BLUE CEDAR HAND SANITIZER- hand sanitizer gel
Anhui Kiwi Biotech 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.

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

Flammable. Keep away from fire or flame.

Forexternal use only.

Do not use this product in or near the eyes. In case of accidental contact, ninse eyes thoroughly with water.

Stop use and ask doctor if iritation or rash appears andlasts.

Keep away from children. If sallowed, get medicalhelp right away,

Kills Harmful Bacteria or Germs

keep out of reach of children.

Place enough product (1.5~-2 mL, about one full pump) in your palm to thoroughly cover your hands.

Rub hands briskly until dry.

Children under 6 years old should be supervised when using this product.

Ethyl Alcohol 75 59% 6(VW) or 65 5%

Water, Carbomer, Glycereth :26, Triethanolamine, Fragrance (Parfum).

65*100mm

BLUE CEDAR HAND SANITIZER

hand sanitizer gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:55444-001
Route of Administration	EXTRACORPOREAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	375 mL in 500 mL

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
.ALPHA.-HEXYLCINNAMALDEHYDE (UNII: 7X6O37OK2I)	
.ALPHA.-TERPINEOL (UNII: 21334LVV8W)	
TERPINEOL (UNII: R53Q4ZWC99)	

.GAMMA.-NONALACTONE (UNII: I1XGH66S8P)	
LINALOOL, (+/-)- (UNII: D81QY6I88E)	
METHYL DIHYDROJASMONATE (SYNTHETIC) (UNII: 3GW44CIE3Y)	
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)	
TROLAMINE (UNII: 9O3K93S3TK)	
GLYCERETH-26 (UNII: NNE56F2N14)	
PHENYLETHYL ALCOHOL (UNII: ML9LGA7468)	

Product Characteristics

Color		**Score**	
Shape		**Size**	
Flavor	FRUIT	**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:55444-001-01	500 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/21/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	03/21/2020	

Labeler - Anhui Kiwi Biotech Co., Ltd. (554444636)

Establishment

Name	Address	ID/FEI	Business Operations
Anhui Kiwi Biotech Co., Ltd.		554444636	manufacture(55444-001)

Revised: 4/2020 Anhui Kiwi Biotech Co., Ltd.