NEW HANSAN HAND SANITIZER- alcohol gel
Bb & Co., Ltd.

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.

Drug Facts

ACTIVE INGREDIENT

ACTIVE INGREDIENT: ALCOHOL 70.0%

INACTIVE INGREDIENT

INACTIVE INGREDIENTS:

Water, Glycerin, Butylene Glycol, Aloe Barbadensis Leaf Extract, Carbomer, Triethanolamine, Calendula Officinalis Extract, Flavor

PURPOSE

PURPOSE: SANITIZER

KEEP OUT OF REACH OF CHILDREN

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

Uses:

Uses : Hand sanitizer to help reduce bacteria on the skin.

WARNINGS

Warnings:

Flammable. Keep away from fire or flame.
For external use only.
When using this product do not use in or near the eyes. In case of contact, rinse eyes thoroughly with water.
Discontinue use if irritation or redness develops. If condition persists for more than 72 hours, consult a doctor.
Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

Directions

Directions:

• Put enough product in your palm to cover hands and rub hands together briskly until dry.
• Children under 6 years of age should be supervised when using this product.

Other Information

Other Information:
• Store below 90°F(32°C)
• May discolor certain fabrics or surfaces

PACKAGE LABEL.PRINCIPAL DISPLAY PANEL

NEW HANSAN HAND SANITIZER

alcohol gel

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:74080-010
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	42.0 g in 60 mL

Inactive Ingredients

Ingredient Name	Strength
Water (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
Butylene Glycol (UNII: 3XUS85K0RA)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)	
Trolamine (UNII: 9O3K93S3TK)	
CALENDULA OFFICINALIS FLOWER (UNII: P0M7O4Y7YD)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:74080-010-02	1 in 1 CARTON	03/01/2020	
1	NDC:74080-010-01	60 mL in 1 CONTAINER; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	03/01/2020	

Labeler - Bb & Co., Ltd. (695732817)

Registrant - Bb & Co., Ltd. (695732817)

Establishment

Name	Address	ID/FEI	Business Operations
KMPHARMACEUTICAL Co.,Ltd.		688679158	manufacture(74080-010)

Revised: 3/2020 Bb & Co., Ltd.