

**PERAFLU D- peracetic acid liquid  
Valley Inc.**

*Disclaimer: This drug has not been found by FDA to be safe and effective, and this labeling has not been approved by FDA. For further information about unapproved drugs, click here.*

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**ACTIVE INGREDIENT**

PERACETIC ACID 0.2%

**INACTIVE INGREDIENTS**

Water, Acetic acid, Hydrogen peroxide

**PURPOSE**

Antifungal

**WARNINGS**

For external use only

Do not use on the human body

Avoid contact with eyes.

Discontinue use if signs of irritation or rashes appear.

Keep out of reach of children

**KEEP OUT OF REACH OF CHILDREN**

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**Uses**

Steriliant

**Directions**

Directions:

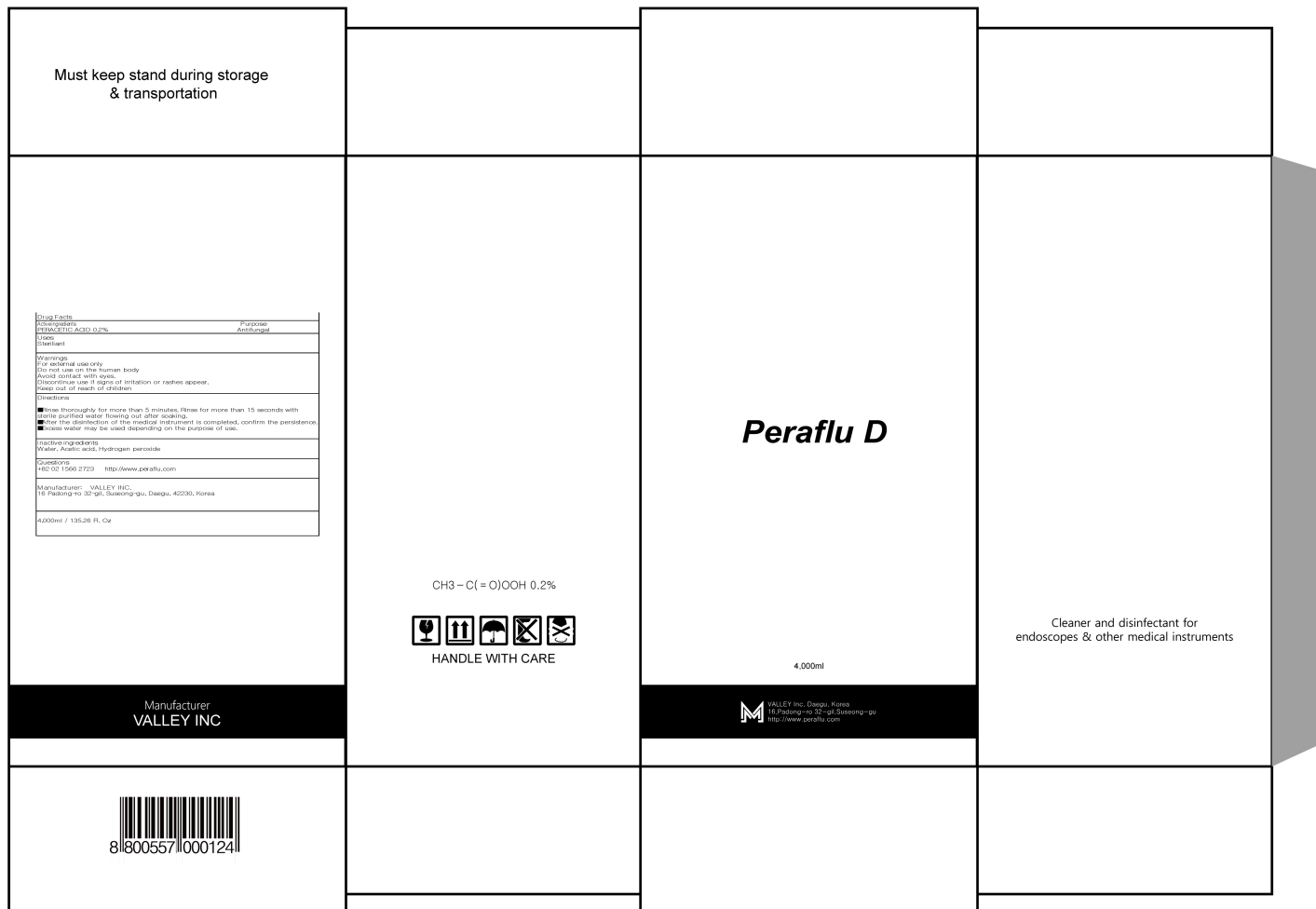
- Rinse thoroughly for more than 5 minutes. Rinse for more than 15 seconds with sterile purified water flowing out after soaking.
- After the disinfection of the medical instrument is completed, confirm the persistence.
- Excess water may be used depending on the purpose of use.

# QUESTIONS

+82 02 1566 2723

<http://www.peraflu.com>

## PACKAGE LABEL.PRINCIPAL DISPLAY PANEL



### PERAFLU D

peracetic acid liquid

#### Product Information

|                                |                |                           |               |
|--------------------------------|----------------|---------------------------|---------------|
| <b>Product Type</b>            | HUMAN OTC DRUG | <b>Item Code (Source)</b> | NDC:81803-020 |
| <b>Route of Administration</b> | TOPICAL        |                           |               |

#### Active Ingredient/Active Moiety

| Ingredient Name  | Basis of Strength | Strength         |
|--|-------------------|------------------|
| PERACETIC ACID (UNII: I6KPI2E1HD) (PERACETIC ACID - UNII:I6KPI2E1HD) | PERACETIC ACID    | 8.0 g in 4000 mL |

**Inactive Ingredients**

| Ingredient Name                       | Strength |
|---------------------------------------|----------|
| <b>Water</b> (UNII: 059QF0KO0R)       |          |
| <b>Acetic acid</b> (UNII: Q40Q9N063P) |          |

**Packaging**

| # | Item Code        | Package Description                                       | Marketing Start Date | Marketing End Date |
|---|------------------|---|----------------------|--------------------|
| 1 | NDC:81803-020-02 | 1 in 1 CARTON   | 11/01/2021           |                    |
| 1 | NDC:81803-020-01 | 4000 mL in 1 CONTAINER; Type 0: Not a Combination Product |                      |                    |

**Marketing Information**

| Marketing Category       | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|--------------------------|--|----------------------|--------------------|
| unapproved drug<br>other |  | 11/01/2021           |                    |

**Labeler** - Valley Inc. (695095445)**Registrant** - Valley Inc. (695095445)**Establishment**

| Name        | Address | ID/FEI    | Business Operations    |
|-------------|---------|-----------|------------------------|
| Valley Inc. |         | 695095445 | manufacture(81803-020) |

Revised: 11/2021

Valley Inc.