VAXOL PURI- phenoxyethanol liquid
OPENKOREA CO., Ltd

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

Phenoxyethanol 1.5%

Purpose

Antiseptic

Use

For use on clothing or people, Spray it at home, at school, at the office, Stores and spaces with a large floating population, Target requiring sterilization and deodorization.

Warnings

Check the label before use, Do not throw or drop, People with damaged skin should be careful not to come into contact for a long time, Don't drink.

Do not use

On open skin wounds, Do not drink.

When using this product

If it gets into your eyes Rinse immediately with water.

Ask Doctor

If skin irritation or red spots appear, seek medical treatment. If you eat or swallow, take first aid and consult your doctor immediately.

Keep out of reach of children

Be careful not to put it in your child's mouth.

Directions

Spray on a target that requires sterilization and deodorization, Use for air sterilization in

crowded places, Use for children's supplies, toys and clothing, Use in stinking or unsanitary spaces.

Inactive ingredient

Purified Water, Hexanediol, d-Limonene, alpha-Pinene, gamma-Terpinene, para-Cymene

Package Label

VAXOL PURI
phenoxyethanol liquid

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:81268-201
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
PHENOXYETHANOL (UNII: HIE492ZZ3T) (PHENOXYETHANOL - UNII:HIE492ZZ3T)	PHENOXYETHANOL	1.5 mg in 100 mL

Inactive Ingredients

Ingredient Name	Strength
.GAMMA.-TERPINENE (UNII: 4YGF4PQP49)	
WATER (UNII: 059QF0KO0R)	
P-CYMENE (UNII: 1G1C8T1N7Q)	
HEXANEDIOL (UNII: ZIA319275I)	
.ALPHA.-PINENE (UNII: JPF3YI7O34)	
LIMONENE, (+)- (UNII: GFD7C86Q1W)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:81268-201-01	50 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	12/17/2020	
2	NDC:81268-201-02	500 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	12/17/2020	
3	NDC:81268-201-03	2000 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	12/17/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		12/17/2020	

Labeler - OPENKOREA CO., Ltd (695742727)

Registrant - OPENKOREA CO., Ltd (695742727)

Establishment

Name	Address	ID/FEI	Business Operations
OPENKOREA CO., Ltd		695742727	manufacture(81268-201)

Revised: 11/2021 OPENKOREA CO., Ltd