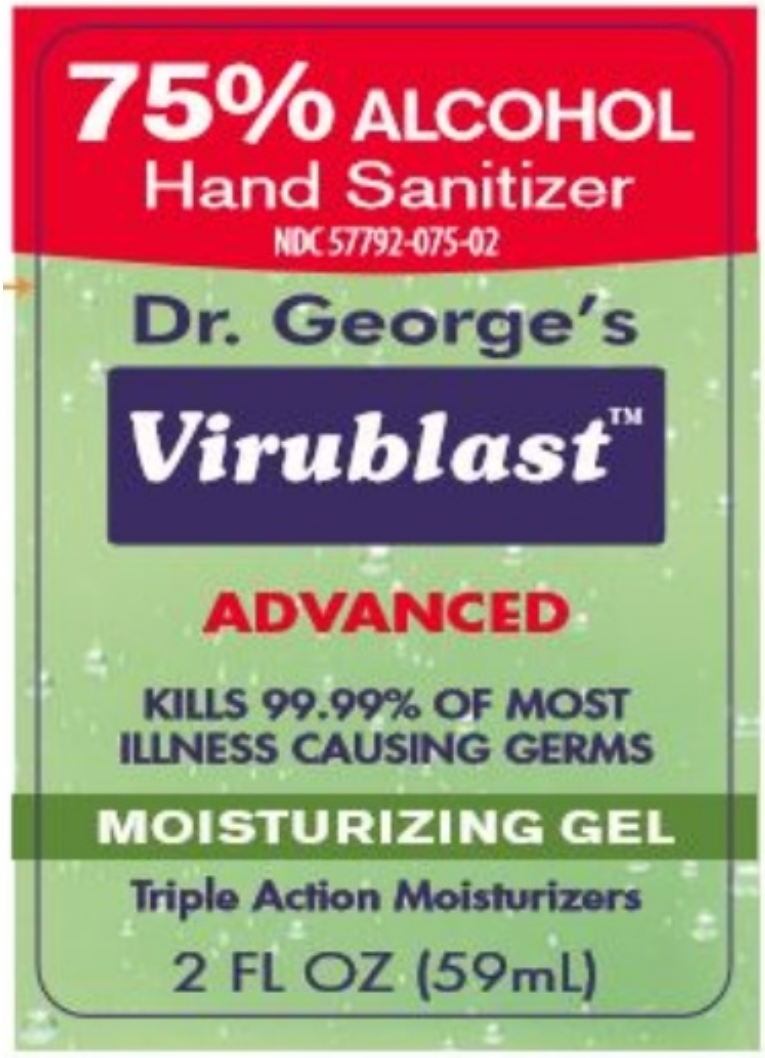
- store below 110F or 43C
- may discolor certain fabrics or surfaces

Inactive ingredients

water, isopropyl alcohol, aloe barbadensis leaf juice, caprylyl glycol, glyercin, isopropyl myristate, tocopheryl acetate, acrylates/c-10-30 alkyl acrylate crosspolymer, aminomethyl propanol, fragrance

Principal display panel

2 fl oz (59 ml)



DR GEORGES VIRUBLAST

ethyl alcohol gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:57792-075
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	75 mL in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**
CARBOMER INTERPOLYMER TYPE A (55000 CPS) (UNII: 59TL3WG5CO)	
CAPRYLYL GLYCOL (UNII: 00YIU5438U)	
WATER (UNII: 059QF0KO0R)	
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)	

.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
GLYCERIN (UNII: PDC6A3C0OX)	
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)	
ISOPROPYL ALCOHOL (UNII: ND2M416302)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:57792-075-02	59 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/01/2020	
2	NDC:57792-075-67	66 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/01/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	05/01/2020	

Labeler - M&M Innovations Inc (614929743)

Establishment

Name	Address	ID/FEI	Business Operations
M&M Innovations Inc		614929743	manufacture(57792-075)

Revised: 8/2020 M&M Innovations Inc