PROVTECT HAND SANITIZER- chlorine dioxide spray
Ignal

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

Chlorine Dioxide 0.01%

INACTIVE INGREDIENT

Water

PURPOSE

Antimicrobial

WARNINGS

For external use only. Avoid direct sunlight and store in a cool place.

Do not use
• in children less than 2 months of age
• internal wound • inhalation by fumigation

When using this product keep out of eyes, ears, and mouth. In case of contact with eyes, rinse eyes thoroughly with water.

Stop use and ask a doctor if irritation or rash occurs. These may be signs of a serious condition.

Keep out of reach of children. If nauseous feeling or sickness occurs, when swallowed, get medical help or contact a Poison Control Center right away.

KEEP OUT OF REACH OF CHILDREN

If nauseous feeling or sickness occurs, when swallowed, get medical help or contact a Poison Control Center right away.

Uses

■ Hand Sanitizer help to reduce germs and inactivate viruses that potentially can cause disease.

Directions

■ Spray on or Wet palms, rub hands thoroughly together until dry.
■ Supervise children under 6.

Other information

■ Store between 5-20C (41-68F)
■ Avoid freezing and excessive heat above 40C (104F)

Questions?

Questions 8231-717-2562
www.ignal.co.kr
ignal.co.kr@gmail.com

PACKAGE LABEL.PRINCIPAL DISPLAY PANEL

PROVTECT HAND SANITIZER

chlorine dioxide spray

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:81754-010
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
Chlorine Dioxide (UNII: 8061YMS4RM) (CHLORINE DIOXIDE - UNII:8061YMS4RM)	Chlorine Dioxide	0.05 g in 500 mL

Inactive Ingredients

Ingredient Name	Strength
Water (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:81754-010-01	500 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	04/01/2021	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		04/01/2021	

Labeler - Ignal (695036204)

Registrant - Ignal (695036204)

Establishment

Name	Address	ID/FEI	Business Operations
Ignal		695036204	manufacture(81754-010)

Revised: 4/2021 Ignal