ACTION INSTANT HAND SANITIZER- alcohol gel MAESTRO LABORATORIES SDN. BHD.

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.

Action Instant Hand Sanitizer

Drug Facts

Active Ingredient:

Alcohol Denat 75%

Purpose

Antimicrobial

Uses:

- Hand sanitizer helps to reduce bacteria on the skin that could cause disease
- Recommended for repeated use

WARNINGS:

FOR EXTERNAL USE ONLY

• Flammable. Keep away from fire or flame.

When using this product

- do not use in or near eyes.
- In case of contact, rinse eyes thoroughly with water.

Stop use and ask a doctor

if irritation or rash appears and lasts

Keep out of reach of children.

• If swallowed, get medical help or contact a Poison Control Center right away.

Direction Of Use:

Squeeze amount in your palm then briskly rub hands together until dry.

Children under 3 years of age should be supervised when using this product

Other Information:

Store below 110F (43C)

Inactive Ingredients:

Aqua, Glycerin, Acrylates/C10-30 Alkyl Acrylate Cross-polymer, Aminomethyl Propanol, Aloe Barbadensis Leaf Juice, Panthenol, Tocopheryl Acetate, Fragrance.

Questions or comments?

Call Customer Care Line 1300 88 4848

Package Labeling:









46x120mm

03-07-2020

ACTION INSTANT HAND SANITIZER

alcohol gel

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:80613-000

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name Basis of Strength Strength

ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M) ALCOHOL 0.75 mL in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
CARBOMER INTERPOLYMER TYPE A (ALLYL SUCROSE CROSSLINKED) (UNII: 59TL3WG5CO)	
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)	
ALOE VERA LEAF (UNII: ZY8 1Z8 3H0 X)	
PANTHENOL (UNII: WV9CM0O67Z)	
.ALPHATO COPHEROL ACETATE (UNII: 9E8X80D2L0)	

Packaging

l	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
ı	1	NDC:80613-000-00	500 mL in 1 BOTTLE; Type 0: Not a Combination Product	10/01/2019	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	10/01/2019	
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