

**MIPASTE PLUS VANILLA- mipaste plus vanilla paste, dentifrice
GC America Inc.**

MIPaste Plus Vanilla

Active Ingredient

Sodium fluoride 0.20% (w/w)

Purpose

Anticavity

Use

- aids in the prevention of dental cavities

Warnings

Keep out of reach of children under 6 years of age. Do not use on patients with a milk protein or hydroxybenzoates allergy. In case of allergic reaction; stop use, rinse mouth with water and seek medical advice. If you accidentally swallow more than used for brushing, get medical help or contact a Poison Control Center right away.

Warnings

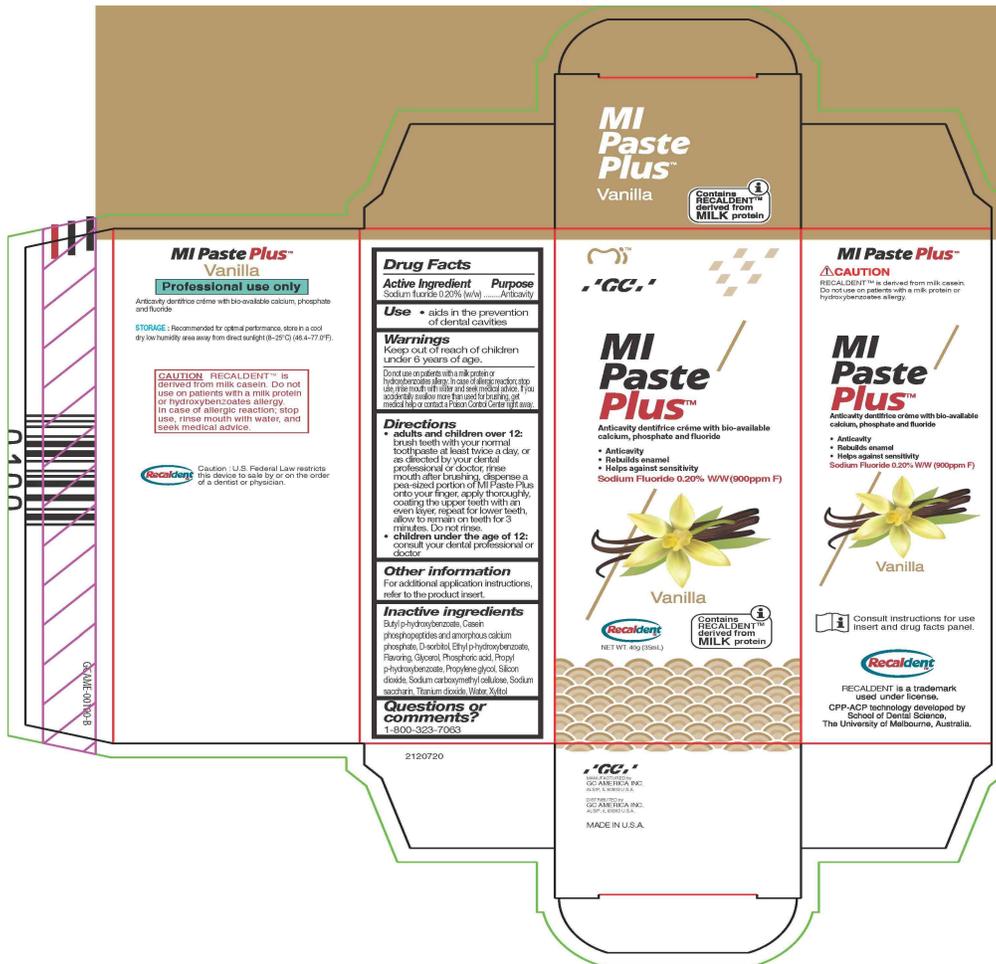
Keep out of reach of children under 6 years of age.

