CYGZYME PLUS- 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.

ACTIVE INGREDIENT

Peracetic acid 0.05%

INACTIVE INGREDIENTS

Water, Lauramine oxide, Sodium bicarbonate, Sodium acetate, anhydrous, Subtilisins, Cellulase, Rizolipase, alpha Amylase, Brilliant Blue, Alcohols, C12-14, ethoxylated, propoxylated, Sodium sesquicarbonate

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

Keep out of reach of children

Uses

Helps prevent sterilant

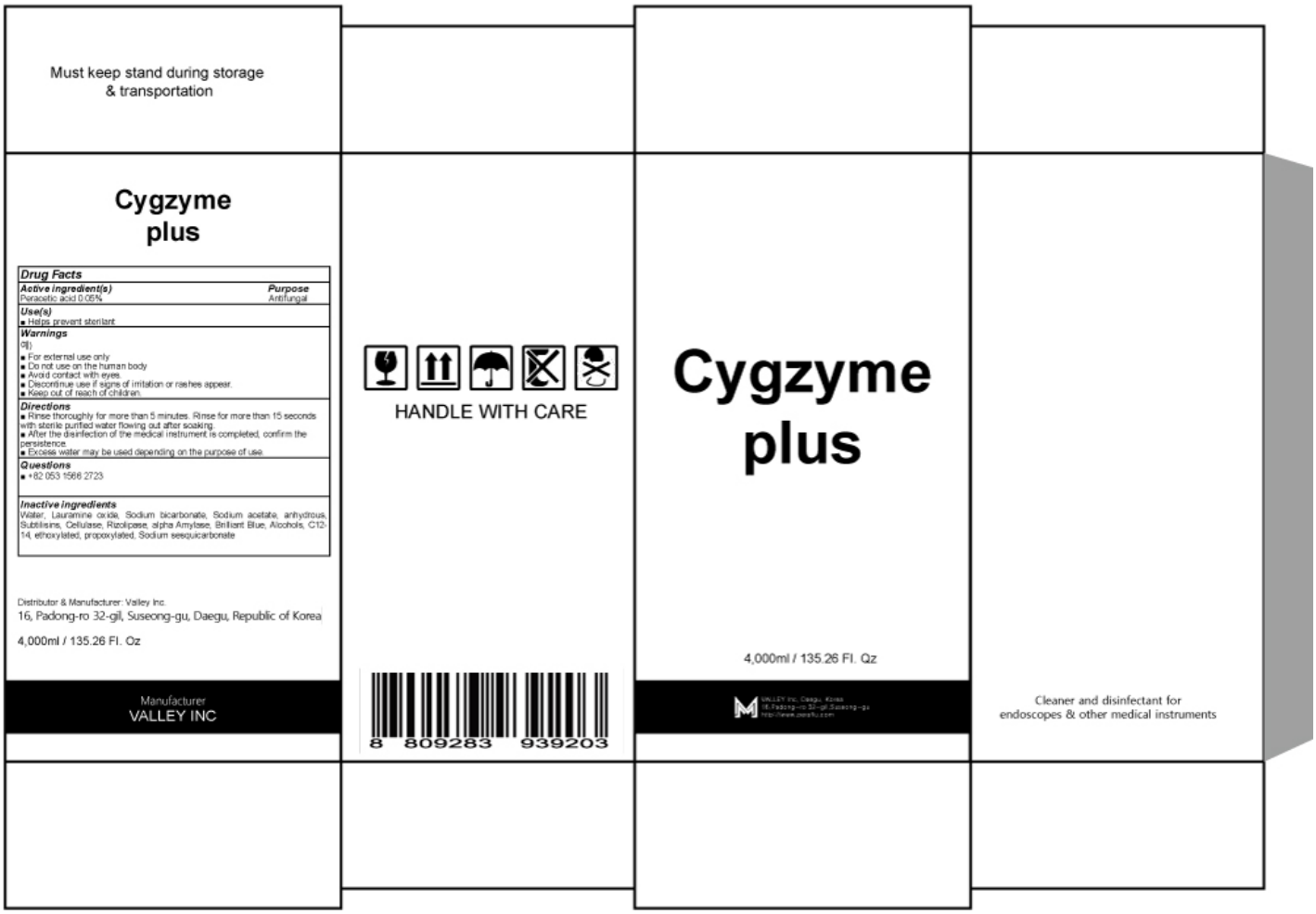
Directions

■ 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

PACKAGE LABEL.PRINCIPAL DISPLAY PANEL



CYGZYME PLUS

peracetic acid liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:81803-050
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
PERACETIC ACID (UNII: I6KPI2E1HD) (PERACETIC ACID - UNII:I6KPI2E1HD)	PERACETIC ACID	0.05 g in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**
Water (UNII: 059QF0KO0R)	
Lauramine oxide (UNII: 4F6FC4MI8W)	

Sodium bicarbonate (UNII: 8MDF5V39QO)	
SODIUM ACETATE (UNII: 4550K0SC9B)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:81803-050-02	1 in 1 CARTON	02/01/2024	
1	NDC:81803-050-01	4000 mL in 1 BOTTLE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		02/01/2024	

Labeler - Valley Inc. (695095445)

Registrant - Valley Inc. (695095445)

Establishment

Name	Address	ID/FEI	Business Operations
Valley Inc.		695095445	manufacture(81803-050)

Revised: 3/2024 Valley Inc.