## GELATO APF- sodium fluoride gel Mycone Dental Supply Co., Inc DBA Keystone Industries and Deepak Products 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.

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### Active ingredient:

**Purpose:** 

Fluoride Ion 1.23% ...... Fluoride 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 Office Use Only. This product is not intended for home or unsupervised consumer 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 twice 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:

• Do not store above 25°C/77°F. Do not freeze.

# Inactive ingredients:

Citric Acid, Flavor, Magnesium Aluminum Silicate, Phosphoric Acid, Polysorbate 20,

Sodium Benzoate, Sodium Saccharin, Titanium Dioxide, Water, Xanthan Gum, Xylitol. May contain blue #1, green #3, red #3, red #40, yellow #5 (tartrazine), yellow #6 as a color additive.



## **GELATO APF**

sodium fluoride gel

Product Information			
Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:68400-106
Route of Administration	DENTAL		

<b>Active</b>	Ingredient/Active	Moiety

Ingredient Name	Basis of Strength	Strength
marealent Name	basis of strength	Juengui

**SODIUM FLUORIDE** (UNII: 8ZYQ1474W7) (FLUORIDE ION - UNII:Q80VPU408O) FLUORIDE ION

5.6 g in 454 g

Inactive Ingredients				
Ingredient Name	Strength			
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)				
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)				
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)				
FD&C GREEN NO. 3 (UNII: 3P3ONR6O1S)				
FD&C RED NO. 3 (UNII: PN2ZH5LOQY)				
FD&C RED NO. 40 (UNII: WZB9127XOA)				
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)				
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)				

Product Characteristics				
Color	red	Score		
Shape		Size		
Flavor	CHERRY	Imprint Code		
Contains				

l	Packaging				
	#	Item Code	tem Code Package Description		Marketing End Date
	1	NDC:68400- 106-15	454 g in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	06/01/2016	

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
unapproved drug other		12/14/2012		
outer				

(014769301)

Establishment			
Name	Address	ID/FEI	Business Operations
Keystone Industries		014769301	manufacture(68400-106) , label(68400-106)

Revised: 1/2022 Mycone Dental Supply Co., Inc DBA Keystone Industries and Deepak Products Inc.