# INSTANT HAND SANITIZING WIPE- ethyl alcohol swab McKesson

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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# **Instant Hand Sanitizing Wipe**

# **Drug Facts**

# Active ingredient

Ethyl Alcohol 65.9% by volume

# **Purpose**

**Antiseptic** 

#### Uses

- For handwashing to decrease bacteria on the skin.
- For sanitizing hands after assisting ill persons, or Prior to contacting with a person under medical care or treatment
- Topical Application to skin to help prevent cross-contamination.

# Warnings

Flammable. Keep away from fire, flame or source of ignition i.e. electrocautery procedures.

For external use only.

Do not use in or contact with eyes.

Discontinue use if irritation and redness develop. If condition persists more than 72 hours, consult a doctor.

**Keep out of reach of children** unless under adult supervision.

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

#### **Directions**

To dispense, remove cover, discard seal from container, pull center sheet from roll, twist wipe to a point and thread through dispenser hole in cover.

Replace cover. Pull out wipe and snap off at 90 degree angle. The next wipe is ready for dispensing. Apply wipe thoroughly onto hands and allow to dry.

No rinsing required. Discard after single use. Dispose of used wipes in trash receptacle. Do not flush. When not in use, keep lid closed to prevent evaporation.

#### Other information

Lot No. and Expiration Date can be found on canister. Store at 53.6 - 93.2\*F (12-34\*C)

# Inactive ingredients

Aminomethyl Propanol, Aloe, Barbadensis Leaft Powder, Carbomer, Glycerin, Propylene Glycol, Tocopheryl Acetate, Water

**Questions?** Call 1-800-777-4908

McKesson

16-3460

Instant Hand Sanitizing Wipe

5.5 in x 7.9 in

(14 cm x 20 cm)

160 wipes per canister





# **INSTANT HAND SANITIZING WIPE**

ethyl alcohol swab

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:68599-5808
Route of Administration	TOPICAL		

# Active Ingredient/Active Moiety Ingredient Name Basis of Strength ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M) ALCOHOL (55.9 mg

Inactive Ingredients			
Ingredient Name	Strength		
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)			
ALOE (UNII: V5VD430YW9)			
GLYCERIN (UNII: PDC6A3C0OX)			

.ALPHATOCOPHEROL ACETATE (UNII: 9E8X80D2L0)		
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)		
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)		
WATER (UNII: 059QF0KO0R)		

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:68599- 5808-6	160 in 1 BOX; Type 0: Not a Combination Product	02/24/2020	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	02/24/2020	
inidi			

# **Labeler -** McKesson (023904428)

Establishment			
Name	Address	ID/FEI	Business Operations
AHC LTD		413138557	manufacture(68599-5808)

Revised: 4/2023 McKesson