CHEMPACE SANI-GEL INSTANT SKIN SANITIZER- ethanol gel Chempace Corporation

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.

chempace SANI-GEL INSTANT SKIN SANITIZER

Drug Facts

Active Ingredient:

Ethanol (60% v/v)

Purpose:

Anti-Microbial Hand Sanitizer

Uses

- Helps reduce bacteria that potentially can cause disease
- Helps prevent cross contamination by hand contact
- Recommended for repeated use

Warnings

- For external use only
- Flammable, keep away from fire, heat, or flame
- Keep out of reach of children.
- Do not use near eyes
- In case of eye contact flush with water for 15 minutes
- If irritation persists stop use of product and get medical attention
- In case of accidental ingestion seek medical attention or contact a poison control center immediately.

Directions

- Use no water or towels
- Apply appropriate amount of product to palm of hand
- Rub until hands are completely covered
- Agitate lightly until dry
- Let air dry for 15 seconds
- Do not rinse or wipe with towel.

Other Information

• Store in a cool dry place below 104° F.

Inactive Ingredients

Water, Carbomer, Triethanolamine, PEG-75 Lanolin, Aloe Vera Gel, Fragrance.

Principal Display Panel

Principal Display Panel – Bottle Label

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SANI-GEL INSTANT SKIN SANITIZER

- Enhanced with Moisturizers
- Kills disease causing germs within seconds

- Effective against MRSA, VRE, E. coli (0157:H7) Staphylococcus, Streptococcus and other organisms
- Assists with OSHA Bloodborne Pathogen Standard Compliance

For Hospital and Professional Use Only See Drug Facts panel for additional information

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Net Contents: 8 Fl. Oz.

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Manufactured for: CHEMPACE CORPORATION 339 arco drive toledo, ohio 43607 1-800-423-5350 515-001-014-305

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ethanol gel

| Product Information | | | | |
|-------------------------------------|--------------------|---------------|-------------|-------------------|
| Product T ype | HUMAN OTC DRUG | Item Code (So | ource) | NDC:64191-515 |
| Route of Administration | TOPICAL | | | |
| | | | | |
| Active Ingredient/Active Moi | a + 1 | | | |
| Active Ingredient/Active Mon | ety | | | |
| Ingredie | nt Name | Basis | of Strength | Strength |
| ethanol (UNII: 3K9958V90M) (ethanol | - UNII:3K9958V90M) | ethanol | e | 500 mL in 1000 mL |
| | | | | |
| Inactive Ingredients | | | | |
| | Strength | | | |
| water (UNII: 059QF0KO0R) | | | | |
| carbomer homopolymer type C (UNII | : 4Q93RCW27E) | | | |
| aloe (UNII: V5VD430YW9) | | | | |

| trolamine (UNII: 9O3K93S3TK) | | | | | | | |
|------------------------------|--|-----------|----------------------|----|--------------------|--|--|
| | | | | | | | |
| Packaging | | | | | | | |
| # Item Code | Package Description | Marketing | Start Date | Ma | rketing End Date | | |
| 1 NDC:64191-515-62 | 237 mL in 1 BOTTLE | | | | | | |
| | | | | | | | |
| | | | | | | | |
| Marketing Information | | | | | | | |
| Marketing Category | Application Number or Monograph Citation | | Marketing Start Date | | Marketing End Date | | |
| OTC monograph not final | part333 | | 05/21/2010 | | | | |
| | | | | | | | |

Labeler - Chempace Corporation (043636471)

| Establishment | | | | | | | |
|----------------------|---------|-----------|----------------------------|--|--|--|--|
| Name | Address | ID/FEI | Business Operations | | | | |
| Canberra Corporation | | 068080621 | MANUFACTURE | | | | |

Revised: 5/2010

Chempace Corporation