GELO-X- sodium fluoride gel, dentifrice Dharma Research, Inc.

Storage and Handling

- Store at 59° 86° F (15° 30° C)
- Protect from freezing
- Product is free of gluten, soy milk, egg, peanut, and tree nut products

Inactive Ingredients

Citric acid, FD&C Red No. 40 (C.I. 16035), Flavor, Hydrofluoric Acid, Magnesium Aluminum Silicate, Phosphoric Acid, Polysorbate 20, Purified Water, Sodium Benzoate, Sodium Saccharine, Sweetness enhancer, Titanium dioxide, Tocopherol acetate, Xanthan gum, Xylitol

Warnings and Precautions

- •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
- •Do not use on patients with an allergy to fluoride
- Product contains artificial color; confirm that no known patient allergies exist
- For professional use only

Instructions for Use

- Shake well before use
- •This is a four minute or one minute fluoride gel for in-office patient use.
- •For best results, use at least twice per year
- •After thorough prophylaxis, fill two single or one dual tray one third full with gel. Air dry teeth and insert trays into mouth.
- •Instruct patient to bite down lightly but firmly for one minute (80% effectiveness) or four minutes (100% effectiveness)
- •Remove trays, instruct patient to expectorate any excess gel and not to eat or drink for at least 30 minutes



APF THIXOTROPIC GEL with Xylitol & Vitamin E

Acidulated Phosphate Fluoride Treatment Gel 1.23% Fluoride Ion

GLUTEN FREE

1 minute or 4 minute Treatment



17 fl. oz. (500ml)

Drug Facts

Active Ingredient

Purpose

Uses Prescription fluoride treatment gel used to prevent dental decay.

- Keep out of the reach of children.
- Do not swallow, If product is accidentally swallowed in quantities greater ■ Do not swarrow. In product is accodentally swarrowed in quantities great than would normally occur with a treatment gel, seek medical help or contact a Poison Control Center right away.

 ■ Do not use on patients with an allergy to fluoride.

 ■ Ionitie APF Gel contains artificial color, confirm that no known patient
- allergies exist.
- For professional use only.

Directions

- Shake well before use.
 This is a four minute or one minute fluoride gel for in-office patient use.
 For best results, use at least twice a year.
 After thorough prophylaxis, fill two single or one dual tray one third full with gel. Air dry teeth and insert trays into the mouth.
 Instruct patient to bite down lightly but firmly for one minute (80% effectiveness) or four minutes (100% effectiveness) or four minutes (100% effectiveness).
- Remove trays, instruct patient to expectorate any excess gel and not to eat or drink for at least 30 minutes.

Other information

- Store at 59° 86°F (15°-30° C),
 Protect from freezing,
 lonite APF Gel is free of gluten, soy milk, egg, peanut and tree nut products

Inactive ingredients

Citric Acid, FD&C Red No. 40 (C.I. 16035), Flavor, Hydrofluoric Acid, Wagnesium Aluminum Silicate, Phosphoric Acid, Polysorbate 20, Purified Water, Sodium Benzoate, Sodium Saccharin, Sweetness Enhancer, Titanium Dioxide, Tocopheryl Acetate, Xantham Gum, Xylitol.

Importer and License Holder: Dentorient Fuss LTD 512834284 P.O.B 2232 Tel Aviv Israel Tel 03-6393640 Fax 03-6393645 www.idental.co.il Israeli Registration Number: 25360003 (medical device)

reorder code: gelox002

expiry date and batch printed on the bottom

Strength

MADE IN USA

GELO-X

sodium fluoride gel, dentifrice

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:53045-209

Route of Administration DENTAL

Active Ingredient/Active Moiety Basis of Ingredient Name Strength Strength

SODIUM FLUORIDE (UNII: 8ZYQ1474W7) (FLUORIDE ION -FLUORIDE ION 10.241 g in 100 g UNII:Q80VPU408O)

	Inactive Ingredients	
	Ingredient Name	
- 1		

CITRIC ACID ISOPROPYL ETHER (UNII: 19776V0481)

FD&C RED NO. 40 (UNII: WZB9127XOA)

MAGNESIUM ALUMINUM SILICATE (UNII: 6M3P64V0NC)	
PHOSPHORIC ACID (UNII: E4GA8884NN)	
POLYSORBATE 20 (UNII: 7T1F30V5YH)	
WATER (UNII: 059QF0KO0R)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
SACCHARIN SODIUM (UNII: SB8ZUX40TY)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
.ALPHATOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
XANTHAN GUM (UNII: TTV12P4NEE)	
XYLITOL (UNII: VCQ006KQ1E)	

Product Characteristics				
Color		Score		
Shape		Size		
Flavor	CHERRY	Imprint Code		
Contains				

ı	Packaging				
	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
l	1	NDC:53045-209- 17	490 g in 1 BOTTLE; Type 0: Not a Combination Product	01/01/2016	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
export only		01/01/2016	

Labeler - Dharma Research, Inc. (078444642)

Registrant - Dharma Research, Inc. (078444642)

Establishment				
Name	Address	ID/FEI	Business Operations	
Dharma Research, Inc.		078444642	manufacture(53045-209)	

Revised: 6/2021 Dharma Research, Inc.