

**DENTI WHOO BLUE TABLETGARGLE- sodium fluoride tablet**  
**SUN LIFE SCIENCE 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.*

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Sodium Fluoride

Sodium Bicarbonate

Tartaric Acid

Hydroxypropylcellulose

Hydroxypropyl methylcellulose

Silica

Zea Mays (Corn) Starch

Sodium Cocoyl Glutamate

Maltitol

Mannitol

Erythritol

Potassium Acesulfame

Magnesium Stearate

Brilliant Blue

Menthol

Yucca Vera Leaf/Root Extract

fragrance

Tooth-decay prevention, Odor removal from the mouth and Oral Cleansing

Keep out of reach of children

USAGE : Put 1 tablet in the mouth and use the tongue and lips to rub the gums, teeth, and tongue, gargle it with water, and rinse it several times

- 1) Observe the specified usage capacity.
- 2) Store in a cool place with no sunlight and less moisture.
- 3) Avoid swallowing.
- 4) This product contains 583 ppm of fluorine.
- 5) This product contains fluorine, so if you swallow a large amount, consult a doctor or dentist immediately.
- 6) When used by children under the age of 6, use under the supervision of their guardians to avoid swallowing.
- 7) Keep out of reach of children under the age of 6

For dental use only



## DENTI WHOO BLUE TABLET GARGLE

sodium fluoride tablet

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:86157-0014
<b>Route of Administration</b>	DENTAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>SODIUM FLUORIDE</b> (UNII: 8ZYQ1474W7) (FLUORIDE ION - UNII:Q80VPU408O)	FLUORIDE ION	1.10037 g

### Inactive Ingredients

Ingredient Name	Strength
<b>TARTARIC ACID</b> (UNII: W4888119H)	
<b>MALTITOL</b> (UNII: D65DG142WK)	
<b>SODIUM BICARBONATE</b> (UNII: 8MDF5V39QO)	

## Product Characteristics

<b>Color</b>	white	<b>Score</b>	no score
<b>Shape</b>	ROUND	<b>Size</b>	15mm
<b>Flavor</b>	MENTHOL	<b>Imprint Code</b>	none
<b>Contains</b>			

## Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:86157-0014-1	8 in 1 PACKAGE; Type 0: Not a Combination Product	04/06/2022	

## Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		04/06/2022	

**Labeler** - SUN LIFE SCIENCE CO LTD (695149648)

**Registrant** - SUN LIFE SCIENCE CO LTD (695149648)

## Establishment

Name	Address	ID/FEI	Business Operations
SUN LIFE SCIENCE CO LTD		695149648	manufacture(86157-0014)

Revised: 5/2022

SUN LIFE SCIENCE CO LTD