GERM WAR HAND SANITIZER GEL- alcohol gel SKINFARM

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.

Drug Facts

Alcohol

Water, Carbomer, Trometharmine, Glycerin, Aloe Barbadensis Leaf Juice, Fragrance

Antiseptic

KEEP OUT OF REACH OF THE CHILDREN

For the external use only

For external use only.

Avoid contact with eyes. If introduced to eyes, flush with water.

Do not ingest.

Discontinue use and contact a doctor if irritation and redness develops and condition persist for more than 72 hours.

Keep out of reach of children. if swallowed, get medial help or contact a Poison Control Center right away

Place a small amount in your palm and rub hands together until dry. Children under 6 years of age should be supervised when using this product.

GALLON REFILL



GW203AG11-ART REV A 190446699118



| GERM WAR HAND SANITIZER GEL | | | | | | | |
|---------------------------------|----------------------------|-----------------------------------|------------------|----------------|----------|--|--|
| lcohol gel | | | | | | | |
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| Product Information | | | | | | | |
| Product T ype | HUMAN OTC DRUG | Item Code (Source) NDC:73793-0010 | | 793-0010 | | | |
| Route of Administration | TOPICAL | | | | | | |
| | | | | | | | |
| | | | | | | | |
| Active Ingredient/Active Moi | ety | | | | | | |
| Ingredient Name | | | Basis of Strengt | h | Strength | | |
| ALCOHOL (UNII: 3K9958V90M) (ALC | ALCOHOL 6 | | 66 | .5 g in 100 mL | | | |
| | | | | | | | |
| 11 . | | | | | | | |
| Inactive Ingredients | | | | | | | |
| Ingredient Name | | | | | Strength | | |
| TROLAMINE (UNII: 903K93S3TK) | | | | | | | |
| CARBOMER HOMOPOLYMER, UNSI | PECIFIED TYPE (UNII: 0A5MM | 307FC) | | | | | |
| GLYCERIN (UNII: PDC6A3C0OX) | | | | | | | |
| WATER (UNII: 059QF0KO0R) | | | | | | | |

| ALOE VERA LEAF (UNII: ZY8 1Z8 3H0 X) | | | | | | | | |
|--------------------------------------|--------------|---|----------------------|--------------------|--|--|--|--|
| | | | | | | | | |
| Packaging | | | | | | | | |
| # Item Co | ode | Package Description | Marketing Start Date | Marketing End Date | | | | |
| 1 NDC:73793 | -0010-1 38 | 800 mL in 1 BOTTLE; Type 0: Not a Combination Product | 09/01/2020 | | | | | |
| | | | | | | | | |
| | | | | | | | | |
| Marketing Information | | | | | | | | |
| Marketing | Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date | | | | |
| OTC monogra | ph not final | l part333A | 09/01/2020 | | | | | |
| | | | | | | | | |

Labeler - SKINFARM (688594873)

Registrant - SKINFARM (688594873)

| Establishment | | | | | | | |
|---------------|---------|-----------|-------------------------|--|--|--|--|
| Name | Address | ID/FEI | Business Operations | | | | |
| SKINFARM | | 688594873 | manufacture(73793-0010) | | | | |

Revised: 9/2020

SKINFARM