Directions

- adults and children over 12: brush teeth with your normal toothpaste at least twice a day, or as directed by your dental professional or doctor, rinse mouth after brushing, dispense a pea-sized portion of MI Paste Plus onto your finger, apply thoroughly, coating the upper teeth with an even layer, repeat for lower teeth, allow to remain on teeth for 3 minutes. Do not rinse.
- children under the age of 12: consult your dental professional or doctor

Inactive ingredients

Butyl p-hydroxybenzoate, Casein phosphopeptides and amorphous calcium phosphate, D-sorbitol, Ethyl p-hydroxybenzoate, Flavoring, Glycerol, Phosphoric acid, Propyl p-hydroxybenzoate, Propylene glycol, Silicon dioxide, Sodium carboxymethyl cellulose, Sodium saccharin, Titanium dioxide, Water, Xylitol



JOB INFORMATION PANEL		GCAME-00100-B.pdf									
Date and time: 07/22/2020 - 08:21:36 AM		L4876									
Design width: 6.86200 inch	Barcodes:	BWR:									
Design height: 8.35000 inch	None -	None									
Prepress Operator: PP	None -	None									
	None -	None									
	None -	None									
	Number of press inks: 4	This proof utilizes process colors to represent the Pantone colors used.									
LIMITATION OF LIABILITY All proofs should be thoroughly inspected by the recipient immediately upon receipt. It is the responsibility of the recipient to examine and proofread all proofs prior to plating. Should the proof be found to be incorrect upon the inspection described above, call Accord Carton. The liability of Accord Carton shall be limited to correcting its own product. Accord Carton will not be liable for any loss or damage including but not limited to cost of the printing plates, overtime, press delays, and or missing of insertion dates.	<table border="1"> <tr> <td></td> <td></td> <td></td> <td></td> </tr> <tr> <td>8.23%</td> <td>10.30%</td> <td>17.73%</td> <td>5.85%</td> </tr> </table>					8.23%	10.30%	17.73%	5.85%		
8.23%	10.30%	17.73%	5.85%								
	Special Inks										
	<table border="1"> <tr> <td></td> <td></td> <td></td> <td></td> </tr> </table>										
Please mark any corrections or changes on this proof. If approved sign and date as indicated below.											
SIGN FOR APPROVAL:											
Signature:	Date:										
		WWW.ACCORDCARTON.COM									

MIPASTE PLUS VANILLA

mipaste plus vanilla paste, dentifrice

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:61596-802
Route of Administration	DENTAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
SODIUM FLUORIDE (UNII: 8ZYQ1474W7) (FLUORIDE ION - UNII:Q80VPU408O)	FLUORIDE ION	0.2 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
BUTYLPARABEN (UNII: 3QPI1U3FV8)	
CALCIUM PHOSPHATE, UNSPECIFIED FORM (UNII: 97Z1W3NDX)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
CARBOXYMETHYLCELLULOSE SODIUM, UNSPECIFIED (UNII: K679OBS311)	
WATER (UNII: 059QF0KO0R)	
XYLITOL (UNII: VCQ006KQ1E)	
SACCHARIN SODIUM (UNII: SB8ZUX40TY)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
SORBITOL (UNII: 506T60A25R)	
ETHYLPARABEN (UNII: 14255EXE39)	
GLYCERIN (UNII: PDC6A3C0OX)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
PHOSPHORIC ACID (UNII: E4GA8884NN)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	

Product Characteristics

Color		Score	
Shape		Size	
Flavor	VANILLA	Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:61596-802-10	10 in 1 BOX	02/03/2022	
1	NDC:61596-802-41	1 in 1 BOX, UNIT-DOSE		
1	NDC:61596-802-40	40 g in 1 TUBE; Type 9: Other Type of Part 3 Combination Product (e.g., Drug/Device/Biological Product)		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M021	02/03/2022	

Labeler - GC America Inc. (005473608)

Registrant - GC America Inc. (005473608)

Establishment

Name	Address	ID/FEI	Business Operations
GC America Inc.		005473608	manufacture(61596-802)

Revised: 12/2025

GC America Inc.