HAND SANITIZER GEL- hand sanitizer gel gel LS Promotions 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.

Hand Sanitizer Gel - Single Use Packet

Drug Facts

Drug Facts

Active Ingredient

Ethanol 62%

PURPOSE

ANTISEPTIC

USES

TO HELP DECREASE BACTERIA ON HANDS

WARNINGS

FOR EXTERNAL USE ONLY. FLAMMABLE, KEEP AWAY FROM FIRE OR FLAME

WHEN USING THIS PRODUCT

KEEP OUT OF EYES. IN CASE OF CONTACT WITH EYES, FLUSH THOROUGHLY WITH WATER AND THEN SEEK MEDICAL ADVICE.

STOP USE AND ASK A DOCTOR

IF IRRITATION OR REDNESS DEVELOPS

KEEP OUT OF REACH OF CHILDREN

IF SWALLOWED, GET MEDICAL HELP OR CONTACT A POISON CONTROL CENTER IMMEDIATELY.

DIRECTIONS

PLACE ENOUGH PRODUCT IN YOUR PALM TO THOROUGHLY COVER HAND. RUB HANDS TOGETHER BRISKLY UNTIL DRY. FOR CHILDREN UNDER 6 YEARS OLD, USE ADULT SUPERVISION.

OTHER INFORMATION

DO NOT STORE ABOVE 100° F.

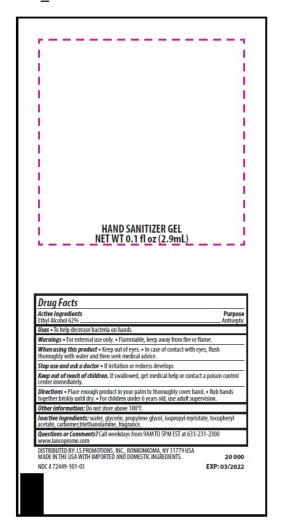
INACTIVE INGREDIENTS

Inactive Ingredients: water, glycerin, propylene glycol, isopropyl myristate, tocopheryl acetate, carbomer, triethanolamine, fragrance

QUESTIONS OR COMMENTS?

CALL WEEKDAYS FROM 9AM TO 5PM EST AT 631-231-2300 WWW.LANCOPROMO.COM

PDP_Packet



HAND SANITIZER GEL hand sanitizer gel gel Product Information Product Type HUMAN OTC DRUG Item Code (Source) NDC:72449-101 Route of Administration TOPICAL Active Ingredient/Active Moiety Ingredient Name Basis of Strength Strength

| Inactive Ingredients | | | | |
|--|----------|--|--|--|
| Ingredient Name | Strength | | | |
| CARBOMER 1342 (UNII: 809 Y72KV36) | | | | |
| WATER (UNII: 059QF0KO0R) | | | | |
| GLYCERIN (UNII: PDC6A3C0OX) | | | | |
| TROLAMINE (UNII: 9O3K93S3TK) | | | | |
| ISOPROPYL MYRISTATE (UNII: 0 RE8 K4LNJS) | | | | |
| .ALPHATO COPHEROL ACETATE, D- (UNII: A7E6112E4N) | | | | |
| PROPYLENE GLYCOL (UNII: 6 DC9 Q16 7 V3) | | | | |

| ı | Packaging | | | | | | | |
|---|--------------------|---|-----------------------------|--------------------|--|--|--|--|
| ı | # Item Code | Package Description | Marketing Start Date | Marketing End Date | | | | |
| ı | 1 NDC:72449-101-01 | 2.9 mL in 1 PACKET; Type 0: Not a Combination Product | 03/16/2020 | | | | | |
| ı | 2 NDC:72449-101-02 | 5.9 mL in 1 PACKET; Type 0: Not a Combination Product | 03/16/2020 | | | | | |

| Marketing Information | | | | | | | |
|-------------------------|--|----------------------|--------------------|--|--|--|--|
| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date | | | | |
| OTC monograph not final | part333E | 03/16/2020 | | | | | |
| | | | | | | | |

Labeler - LS Promotions Inc. (024818391)

Registrant - LS Promotions Inc. (024818391)

| Establishment | | | | | | |
|--------------------|---------|-----------|------------------------|--|--|--|
| Name | Address | ID/FEI | Business Operations | | | |
| LS Promotions Inc. | | 024818391 | manufacture(72449-101) | | | |

Revised: 3/2020 LS Promotions Inc.