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.

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

Must keep stand during storage & transportation				
String Facts Anniverse Anniverse Anniverse Anniverse CHYPACTET ACID GIPS. Anniverse CHYPACT ACID GIPS. Anniverse CHYPACTET ACID GIPS. Anniverse CHYPACTET A		Peraflu D		
Manufacturer VALLEY INC. 4,000ml / 135.26 R. Cx Manufacturer VALLEY INC	CH3 – C(= 0)OOH 0.2% This is a second of the control of the contr	4,000ml VALEY Inc., Desgoi, Konst It is delayer to 31 or Subsensor and It is clearly small blockers.	Cleaner and disinfectant for endoscopes & other medical instruments	
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peracetic acid liquid

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:81803-020	
Route of Administration	TOPICAL			

Active Ingredient/Active Moiety				
Ingredient Name	Basis of Strength	Strength		
PERACETIC ACID (UNII: 16KP12E1HD) (PERACETIC ACID - UNII:16KP12E1HD)	PERACETIC ACID	8.0 g in 4000 mL		

Inactive Ingredients				
Ingredient Name	Strength			
Water (UNII: 059QF0KO0R)				
Acetic acid (UNII: Q40Q9N063P)				

ı	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 other		11/01/2021	
		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.