FELIX HAND SANITIZER GEL- alcohol gel Ester 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.

[Ester Co., Ltd.-FELIX HAND SANITIZER GEL] 500 ml: BOTTLE+PUMP: 74004-702-01



FELIX HAND SANITIZER GEL

alcohol gel

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:74004-702	
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		
ALLANTO IN (UNII: 344S277G0Z)			
GREEN TEA LEAF (UNII: W2ZU1RY8B0)			
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)			
EDETOL (UNII: Q4R969U9FR)			
CARBOMER HO MO PO LYMER, UNSPECIFIED TYPE (UNII: 0 A5MM307FC)			
WATER (UNII: 059QF0KO0R)			

ALOE ARBORESCENS LEAF (UNII: 09TD8L5SQV)

GLYCERIN (UNII: PDC6A3C0OX)

ı	Packaging				
	# It	em Code	Package Description	Marketing Start Date	Marketing End Date
	1 NDC 01	:74004-702-	500 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	05/21/2020	

Marketing Information					
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date		
OTC monograph not final	part333A	05/21/2020			

Labeler - Ester Co., Ltd. (688425766)

Establishment				
Name	Address	ID/FEI	Business Operations	
Ester Co., Ltd.		688425766	manufacture(74004-702)	

Revised: 5/2020 Ester Co., Ltd.