

G AND H PROTECT HAND SANITIZER- alcohol gel
Access Business Group LLC

G&H Protect Hand Sanitizer

Drug Facts

Active Ingredient

Ethyl alcohol 65.6%

Purpose

Antiseptic

Uses

Hand sanitizer to help reduce bacteria on the skin that potentially can cause disease. For use when soap and water are not available.

Warnings

For external use only. Flammable. Keep away from heat or flame.

Do not use

- In children less than 2 years of age.
- On open skin wounds.

When using this product

keep out of eyes, ears, and mouth. In case of contact with eyes, rinse eyes thoroughly with water.

Stop use and ask a doctor

if irritation or rash occurs. These may be signs of a serious condition.

Keep out of reach of children.

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

Directions

- Place enough product on hands to cover all surfaces. Rub thoroughly into hands for at least 30 seconds. Allow to dry.
- Supervise children under 6 years of age when using this product to avoid swallowing.

Other information

- Store between 15-30°C (59-86°F)
- Avoid freezing and excessive heat above 40°C (104°F).

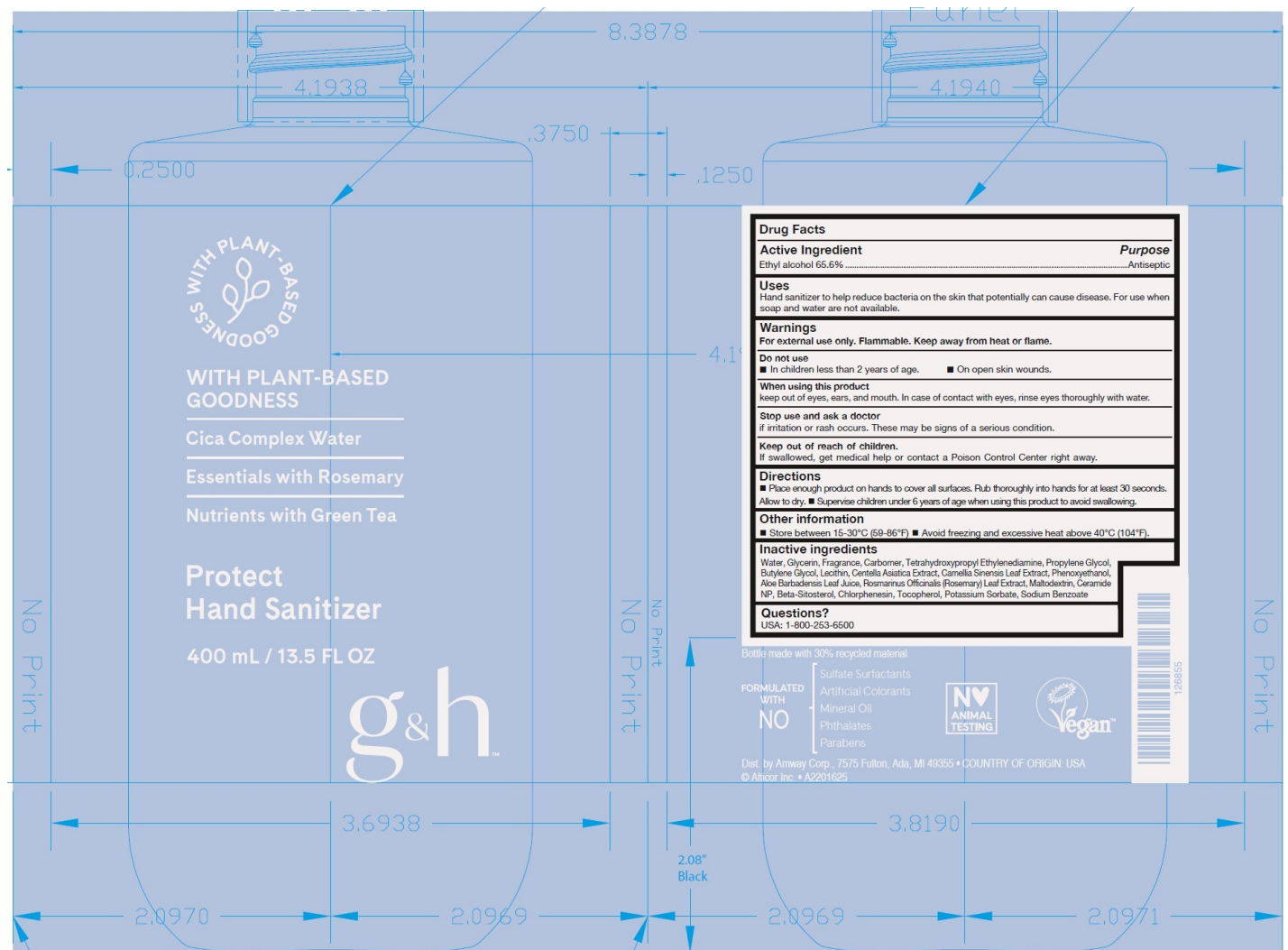
Inactive ingredients

Water, Glycerin, Fragrance, Carbomer, Tetrahydroxypropyl Ethylenediamine, Propylene Glycol, Butylene Glycol, Lecithin, Centella Asiatica Extract, Camellia Sinensis Leaf Extract, Phenoxyethanol, Aloe Barbadensis Leaf Juice, Rosmarinus Officinalis (Rosemary) Leaf Extract, Maltodextrin, Ceramide NP, Beta-Sitosterol, Chlorphenesin, Tocopherol, Potassium Sorbate, Sodium Benzoate

Questions?

USA: 1-800-253-6500

Package Labeling:



G AND H PROTECT HAND SANITIZER

alcohol gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:10056-532
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	65.6 mL in 100 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0K00R)	
GLYCERIN (UNII: PDC6A3C00X)	
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)	
EDETOL (UNII: Q4R969U9FR)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	
CENTELLA ASIATICA TRITERPENOIDS (UNII: 4YS74Q4G4J)	
GREEN TEA LEAF (UNII: W2ZU1RY8B0)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
ROSEMARY (UNII: IJ67X351P9)	
MALTODEXTRIN (UNII: 7CVR7L4A2D)	
CERAMIDE NP (UNII: 4370DF050B)	
.BETA.-SITOSTEROL (UNII: S347WMO6M4)	
CHLORPHENESIN (UNII: I670DAL4SZ)	
TOCOPHEROL (UNII: R0ZB2556P8)	
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:10056-532-00	400 mL in 1 BOTTLE; Type 0: Not a Combination Product	11/11/2023	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	11/11/2023	

Labeler - Access Business Group LLC (839830713)

