

HOO GARGLE (APPLE SPEARMINT FLAVOR)- sodium flouride liquid
MIKO CO LTD

Drug Facts

sodium fluoride

menthol

xylitol

Sodium acetate

Allanine

Acetic acid

Propolis

Chitosan

Green tea extract

Stevia

Cyclodextrin

Benzoic acid

Red ginsengs extract

Caramel

Grapefruit seed extract

Maple flavor

Apple spearmint flavor

Purified Water

dental care

KEEP OUT OF REACH OF THE CHILDREN

1. Put one cup in mouth.
2. Gargle for 30 sec ~ 1 minute
3. Spit it out.
5. Rinse with water.

Be careful not to swallow it, and rinse your mouth thoroughly after use.

dental use only

Drug Facts

Sodium fluoride 0.02% (0.01% w/v Fluoride Ion)

Purpose

Anticavity, Remove bad breath

Use

Aids in the prevention of dental cavities

Warnings

Keep out of reach of children.

If more than used for rinsing is accidentally swallowed, get medical help or contact a Poison Control Center right away

Directions

- * Adults and children 6 years of age and older
- * Use before or after brushing your teeth with a toothpaste or use whenever you want but, do not use more than 3 times a day
- * Vigorously swish 10ml (2 teaspoonfuls) of rinse between your teeth for 1 minute and then spit out
- * Do not swallow the rinse
- * Instruct children under 12 years of age in good rinsing habits (to minimize swallowing)
- * Supervise children as necessary until capable of using without supervision
- * Children under 6 years age : consult a dentist or doctor

Other Information

Store at room temperature

Inactive ingredients

Water, L-Menthol, Xylitol, Sodium acetate, Allanine, Acetic acid, Propolis, Chitosan, Green tea extract, Stevia, Cyclodextrin, Benzoic acid, Red ginseng extract, Caramel, Grapefruit seed extract, Maple flavor, Apple Spearmint flavor

ALCOHOL FREE

Made in Korea

Questions?

Call toll-free +1 213-908-5154

Distributed By Flax-Protection Health&Beauty
1331 W 7th street Los Angeles CA 90017

Expiration Date : AUG 17 2025



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HOO GARGLE (APPLE SPEARMINT FLAVOR)

sodium flouride liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:81840-3333
Route of Administration	DENTAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
SODIUM FLUORIDE (UNII: 8ZYQ1474W7) (FLUORIDE ION - UNII:Q80VPU4080)	SODIUM FLUORIDE	0.02 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:81840-3333-5	100 in 1 PACKAGE	09/20/2022	
1	NDC:81840-3333-4	50 in 1 PACKAGE		
1	NDC:81840-3333-3	30 in 1 PACKAGE		
1	NDC:81840-3333-2	15 in 1 PACKAGE		
1	NDC:81840-3333-1	10 in 1 PACKAGE		
1	NDC:81840-3333-0	10 mL in 1 POUCH; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M021	01/01/2022	

Labeler - MIKO CO LTD (689973854)

Registrant - MIKO CO LTD (689973854)

Establishment

Name	Address	ID/FEI	Business Operations
MIKO CO LTD		689973854	manufacture(81840-3333)

Revised: 10/2023

MIKO CO LTD