ULINE HAND SANITIZER- alcohol gel
Uline

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.

Uline Hand Sanitizer

Active Ingredient

Active Ingredient

Ethyl Alcohol 70%

Uses

Uses

• Hand sanitizer to reduce microorganisms on the skin.

• Use this product when soap and water are not available.

Purpose

Purpose

Antiseptic

Warnings

Warnings

• For external use only.

• Avoid contact with eyes. If contact occurs, rinse thoroughly with water.

• FLAMMABLE. This product contains ethyl alcohol. Keep away from sources of ignition.

• Discontinue use if irritation or redness develops.

• If irritation persists for more than 72 hours, consult a physician.

• KEEP OUT OF REACH OF CHILDREN.

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

Directions

Directions

• Read the entire label before using this product.

• Place enough product on your palm to thoroughly cover your hands.

• Rub hands together briskly until dry.

Inactive Ingredients

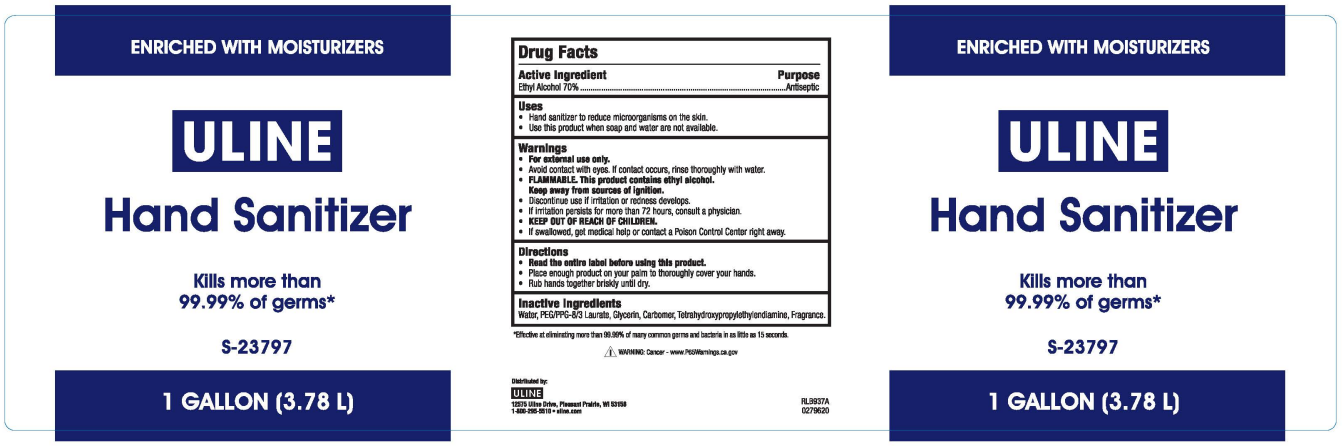
Inactive Ingredients

Water, PEG/PPG-8/3 Laurate, Gycerin, Carbomer, Tetrahydroxypropylethylendiamine, Fragrance.

Warnings

KEEP OUT OF REACH OF CHILDREN.

Principal Display Panel



3780 ml 69790-801-04

ULINE HAND SANITIZER

alcohol gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:69790-801
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	0.7 mL in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
CARBOMER INTERPOLYMER TYPE A (ALLYL SUCROSE CROSSLINKED) (UNII: 59TL3WG5CO)	
DIMETHICONE PEG-8 LAURATE (UNII: 72MF9C2A18)	
GLYCERIN (UNII: PDC6A3C0OX)	
EDETOL (UNII: Q4R969U9FR)	

Packaging				
			Marketing Start	Marketing End

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69790-801-04	3780 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	06/08/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	06/08/2020	

Labeler - Uline (039612668)

Registrant - Betco Corporation, Ltd (024492831)

Establishment

Name	Address	ID/FEI	Business Operations
Betco Corporation, Ltd		024492831	manufacture(69790-801) , label(69790-801)

Revised: 6/2020 Uline