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:

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 Blue #1, FD&C Red#3, Flavor, Magnesium Aluminum Silicate, Phosphoric Acid, Polysorbate 20, Sodium Benzoate, Sodium Saccharin, Titanium Dioxide, Water, Xanthan Gum, Xylitol.

NRG

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APF Gel Thixotropic with Xylitol

Acidulated Phosphate Fluoride Treatment Gel 1.23% Fluoride Ion

GLUTEN FREE

Grape

Re-order#: NRGAPFG-GR

Ronly

IMPORTANT: READ DIRECTIONS FOR PROPER USE

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

Distributed by IQ Dental				
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 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 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: • Store at controlled room temperature 59° to 86°F (15°-30°C). Protect from freezing.				
Inactive Ingredients: Citric Acid, FD&C Blue #1, FD&C Red #3, 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						
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Product Information						
Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)		NDC:	NDC:42756-1114	
Route of Administration	DENTAL					
Active Ingredient/Active	Majatu					
Active Ingredient/Active	Molety					
Ingre	Basis of Strength		Strengtl	h		
SODIUM FLUORIDE (UNII: 8ZYQ14	FLUORIDE ION		5.6 g in 454	4 g		

	edients				Strength		
Ingredient Name							
CITRIC ACID MO							
FD&C BLUE NO.	· ·						
FD&C RED NO. 3							
		E (UNII: 6M3P64V0NC)					
PHOSPHORIC AC							
POLYSORBATE 2	· ·	•					
SODIUM BENZO							
SACCHARIN SOD							
		232)					
WATER (UNII: 059QF0KO0R) XANTHAN GUM (UNII: TTV12P4NEE)							
		1					
XYLITOL (UNII: VO							
Product Cha	racteristics						
Color		purple	Score				
Shape		parpie	Size				
Flavor		GRAPE	Imprint Code				
Contains							
contains							
Packaging							
					Marketing En		
# Item Code	Pa	ckage Description		Marketing Start Date	Date		
1 NDC:42756-		LE, PLASTIC; Type 0: N	Not a	08/01/2013			
1114-7	L114-7 Combination Product			00/01/2015			
Marketing	Informat	ion					
Marketing Category	Applicat	tion Number or Monograph Citation		Marketing Start Date	Marketing End Date		
unapproved drug				08/01/2013			

Labeler - IQ Dental Supply, LLC (800349763)

Revised: 1/2022

IQ Dental Supply, LLC