

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.

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

kills 99% of germs

eco
SANITIZER
Get your hands on it!

HAND SANITIZER GEL
désinfectant en gel pour les mains

Contains 70% Ethyl Alcohol Contient 70% D'Alcool Ethylique

Fragrance free | Hypoallergenic
San parfum | Hypoallergénique

npn #80100135

1L/33.8 fl oz

Enriched Gel with Glycerin | Can be used as is or as a refill
Gel enrichi avec glycérine | Utiliser tel quel ou comme recharge

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. **Mode d'emploi:** Surveillez les enfants lorsqu'ils utilisent ce produit. Pour usage domestique occasionnel et personnel. Frottez soigneusement dans vos mains pendant au moins 30 secondes. Laisser sécher.

Medicinal Ingredients: Ethyl Alcohol (70% v/v). **Ingrédients médicinaux:** Alcool Ethylique* (70% v/v). **Ingredients:** Aqua, Glycerin, Hydroxypropylcellulose. **Ingrédients:** Aqua, Glycérine*, Hydroxypropylcellulose*. **Ingrédient à base de plantes.**

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. **Avertissement:** Pour usage externe seulement. Tenir hors de la portée des enfants. Inflammable. Tenir loin des flammes nues et des sources de chaleur. Eviter tout contact avec les yeux. En cas de contact avec les yeux, rincer avec de l'eau. En cas d'irritation, cesser l'emploi et consulter un médecin. Conserver entre 15°C/59°F et 30°C/86°F.

Contraindications: Do not use on children/infants less than 2 years of age (unless directed by a health care practitioner). **Contre-indications:** Ne pas utiliser sur les enfants/nourrissons de moins de 2 ans (sauf indication contraire d'un professionnel de la santé).

caution
flammable/
inflammable
HDPE

Made in Canada
from Domestic and Imported products.
Fabriqué au Canada
à partir de produits locaux et importés.

Eco Sanitizer | ecosanitizer.ca

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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: 059QF0K00R)	
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
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OTC monograph not final	part333A	10/08/2020	
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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.