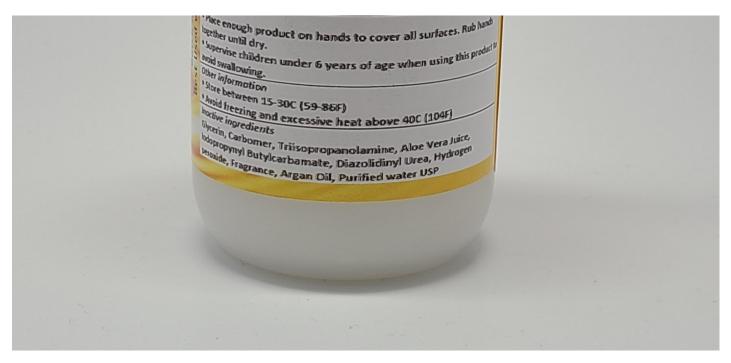
THE ESSENCE OF EVA- hand sanitizing lotion lotion F & E Group LLP

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.

Back Label NDC 79206-003-06





Active ingredient[s] Purpose

Isopropyl alcohol 70%v/v.....Antiseptic

Use[s]

Hand sanitizer to help reduce bacteria 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 months 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 hands together until dry.
- Supervise children under 6 years of age when using this product to avoid swallowing.

Other information

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

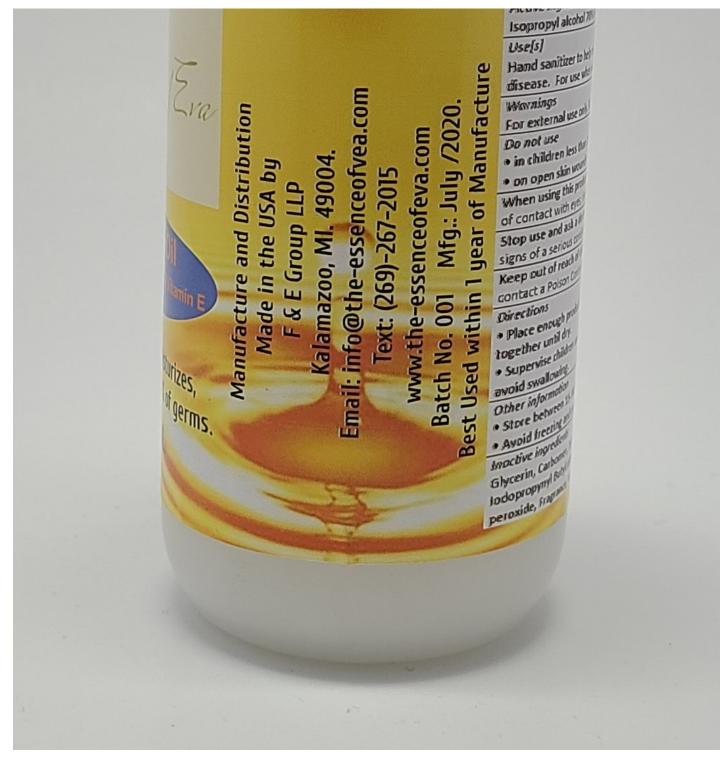
NDC 79206-003

Inactive ingredients

Glycerin, Carbomer, Triisopropanolamine, Aloe Vera Juice, Iodopropynyl Butylcarbamate, Diazolidinyl Urea, hydrogen peroxide, fragrance,Argan Oil, purified water USP

Manufacture and Distribution Made in the USA by F & E Group LLP Kalamazoo, MI. 49004. Email: info@the-essenceofvea.com Text: (269)-267-2015 www.the-essenceofeva.com Batch No. 001 Mfg.: June /2020. Best Used within 1 year of Manufacture





Directions

- Place enough product on hands to cover all surfaces. Rub hands together until dry.
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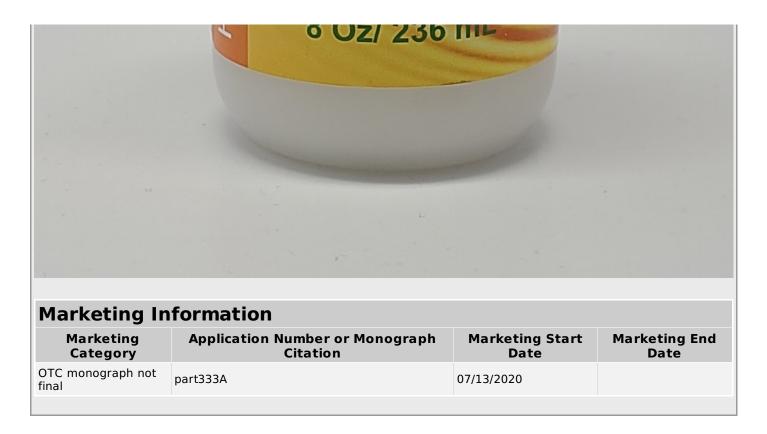
Primary label front NDC 79206-003-06 236 mL





THE ESSENCE OF EVA hand sanitizing lotion lotion **Product Information Product Type** HUMAN OTC DRUG Item Code (Source) NDC:79206-003 **Route of Administration** TOPICAL **Active Ingredient/Active Moiety Basis of Ingredient Name** Strength Strength ISOPROPYL ALCOHOL (UNII: ND2M416302) (ISOPROPYL ALCOHOL -ISOPROPYL 70 mL UNII:ND2M416302) ALCOHOL in 100 mL **Inactive Ingredients** Strength **Ingredient Name** 35 mL in 100 mL ALOE (UNII: V5VD430YW9) HYDROGEN PEROXIDE (UNII: BBX060AN9V) 0.125 mL in 100 mL 2 mL in 100 mL GLYCERIN (UNII: PDC6A3C0OX) 15 mL in 100 mL WATER (UNII: 059QF0KO0R) 5 mL in 100 mL ARGAN OIL (UNII: 4V59G5UW9X) **Product Characteristics** Color Score Shape Size Flavor **Imprint Code** Contains Packaging **Marketing Start Marketing End** # Item Code **Package Description** Date Date 1 NDC:79206-236 mL in 1 BOTTLE, PUMP; Type 0: Not a 07/13/2020 003-06 Combination Product





Labeler - F & E Group LLP (117556801)

Revised: 12/2021

F & E Group LLP