CLARIO INSTANT HAND SANITIZER- alcohol soap
Betco Corporation, 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.

Clario Instant Hand Sanitizer

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Active Ingredient

Ethyl Alcohol 62%

Clario Instant Hand Sanitizer

Uses

- Hand sanitizer to reduce microorganisms on the skin.
- Use this product when soap and water are not available.

Clario Instant Hand Sanitizer

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 is 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 the poison control center right away.

Clario Instant Hand Sanitizer

Directions

- **Read the entire label before using this product.**
- Dispense two pumps of product onto palm of hand and rub thoroughly over all surfaces for both hands until dry.

Clario Instant Hand Sanitizer

Inactive Ingredients

Water, PEG/PPG-8/3 Laurate, Tetrahydroxypropyl Ethelendiamine, Carbomer, Fragrance.

Clario Instant Hand Sanitizer

Purpose

Antiseptic

Clario Instant Hand Sanitizer

KEEP OUT OF REACH OF CHILDREN

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Ethanol Based Gel Hand Sanitizer

KILLS 99.9% of Germs

Net Contents: 8 fl.oz. (236 mL)

Hand Sanitizer 776

RLB0334 0277614

CLARIO INSTANT HAND SANITIZER

alcohol soap

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:65601-776
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	0.62 mL in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
EDETOL (UNII: Q4R969U9FR)	
CARBOMER HOMOPOLYMER TYPE C (UNII: 4Q93RCW27E)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:65601-776-29	1000 mL in 1 BAG; Type 0: Not a Combination Product	05/01/2013	
2	NDC:65601-776-24	118 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	05/01/2013	03/26/2019

3	NDC:65601-776-68	236 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	05/01/2013	03/26/2019
4	NDC:65601-776-69	473 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	05/01/2013	03/26/2019
5	NDC:65601-776-57	550 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	05/01/2013	
6	NDC:65601-776-19	900 mL in 1 BOX; Type 0: Not a Combination Product	05/01/2013	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	05/01/2013	

Labeler - Betco Corporation, Ltd. (024492831)

Registrant - Betco Corporation, Ltd. (024492831)

Establishment

Name	Address	ID/FEI	Business Operations
Betco Corporation, Ltd.		024492831	manufacture(65601-776) , pack(65601-776) , label(65601-776)

Revised: 3/2019 Betco Corporation, Ltd.