

AP-24 WHITENING- sodium monofluorophosphate paste
NSE Products, Inc.

AP-24® Whitening Toothpaste

Drug Facts

Active Ingredients

Sodium Monofluorophosphate 0.76% (0.15% w/v fluoride ion)

Purpose

anticavity

Use

Aids in the prevention of dental cavities.

Warning

- **Keep out of the reach of children under 6 years of age.** If more than used for brushing is accidentally swallowed, get medical help or contact a Poison Control Center immediately.

Directions

- Adults and children 2 years of age and older: Brush teeth thoroughly, preferably after each meal or at least twice a day, or as directed by a dentist or doctor.
- Instruct children under 6 years of age on 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 (Aqua), Hydrated Silica, Sorbitol, Alumina, Glycerin, Tetrasodium Pyrophosphate, PEG-12, Dimethicone¹, Poloxamer 338¹, Poloxamer 407¹, Cellulose Gum, Sodium Lauryl Sulfate, Flavor (Aroma), Sodium Saccharin, Titanium Dioxide (CI 77891).

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 during brushing and helps reduce plaque buildup.

Questions

1-888-742-7626

PRINCIPAL DISPLAY PANEL - 110 g Tube Label

AP24®

Whitening
Fluoride
Toothpaste

NU SKIN®

110 g e (4 oz.)



AP₂₄®

Whitening Fluoride Toothpaste

 NU SKIN®

110 g e (4 oz.)

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Inactive Ingredients: Water (Aqua), Hydrated Silica, Sorbitol, Aluminum Hydroxide, Glycerin, Tetrasodium Pyrophosphate, PEG-12, Dimethicone*, Poloxamer 338*, Poloxamer 407*, Cellulose Gum, Sodium Lauryl Sulfate, Flavor (Aroma), Sodium Saccharin, Titanium Dioxide (CI 77891).

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Nu Skin Enterprises
Authorized Distributors

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NSE Products, Inc., Provo, UT 84601
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nuskin.com
01 111155 90016033/7



AP-24 WHITENING

sodium monofluorophosphate paste

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:62839-1155
Route of Administration	DENTAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
Sodium Monofluorophosphate (UNII: C810JCZ56Q) (Fluoride Ion - UNII:Q80VPU408O)	Fluoride Ion	7.6 mg in 1 g

Inactive Ingredients

Ingredient Name	Strength
Water (UNII: 059QF0KO0R)	
Hydrated Silica (UNII: Y607T4G8P9)	
Sorbitol (UNII: 506T60A25R)	
Aluminum Hydroxide (UNII: 5QB0T2IUN0)	
Glycerin (UNII: PDC6A3C0OX)	
Sodium Pyrophosphate (UNII: O352864B8Z)	
Polyethylene Glycol 600 (UNII: NL4J9F21N9)	
Dimethicone (UNII: 92RU3N3Y1O)	
Poloxamer 338 (UNII: F75JV2T505)	
Poloxamer 407 (UNII: TUF2IVW3M2)	
CARBOXYMETHYLCELLULOSE SODIUM, UNSPECIFIED (UNII: K679OBS311)	
Sodium Lauryl Sulfate (UNII: 368GB5141J)	
Saccharin Sodium (UNII: SB8ZUX40TY)	
Titanium Dioxide (UNII: 15FIX9V2JP)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:62839-1155-1	110 g in 1 TUBE; 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)

