

BYE ZERO- magnesium liquid
Korea Life Science

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.

Sodium Chloride

Kelp extract, Seaweed extract, magnesium extract, calcium extract, potassium extract, sulfate ion extract, water

Hand sanitizer to help reduce bacteria that potentially can cause disease. for use when soap and water are not available.

keep out of reach of the children

place enough product on hadns to cover all surfaces. rub hands together until dry.

supervise children uner 6 years of age when using this product to avoid swallowing.

Stop use and ask a doctor if irritation or rash occurs.

Keep out of reach of children.

for external use only

Drug Facts	
Active Ingredient	Purpose
Sodium Chloride 0.99%	Antiseptic
Uses(용도)	
■ hand sanitizer to help reduce bacteria that potentially can cause disease. For use when soap and water are not available	
Warnings(경고)	
■ stop use and ask a doctor if irritation or rash occurs. ■ keep out of reach of children.	
Directions	
■ place enough product on hands to cover all surfaces. rub hands together until dry. ■ supervise children under 6 years of age when using this product to avoid swallowing	
Other Information	
■ store between 10-35C (50-95F) ■ avoid freezing and excessive heat above 40C(104F)	
Inactive Ingredients	
Calcium, Kelp, Magnesium, Potassium, Seaweed, Sulfate	
Questions or Comments? Call 82-10-3585-8580 or email: biotoc@naver.com	

BYE ZERO
magnesium liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:80762-0020
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
MAGNESIUM (UNII: I38ZP9992A) (MAGNESIUM - UNII:I38ZP9992A)	MAGNESIUM	4.95 g in 500 mL

Inactive Ingredients

Ingredient Name	Strength
NORI (UNII: 477TV3P5UX)	
CALCIUM (UNII: SY7Q814VUP)	
POTASSIUM (UNII: RWP5GA015D)	
SULFATE ION (UNII: 7IS9N8KPMG)	
WATER (UNII: 059QF0K00R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:80762-0020-1	90 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	06/20/2023	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		10/06/2020	

Labeler - Korea Life Science (695504093)**Registrant** - Korea Life Science (695504093)**Establishment**

Name	Address	ID/FEI	Business Operations
Korea Life Science		695504093	manufacture(80762-0020)

Revised: 10/2023

Korea Life Science