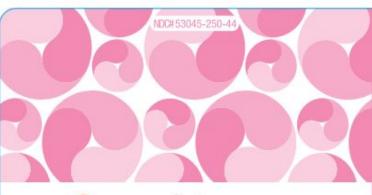
IONITE APF FOAM- sodium fluoride aerosol, foam Dharma Research, Inc.

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.

Castor Oil, Decyl Glucoside, Distilled Water, Flavor, Hydrofluoric Acid, Phosphoric Acid, Poloxamer, Propellant A31, Sodium Benzoate, Sodium Laureth Sulface, Sodium Saccharne, Triethanolamine, Xylitol

- Do not swallow.
- Keep out of reach of children.
- Contents under pressure.
- Do not place in hot water or near radiators, stoves or other sources of heat.
- Do not puncture or incinerate container. Do not spray towards open flames.
- For professional use only.
- Remove cap from can. Prior to each use, shake can thoroughly for at least 15 seconds.
- To dispense, invert the can completely upside down. Slowly depress nozzle to dispense foam into a fluoride tray (foam will expand slightly higher than fluoride tray).
- Air dry teeth thoroughly and inset tray(s) into patient's mouth. Instruct patient to bite down and leave the tray in contact with the teeth between 1 4 minutes.
- Use a saliva ejector during treatment to minimize ingestion of product.
- Remove the tray(s) and have patient expectorate.
- Instruct patient not to eat, drink or rinse for 30 minutes after treatment.

Store at a controlled room temperature 59°-86°F (15°-30°C)





APF FOAM with Xylitol

Bubble Gum

Acidulated Phosphate Fluoride Treatment Foam 1.23% Fluoride Ion

GLUTEN FREE

1 minute or 4 minute Treatment



4.4 fl.oz. (130 ml)

Drug Facts

Active Ingredient

Purpose

Sodium Fluoride 2,72%...

..Anticaries

Uses This is a prescription fluoride treatment foam used to help prevent dental decay.

Warnings

- Do not swallow.
- Keep out of the reach of children,
- Contents under pressure.
- Do not place in hot water or near radiators, stoves or other sources of heat.
- Do not puncture or incinerate container, Do not spray toward open flame,
- For Professional Use Only,

Directions

- Remove cap from can, Prior to each use, shake can thoroughly for at least 15 seconds
- To dispense, invert the can completely upside down. Slowly depress nozzle to dispense foam into a fluoride tray (foam will expand slightly higher than fluoride tray),
- Air dry teeth thoroughly and insert tray(s) into patient's mouth, Instruct patient to bite down and leave the tray in contact with the teeth between 1 - 4 minutes,
- Use a saliva ejector during treatment to minimize ingestion of product,
 Remove tray(s) and have patient expectorate.
- Instruct patient not to eat, drink or rinse for 30 minutes after treatment,

Inactive ingredients

Castor Oil, Decyl Glucoside, Distilled Water, Flavor, Phosphoric Acid, Poloxamer, Propellant A31, Sodium Benzoate, Sodium Laureth Sulfate, Sodium Saccharine, Triethanolamine, Xylitol,

Other information

■ Store at controlled room temperature 59°- 86°F (15°-30° C)

Shake well before each use

INVERT CAN COMPLETELY AND DEPRESS NOZZLE TO DISPENSE



Rev. B, Date 10/12/2020

Manufactured by:

DHARMA
RESEARCH, INC.
WWW.DHARMARESEARCH.COM

5220 N.W 72 Avenue Unit 15, Miami, FI, 33168 1-877-833-3725

Re-order#: 56-00100



MADE IN USA

IONITE APF FOAM

sodium fluoride aerosol, foam

Product Information

Product Type HUMAN PRESCRIPTION DRUG Item Code (Source) NDC:53045-250

Route of Administration DENTAL, TOPICAL, ORAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
SODIUM FLUORIDE (UNII; 8ZYQ1474W7) (FLUORIDE ION - UNII:Q80 VPU408O)	FLUORIDE ION	1.5498 g in 126 g

Inactive Ingredients		
Ingredient Name	Strength	
CASTOR OIL (UNII: D5340 Y219 G)		
DECYL GLUCOSIDE (UNII: Z17H97EA6Y)		
WATER (UNII: 059QF0KO0R)		
HYDRO FLUO RIC ACID (UNII: RGL5YE86CZ)		
PHO SPHO RIC ACID (UNII: E4GA8884NN)		
SODIUM BENZOATE (UNII: OJ245FE5EU)		
SO DIUM LAURETH SULFATE (UNII: BPV390 UAP0)		
SACCHARIN SO DIUM (UNII: SB8 ZUX40 TY)		
XYLITOL (UNII: VCQ006KQ1E)		
POLOXAMER 407 (UNII: TUF2IVW3M2)		
TROLAMINE (UNII: 903K93S3TK)		

Product Characteristics		
Color		Score
Shape		Size
Flavor	BUBBLE GUM	Imprint Code
Contains		

ı	Packaging			
ı	# Item Code	Package Description	Marketing Start Date	Marketing End Date
ı	1 NDC:53045-250-44	126 g in 1 BOTTLE; Type 0: Not a Combination Product	04/22/2013	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		04/22/2013	

Labeler - Dharma Research, Inc. (078444642)

Registrant - Dharma Research, Inc. (078444642)

Establishment				
Name	Address	ID/FEI	Business Operations	
Dharma Research, Inc.		078444642	manufacture(53045-250)	

Revised: 10/2020 Dharma Research, Inc.