STRAWBERRY FLAVOR SODIUM FLUORIDE ANTICAVITY TOOTHPAST- sodium fluoride gel HISMILE PTY LTD

Strawberry Flavor Sodium Fluoride Anticavity Toothpaste

Active ingredient

Sodium Fluoride 0.243% (0.14% w/v fluoride ion)

Purpose

Anticavity

Use

Aids in the prevention of dental cavities.

Warnings

Warnings

When using this product

- Do not swallow
- Do not use in eye area

Keep out of reach of children under 6 years

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

Directions

- Adults and children 2 years of age and older: Brush teeth thoroughly, preferably after each meal or at least twice a day, or use as directed by a dentist or doctor.
- Instruct children under 6 years of age in good brushing and rinsing habits (to minimize swallowing).
- Supervise children as necessary until capable of using without supervision.
- Children under 2 years of age: Consult a dentist or doctor.

Inactive ingredients

Water

Glycerin

Hydrated Silica Xylitol Sorbitol Cellulose Gum Flavor 1,2-Hexanediol Sodium Phosphate Sodium Lauroyl Sarcosinate Tetrapotassium Pyrophosphate Sucralose Caprylyl Glycol Lauryl Glucoside

Methyl Diisopropyl Propionamide

Sodium Gluconate

Zinc Lactate

Red 33

Label





SODIUM FLUORIDE ANTICAVITY TOOTHPASTE Net wt. 3.5 oz (100g)

Drug Facts

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Sodium Fluoride 0.243% (0.14% w/v fluoride ion) Anticavity

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- hildren under 2 years of age: Consult a dentist or doctor

Inactive ingredients Water, Glycerin, Hydrated Silica, Xvlitol, Sorbitol, Cellulose Gum, Flavor, 1,2-Hexanediol, Sodium Phosphate, Sodium Lauroyl Sarcosinate, Tetrapotassium Pyrophosphate, Sucralose, Caprylyl Glycol, Lauryl Glucoside, Methyl Diisopropyl Propionamide, Sodium Gluconate, Zinc Lactate, Red 33.

Questions or comments?

+1-302-440-3558

You may also report serious side effects to this phone number.

Distributed by: Waters, QLD 4220 AU.

Made in USA from globally sourced ingredients and components

STRAWBERRY FLAVOR SODIUM FLUORIDE ANTICAVITY TOOTHPAST

sodium fluoride gel

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:83013-339

Route of Administration TOPICAL

Active Ingredient/Active Moiety Ingredient Name Basis of Strength Strength

SODIUM FLUORIDE (UNII: 8ZYQ1474W7) (FLUORIDE ION - UNII:Q80VPU408O)

SODIUM FLUORIDE 0.243 g in 100 g

Inactive Ingredients				
Ingredient Name	Strength			
CAPRYLYL GLYCOL (UNII: 00YIU5438U)				
METHYL DIISOPROPYL PROPIONAMIDE (UNII: 6Q0P5A9489)				
1,2-HEXANEDIOL (UNII: TR046Y3K1G)				
CELLULOSE GUM (UNII: K679OBS311)				
SODIUM LAUROYL SARCOSINATE (UNII: 632GS99618)				
RED 33 (UNII: 9DBA0SBB0L)				
XYLITOL (UNII: VCQ006KQ1E)				
SODIUM PHOSPHATE (UNII: SE337SVY37)				
LAURYL GLUCOSIDE (UNII: 76LN7P7UCU)				
WATER (UNII: 059QF0KO0R)				
ZINC LACTATE (UNII: 2GXR25858Y)				
GLYCERIN (UNII: PDC6A3C0OX)				
HYDRATED SILICA (UNII: Y6O7T4G8P9)				
SUCRALOSE (UNII: 96K6UQ3ZD4)				
SORBITOL (UNII: 506T60A25R)				
TETRAPOTASSIUM PYROPHOSPHATE (UNII: B9W4019H5G)				
SODIUM GLUCONATE (UNII: R6Q3791S76)				

Packaging					
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:83013-339- 10	100 g in 1 TUBE; Type 0: Not a Combination Product	10/31/2025		

Marketing Information					
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date		
OTC Monograph Drug	M021	10/31/2025			

Labeler - HISMILE PTY LTD (758083252)

Establishment							
Name	Address	ID/FEI	Business Operations				
Nutrix International, LLC.		117341868	manufacture(83013-339)				

Revised: 11/2025 HISMILE PTY LTD