

HAND SANITIZER- alcohol liquid
K7 Design Group Inc.

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.

K7 BBC Aloe Hand Sanitizer 62 Blstr

Active Ingredient

Alcohol Denat. 62%

Purpose

Antiseptic

Use

for hand-washing to help decrease bacteria on the skin, only when water is not available

Warnings

For external use only. Flammable, keep away from fire and flames.

When using this product

- do not get into eyes.
- if contact occurs, rinse eyes thoroughly with water.

Stop use and ask a doctor if

irritation and redness develops and persists.

Keep out of reach of children

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

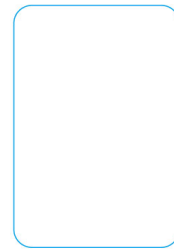
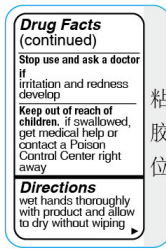
Directions

wet hands thoroughly with product and allow to dry without wiping

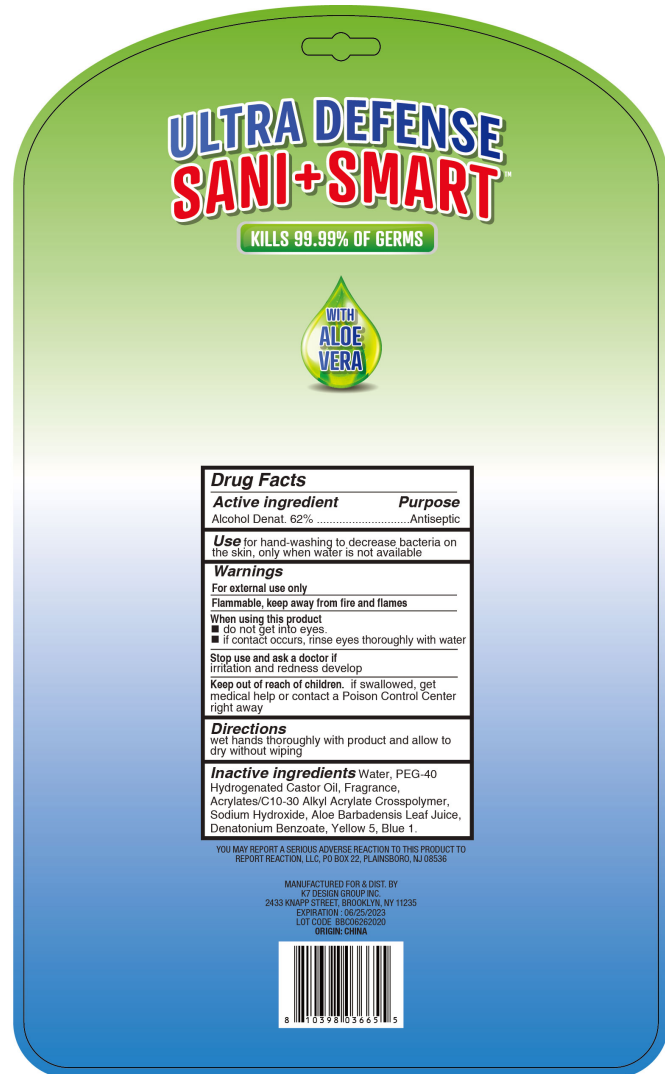
Inactive ingredients

Water, PEG-40 Hydrogenated Castor Oil, Fragrance, Acrylates/C10-30 Alkyl Acrylates Crosspolymer, Sodium Hydroxide, Aloe Barbadensis Leaf Juice, Denatonium Benzoate, Yellow 5, Blue 1

Inner Package Label



Blister Package Label



HAND SANITIZER

alcohol liquid

Product Information

| | | | |
|-------------------------|----------------|--------------------|---------------|
| Product Type | HUMAN OTC DRUG | Item Code (Source) | NDC:74177-028 |
| Route of Administration | TOPICAL | | |

Active Ingredient/Active Moiety

| Ingredient Name | Basis of Strength | Strength |
|--|-------------------|-----------------|
| ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M) | ALCOHOL | 62 mL in 100 mL |

Inactive Ingredients

| Ingredient Name | Strength |
|--|----------|
| ALOE VERA LEAF (UNII: ZY81Z83H0X) | |
| GLYCERIN (UNII: PDC6A3C0OX) | |
| WATER (UNII: 059QF0KO0R) | |
| DENATONIUM BENZOATE (UNII: 4YK5Z54AT2) | |
| FD&C BLUE NO. 1 (UNII: H3R47K3TBD) | |
| POLYOXYL 40 HYDROGENATED CASTOR OIL (UNII: 7YC686GQ8F) | |
| SODIUM HYDROXIDE (UNII: 55X04QC32I) | |
| FD&C YELLOW NO. 5 (UNII: I753WB2F1M) | |

Packaging

| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
|---|------------------|---|----------------------|--------------------|
| 1 | NDC:74177-028-50 | 6 in 1 BLISTER PACK | 06/22/2020 | |
| 1 | NDC:74177-028-02 | 50 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product | | |

Marketing Information

| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|-------------------------|--|----------------------|--------------------|
| OTC monograph not final | part333A | 06/22/2020 | |

Labeler - K7 Design Group Inc. (080357784)

Revised: 7/2020

K7 Design Group Inc.