

KIONI HAND SANITIZER- alcohol gel
YOON JI CORPORATION 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.

KIONI HAND SANITIZER 1.5 ml x 50 : 75024-700-50



KIONI HAND SANITIZER

alcohol gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:75024-700
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	1.05 mL in 1.5 mL

Inactive Ingredients

Ingredient Name	Strength
GREEN TEA LEAF (UNII: W2ZU1RY8B0)	

GLYCERIN (UNII: PDC6A3C0OX)	
CARBOMER HOMO POLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)	
WATER (UNII: 059QF0KO0R)	
PANTHENOL (UNII: WV9CM0O67Z)	
TROMETHAMINE (UNII: 023C2WHX2V)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:75024-700-50	50 in 1 BOX	09/22/2020	
1		1.5 mL in 1 POUCH; 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	part333A	09/22/2020	

Labeler - YOON JI CORPORATION CO.,LTD (688700744)

Establishment

Name	Address	ID/FEI	Business Operations
YOON JI CORPORATION CO.,LTD		688700744	manufacture(75024-700)

Revised: 9/2020

YOON JI CORPORATION CO.,LTD