NRG APF- sodium fluoride gel IQ Dental Supply, LLC

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.

Active Ingredient:	Purpose:
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Fluoride Ion 1.23%......Flouride Treatment Gel

Available from 2.09% Sodium Fluoride and Hydrofluoric Acid

Indications and Usage:

- A stable thixotropic fluoride treatment gel used to help prevent dental decay.
- For Professional Use Only. This product is not intended for home or unsupervised use.

Warnings:

- Keep out of reach of children.
- Do not swallow. If product is accidentally swallowed in quantities greater than would normally occur with a treatment gel, seek medical help or contact a Poison Control Center right away.
- Read directions carefully before using.

Dosage and Administration:

Shake well before use. This is a one minute or four minute fluoride gel for in-office patient use. It is normally used as a preventative caries treatment two times a year.

- 1. After thorough prophylaxsis, fill two single or one dual tray, one third full with gel. Air dry teeth and insert trays into the mouth.
- 2. Instruct patient to bite down lightly but firmly for one minute (80% effectiveness) or four minutes (100% effectiveness).
- 3. Remove trays. Instruct patient to expectorate any excess gel and not to eat or drink for at least 30 minutes.

Other Information:

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

Protect from freezing.

Inactive Ingredients:

Citric Acid, FD&C Green #3, FD&C Yellow #5 (tartrazine) as a color additive, Flavor, Magnesium Aluminum Silicate, Phosphoric Acid, Polysorbate 20, Sodium Benzoate, Sodium Saccharin, Titanium Dioxide, Water, Xanthan Gum, Xylitol.

NRG

Distributed by IQ Dental

APF Gel Thixotropic with Xylitol

Acidulated
Phosphate Fluoride
Treatment Gel
1.23% Fluoride Ion

GLUTEN FREE

Mint

Re-order#: NRGAPFG-MN

Ronly

IMPORTANT: READ DIRECTIONS FOR PROPER USE

MADE IN USA Net Wt. 16 oz (454 g)

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- Do not swallow. If product is accidentally swallowed in quantities greater than would normally occur with a treatment gel, seek medical help or contact a Poison Control Center right away.
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- After thorough prophylaxis, fill two single or one dual tray, one third full with gel. Air dry teeth and insert trays into the mouth.
- 2. Instruct patient to bite down lightly but firmly for one minute (80% effectiveness) or four minutes (100% effectiveness).
- 3. Remove trays. Instruct patient to expectorate any excess gel and not to eat or drink for at least 30 minutes.

Other Information:

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Inactive Ingredients:

Citric Acid, FD&C Green #3, FD&C Yellow #5 (tartrazine) as a color additive, Flavor, Magnesium Aluminum Silicate, Phosphoric Acid, Polysorbate 20, Sodium Benzoate, Sodium Saccharin, Titanium Dioxide, Water, Xanthan Gum, Xylitol.



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NRG APF

sodium fluoride gel

Product Information

Product Type HUMAN PRESCRIPTION DRUG Item Code (Source) NDC:42756-1113

Route of Administration DENTAL

Active Ingredient/Active Moiety

	Ingredient Name	Basis of Strength	Strength
l	SODIUM FLUORIDE (LINII: 87 YO 1474W7) (FLUORIDE ION - LINII: O80VPLI408O)	FLUORIDE ION	5 6 a in 454 a

Inactive Ingredients		
Ingredient Name	Strength	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)		
FD&C GREEN NO. 3 (UNII: 3P3ONR6O1S)		
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)		
MAGNESIUM ALUMINUM SILICATE (UNII: 6M3P64V0NC)		
PHOSPHORIC ACID (UNII: E4GA8884NN)		
POLYSORBATE 20 (UNII: 7T1F30V5YH)		
SODIUM BENZOATE (UNII: OJ245FE5EU)		
SACCHARIN SODIUM (UNII: SB8ZUX40TY)		
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)		
WATER (UNII: 059QF0KO0R)		
XANTHAN GUM (UNII: TTV12P4NEE)		
XYLITOL (UNII: VCQ006KQ1E)		

Product Characteristics		
Color	green	Score
Shape		Size
Flavor	MINT	Imprint Code
Contains		

P	Packaging			
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
		454 g in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	08/01/2013	

Marketing Information				
Application Number or Monograph Citation	Marketing Start Date	Marketing End Date		
	08/01/2013			
	Application Number or Monograph	Application Number or Monograph Marketing Start Citation Date		

Labeler - IQ Dental Supply, LLC (800349763)

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