

AP-24- sodium fluoride mouthwash
NSE Products, Inc

AP-24[®] Mouthwash

Drug Facts

Active Ingredient

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

Purpose

anticavity

Use

Aids in the prevention of dental cavities.

Warning

- **Keep out of the reach of children.** If more than used for rinsing is accidentally swallowed, get medical help or contact a Poison Control Center right away.
- **Do not use if safety seal is broken.**

Directions

- Adults and children 6 years of age and older: Use once a day after brushing your teeth with a toothpaste.
- Vigorously swish 10 milliliters of rinse between your teeth for 1 minute and then spit out.
- Do not swallow the rinse.
- Do not eat or drink for 30 minutes after rinsing.
- 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 of age: Consult a dentist or doctor.

Inactive Ingredients

Water (Aqua), Sorbitol, Glycerin, Poloxamer 338,¹ Poloxamer 407,¹ Dimethicone,¹ Sodium Phosphate, Phosphoric Acid, Disodium EDTA, Sodium Saccharin, Sodium Benzoate, Flavor (Aroma).

¹ AP-24,[®] the patented ultra-emulsion of medical grade, high molecular weight Dimethicone and the surfactants, Poloxamer 338 and Poloxamer 407, helps to remove plaque and helps reduce

plaque buildup.

Questions?

1-888-742-7626

PRINCIPAL DISPLAY PANEL - 500 ml Bottle Label

AP24®

Anti-Plaque

Fluoride

Mouthwash

NU SKIN®

500 ml e (16.9 Fl. Oz.)

AP²⁴®

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**Sold Exclusively by Nu Skin Enterprises
Authorized Distributors**

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AP-24

sodium fluoride mouthwash

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:62839-1152	
Route of Administration	ORAL			
Active Ingredient/Active Moiety				
Ingredient Name		Basis of Strength	Strength	
Sodium Fluoride (UNII: 8ZYQ1474W7) (Fluoride Ion - UNII:Q80VPU408O)		Fluoride Ion	0.2 mg in 1 mL	
Inactive Ingredients				
Ingredient Name			Strength	
Water (UNII: 059QF0KO0R)				
Sorbitol (UNII: 506T60A25R)				
Glycerin (UNII: PDC6A3C00X)				
Poloxamer 338 (UNII: F75JV2T505)				
Poloxamer 407 (UNII: TUF2IVW3M2)				
Dimethicone (UNII: 92RU3N3Y1O)				
Sodium Phosphate (UNII: SE337SVY37)				
Phosphoric Acid (UNII: E4GA8884NN)				
Edetate Disodium Anhydrous (UNII: 8NLQ36F6MM)				
Saccharin Sodium (UNII: SB8ZUX40TY)				
Sodium Benzoate (UNII: OJ245FE5EU)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:62839-1152-1	500 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/08/2017	
Marketing Information				
Marketing Category	Application Number or Monograph Citation		Marketing Start Date	Marketing End Date
OTC MONOGRAPH DRUG	M021		05/08/2017	

Labeler - NSE Products, Inc (803486393)