**PSR HAND SANITIZER- ethyl alcohol spray**
**D-Time Limited Liability Company**

*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.*

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## Active Ingredient

Ethyl alcohol 80%

## Purpose

Antiseptic skin cleanser

## Uses

For personal hand hygiene to help prevent the spread of bacteria.

## Warnings

**For external use only. Flammable. Keep away from heat and flame**

**When using this product** avoid contact with eyes. If contact occurs, rinse thoroughly with water.

Stop use and consult a healthcare professional if irritation develops.

## Keep out of reach of children.

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

## Directions

- Adults and children over 2 years:
- For occasional and personal domestic use
- Supervise children when they use this product •
- Spray onto hands and rub thoroughly for at least 30 seconds. Allow to dry

## Inactive ingredients

Water (Aqua), Glycerin, Apricot kernel oil, Bergamot leaf oil, Hydrogen peroxide, Sodium pyruvate

## Other information

Store at 68° to 70° F (20° to 25° C).
May discolor certain fabrics or surfaces.

**Questions?** 1-844-800-6858

## Package


# PSR HAND SANITIZER

ethyl alcohol spray

| Product Information | | | |
|---|---|---|---|
| **Product Type** | HUMAN OTC DRUG | **Item Code (Source)** | NDC:75306-009 |
| **Route of Administration** | TOPICAL | | |

| Active Ingredient/Active Moiety | | |
|---|---|---|
| **Ingredient Name** | **Basis of Strength** | **Strength** |
| **ALCOHOL** (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M) | ALCOHOL | 80 ng in 100 mL |

| Inactive Ingredients | |
|---|---|
| **Ingredient Name** | **Strength** |
| **WATER** (UNII: 059QF0KO0R) | |
| **GLYCERIN** (UNII: PDC6A3C0OX) | |
| **APRICOT KERNEL OIL** (UNII: 54JB35T06A) | |
| **BERGAMOT OIL** (UNII: 39W1PKE3JI) | |
| **HYDROGEN PEROXIDE** (UNII: BBX060AN9V) | |
| **SODIUM PYRUVATE** (UNII: POD38AIF08) | |

## Packaging

| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
|---|---|---|---|---|
| 1 | NDC:75306-009-01 | 30 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product | 06/01/2020 | |
| 2 | NDC:75306-009-02 | 50 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product | 06/01/2020 | |
| 3 | NDC:75306-009-03 | 60 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product | 06/01/2020 | |
| 4 | NDC:75306-009-04 | 100 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product | 06/01/2020 | |
| 5 | NDC:75306-009-05 | 120 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product | 06/01/2020 | |
| 6 | NDC:75306-009-06 | 150 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product | 06/01/2020 | |
| 7 | NDC:75306-009-07 | 160 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product | 06/01/2020 | |
| 8 | NDC:75306-009-08 | 200 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product | 06/01/2020 | |
| 9 | NDC:75306-009-09 | 250 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product | 06/01/2020 | |
| 10 | NDC:75306-009-10 | 500 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product | 06/01/2020 | |
| 11 | NDC:75306-009-11 | 1000 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product | 06/01/2020 | |
| 12 | NDC:75306-009-12 | 3785 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product | 06/01/2020 | |
| 13 | NDC:75306-009-13 | 18927 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product | 06/01/2020 | |
| 14 | NDC:75306-009-14 | 5000 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product | 06/01/2020 | |
| 15 | NDC:75306-009-15 | 10000 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product | 06/01/2020 | |
| 16 | NDC:75306-009-16 | 15000 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product | 06/01/2020 | |
| 17 | NDC:75306-009-17 | 20000 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product | 06/01/2020 | |

## Marketing Information

| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|---|---|---|---|
| OTC monograph not final | part333A | 06/01/2020 | |

## Labeler - D-Time Limited Liability Company (081728006)

## Establishment

| Name | Address | ID/FEI | Business Operations |
|---|---|---|---|
| D-Time Limited Liability Company | | 081728006 | manufacture(75306-009) |

Revised: 11/2021 D-Time Limited Liability Company