ECO SANITIZER GEL- ethyl alcohol gel
0966452 B.C. 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.

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eco SANITIZER HAND SANITIZER GEL

Purpose: Antiseptic

Use: Hand sanitizer to help reduce bacteria on skin

Directions for Use: Supervise children when they use this product. For occasional and personal domestic use. Rub thoroughly into hands for at least 30 seconds. Allow to dry.

Medicinal Ingredients: Ethyl Alcohol* (70% v/v).

Ingredients: Aqua, Glycerin,* Hydroxypropylcellulose*. *Plant based ingredient.

Warning: For external use only. Keep out of reach of children. Flammable. Keep away from open flame and sources of heat. Avoid contact with eyes. Should this occur, flush eyes with water. If irritation develops, discontinue use and contact a health care practitioner. Store between 15°C/59°F and 30°C/86°F.

Contraindications: Do not use on children/infants less than 2 years of age (unless directed by a health care practitioner).

Keep out of reach of children.

kills 99% of germs

Get your hands on it!

Fragrance free | Hypoallergenic

Enriched Gel with Glycerin | Can be used as is or as a refill

Made in Canada from Domestic and Imported products.

Eco Sanitizer | ecosanitizer.ca

Packaging

ECO SANITIZER GEL

ethyl alcohol gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:80731-002
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	70 mL in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
HYDROXYPROPYL CELLULOSE, UNSPECIFIED (UNII: 9XZ8H6N6OH)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:80731-002-01	50 mL in 1 BOTTLE; Type 0: Not a Combination Product	10/08/2020	
2	NDC:80731-002-05	320 mL in 1 BOTTLE; Type 0: Not a Combination Product	10/27/2020	
3	NDC:80731-002-03	1000 mL in 1 BOTTLE; Type 0: Not a Combination Product	10/08/2020	
4	NDC:80731-002-04	4000 mL in 1 BOTTLE; Type 0: Not a Combination Product	10/08/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date

OTC monograph not final	part333A	10/08/2020	

Labeler - 0966452 B.C. Ltd. (204193445)

Establishment

Name	Address	ID/FEI	Business Operations
0966452 B.C. Ltd.		204193445	manufacture(80731-002)

Revised: 10/2020 0966452 B.C. Ltd.