ALLVIRUSU- hypochlorous acid liquid StarParks

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.

80767-201

Active ingredient[s]

Hypochlorous acid (0.002%)

Purpose

Antiseptic

Use[s]

To reduce microorganism

Warnings

For external use only.

When using this product

do not drink.

Ask a doctor

if you drank a lot.

Keep out of reach of children.

Directions

Spray, pump or drop adequate amount, and let it dry or remove it in 1 minute.

Other information

Avoid direct sunlight, and store in a cool and dry place.

Inactive ingredients

Package Label

3종 국가허가

■식품첨가물 [식약처] ■일반소독제 [환경부] ■탈취제 [환경부]

미국환경청무해성인증: HOCL(차아염소산) 인증

대 부 안 심 보 습: 피부(pH5)보다 중성, 피부트러블(X) 품 질 보 존 력: 2년 장기보존 각 종 세 균 소 독: HOCL 25ppm 이상, 바이러스 등 각종

미생물 강력소독

취 상 쾌: 끈적임(X), 수돗물 수준 깔끔, 무거품 용 편 리: 펌프식, 스프레이식, 10배 수돗물과 혼합사용, 헹굼 물로도 사용

인 체 생 성 소 독 제 : 세균 침입 시 인체 백혈구가 HOCL을

생산하여 살균 경 : 살균작용 후 물로 변함, 발암물질(X)

☑NO알콜 ☑NO락스 ☑피부안심 ☑탈취



FOOD ADDITIVE

Active ingredient[s] Hypo chlorous acid (0.002%) To reduce microorganism Warnings For external use only. When using this product, do not drink. Ask a doctor if you drank a lot. Keep out of reach of children.

Directions

Drug Facts

Spray, pump or drop adequate amount, and let it dry or remove it in 1 minute.

Slightly acidic premium disinfectant

Purpose

Other information

Avoid direct sunlight, and store in a cool and dry place.

Inactive ingredients Water (99.99%)













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ALLVIRUSU

hypochlorous acid liquid

Product Information

NDC:80767-201 **Product Type** HUMAN OTC DRUG **Item Code (Source)**

TOPICAL Route of Administration

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
HYPOCHLOROUS ACID (UNII: 712K4CDC10) (HYPOCHLOROUS ACID - UNII:712K4CDC10)	HYPOCHLOROUS ACID	0.002 g in 100 mL

Inactive Ingredients Ingredient Name Strength WATER (UNII: 059QF0KO0R)

Packaging						
#	Item Code	Package Description	Marketing Start Date	Marketing End Date		
1	NDC:80767-201- 01	500 mL in 1 BOTTLE; Type 0: Not a Combination Product	10/12/2020			

Marketing Information					
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date		
unapproved drug other		10/12/2020			

Labeler - StarParks (688505276)

Registrant - StarParks (688505276)

Establishment						
Name	Address	ID/FEI	Business Operations			
StarParks		688505276	manufacture(80767-201)			

Revised: 12/2022 StarParks