IQ CARE- alcohol gel D D Office Products

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.

ACTIVE INGREDIENT

ALCOHOL 70%

INACTIVE INGREDIENTS

Water, Glycerin, Butylene Glycol, Triethanolamine, Carbomer

PURPOSE

Antiseptic

WARNINGS

Flammable. Keep away from flame and fire. For external use only.

When using this product

Keep out of eyes. In case of contact, rinse eyes thoroughly with water.

Stop use and ask a doctor if irritation or redness develops and lasts.

KEEP OUT OF REACH OF CHILDREN

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

Uses

Hand sanitizer to help reduce bacteria on the skin. *Kills most common germs that may cause illness.

Directions

Put enough product in your palm and rub hands together until dry. For children under 6 years, use only under adult supervision.

Other information

Store below 110°F (43°C) May discolor some fabrics or surfaces

PACKAGE LABEL.PRINCIPAL DISPLAY PANEL





STERILIZING EFFECT 70% 16.9 fl. oz. (500ml)

Q Care GEL

DRUG FACTS

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IQ CARE

alcohol gel

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:77081-010

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name Basis of Strength Strength ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M) ALCOHOL 350 mL in 500 mL

Inactive Ingredients

Ingredient Name Strength Water (UNII: 059QF0KO0R) Glycerin (UNII: PDC6A3C0OX) Butylene Glycol (UNII: 3XUS85K0RA) TROLAMINE (UNII: 903K93S3TK) CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0 A5MM307FC)

Packaging

	# Item Code	Package Description	Marketing Start Date	Marketing End Date
l	1 NDC:77081-010- 01	500 mL in 1 BOTTLE, PUMP; 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	part333E	05/01/2020	

Labeler - D D Office Products (023854701)

Registrant - Handock Cosmetics Co., Ltd. (688037204)

Establishment								
Name	Address	ID/FEI	Business Operations					
Handock Cosmetics Co., Ltd.		688037204	manufacture(77081-010)					

Revised: 5/2020 D D Office Products