

PURISTIC- sodium chlorite liquid
MEDI K 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.

82422-001 PURISTIC

Active Ingredients

Sodium chlorite 4.6%

Purpose

Disinfecting Agent

Uses

- remove 99.9% of viruses, bacteria & fungi
- Deodorizes the immediate area round it

Warning

For external use only

- If swallowed, get medical help or contact a Poison Control Center right away.

When using this product

- if following abnormal symptoms persist, discontinue use

Do not use

- in children less than 2 months of age
- on open skin wounds

■ Do not use the product for a long time in the same area as swelling, inflammation or sickness may occur due to absorption through the skin. It is not recommended to use this one area that have been medically treated with a cast or bandage.

- Do not use in combination with soap or antibacterial cleansing agents.

■ Stop immediately and consult a doctor if you experience

- 1) Hypersensitivity symptoms such as erythema, itching and dermatitis.
- 2) Skin Irritation

3) Following Instructions when using medication

(1) For external use only (Do not use internally)

(2) Avoid getting into the eyes (if contact occurs, wash well with clean water)

■ Be careful not to inhale or use excessively for a long time (ingesting ethanol repeatedly causes irritation to mucous membranes and headaches or other symptoms may appear. When used repeatedly in the same area, skin irritation may occur.

■ Do not use the product for a long time in the same area as swelling, inflammation or sickness may occur due to absorption through the skin.

It is not recommended to use this one area that have been medically treated with a cast or bandage.

- in children less than 2 months of age

Directions

■ tap lightly downward, gently bend the dot of ampoule

■ content turn yellow, to activate and emit through the body surface

■ content turn white, to be discarded in regular trash

Other information

- read the directions and warnings before use

- avoid freezing and excessive heat above 40 degree C (104 degree F)

Inactive Ingredients

Water, Citric acid

Package Label

■유통기한: 2021.02
■제조일자: 2021.02

■품 목: 일반생활화학제품(소독제, 탈취제) ■종 류: 훈증형, 개방공간 ■모델명: 퓨리스틱(팜이톡) ■생산국명: 대한민국
■판매원 및 주소: (주)메디케이 / 광주광역시 북구 첨단과기로 313, 광주하이테크센터 A동 203,204호 ■고객센터: 1833-7208
■제조회사 및 주소: (주)푸르고팜 / 경기 용인시 처인구 모현읍 새래로 116
■성분(기능): 아염소산나트륨(반응제: 2m), 시트르산(구연산)(반응제: 4m) ■유효성분 : 이산화염소(항균, 살균, 탈취) ■용량: 6ml
-화학물질의 등록 및 평가 등에 관한 법률에 의한 위해우려제품 안전기준에 적합함
-자가검사번호 -소독제: C-B01B-H00010001-A151 -탈취제: C-A10B-H00130001-A151
-자가검사번호는 이 제품이 소독제, 탈취제 안전기준에 적합하였음을 의미하며, 타 용도와는 무관합니다.



이주 메디케이

PURISTIC
코로나 바이러스 제거
Sterilizing Disinfectant

■Product Type: Disinfectant, Deodorant ■Model Name: Puristic
■Seller: MEDI K Co.,Ltd A-203, A-204, Gwangju Hightech center, 313 Cheomdangwagi-ro, Buk-gu, Gwangju, Korea
■Tel: +82-1833-7208 ■Manufacturer: PURGOFARM Co, Ltd 116, SaeRaeRo, Cheningu Yongin, GyeongGi, Korea
■Ingredients: Chlorine Dioxide ■Packaging Material: LDPE(REACH Certified) ■Date of Manufacture: Faburary 2021
■PURISTIC is the Consumer Commodity sold as the disinfectant and deodorant which has the certificate of free sales from The Korea Conformity Laboratories (KCL) ■Expiry Date: 5yrs from MFG

Drug Facts

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1) Hypersensitivity symptoms such as erythema, itching and dermatitis. 2) Skin Irritation 3) Following Instructions when using medication (1) For external use only (Do not use internally) (2) Avoid getting into the eyes (if contact occurs, wash well with clean water) ■ Be careful not to inhale or use excessively for a long time (ingesting ethanol repeatedly causes irritation to mucous membranes and headaches or other symptoms may appear. When used repeatedly in the same area, skin irritation may occur. ■ Do not use the product for a long time in the same area as swelling, inflammation or sickness may occur due to absorption through the skin. It is not recommended to use this one area that have been medically treated with a cast or bandage. ■ Do not use in combination with soap or antibacterial cleansing agents. ■ If swallowed, get medical help or contact a Poison Control Center right away.

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Questions or Comments?

Purpose

Antimicrobial

PURISTIC

sodium chlorite liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:82422-001
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
SODIUM CHLORITE (UNII: G538EBV4VF) (CHLORITE ION - UNII:Z63H374SB6)	SODIUM CHLORITE	4.6 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0K00R)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:82422-001-01	6 mL in 1 BOTTLE; Type 0: Not a Combination Product	11/27/2021	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		11/27/2021	

Labeler - MEDI K Co., Ltd. (695158681)

Registrant - MEDI K Co., Ltd. (695158681)

Establishment

Name	Address	ID/FEI	Business Operations
MEDI K Co., Ltd.		695158681	manufacture(82422-001)

Revised: 11/2021

MEDI K Co., Ltd.