DRS CLEAN HAND- alcohol gel THE NADREE Co., Ltd.

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, Glycerin, Aloe Extract, etc.

Antiseptic

KEEP OUT OF REACH OF THE CHILDREN

For the external use only

For external use only.

Flammable, keep away from fire or flame.

When using this product keep out of eyes. If contact with eyes occurs, rinse promptly and thoroughly with water.

Stop use and ask a doctor if significant irritation or sensitization develops.

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

Put suitable amount in your palm to cover hands and rub hands together briskly until dry.





DRS CLEAN HAND

alcohol gel

| Product Information | | | |
|-------------------------|----------------|--------------------|----------------|
| Product Type | HUMAN OTC DRUG | Item Code (Source) | NDC:74442-0005 |
| Route of Administration | TOPICAL | | |

| Active Ingredient/Active Moiety | | | |
|--|-------------------|----------------|--|
| Ingredient Name | Basis of Strength | Strength | |
| ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M) | ALCOHOL | 70 g in 100 mL | |

| Inactive Ingredients | | | |
|------------------------------------|----------|--|--|
| Ingredient Name | Strength | | |
| ALOE (UNII: V5VD430 YW9) | | | |
| WATER (UNII: 059QF0KO0R) | | | |
| BUTYLENE GLYCOL (UNII: 3XUS85K0RA) | | | |

| Packaging | | | | | |
|-----------|------------|-------------------|---|----------------------|--------------------|
| | # Item (| Code | Package Description | Marketing Start Date | Marketing End Date |
| | 1 NDC:7444 | 2-0005-1 60 mL in | 1 BOTTLE; Type 0: Not a Combination Produ | ct 04/01/2020 | |

| Marketing Information | | | | |
|-------------------------|--|----------------------|--------------------|--|
| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date | |
| OTC monograph not final | part333A | 04/01/2020 | | |
| | | | | |

Labeler - THE NADREE Co., Ltd. (690420363)

Registrant - THE NADREE Co., Ltd. (690420363)

| Establishment | | | | |
|---------------|---------|-----------|-------------------------|--|
| Name | Address | ID/FEI | Business Operations | |
| EQMAXON Corp. | | 557821534 | manufacture(74442-0005) | |

| Establishment | | | | |
|----------------------|---------|-----------|---------------------|--|
| Name | Address | ID/FEI | Business Operations | |
| THE NADREE Co., Ltd. | | 690420363 | label(74442-0005) | |

Revised: 4/2020 THE NADREE Co., Ltd.