GIANT EAGLE WINTERMINT ZERO ALCOHOL- sodium fluoride mouthwash Apollo Health and Beauty Care Inc.

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

Drug Facts

Active ingredient

Sodium Fluoride 0.02% (0.01% w/v Fluoride ion)

Purpose

Anticavity

Uses

aids in the prevention of dental cavities or decay

Warnings

Keep out of reach of children.

If more than used for rinsing is accidentally swallowed, get medical help or contact a Poison Control Center immediately.

Directions

- Adults and children 6 years of age and older: use twice a day after brushing your teeth with a toothpaste. Vigorously swish 10 mL of rinse between your teeth for 1 minute then spit out. Do not swallow the rinse. Do not eat or drink for 30 minutes after rinsing.
- Instruct children 12 years of age in good rinsing habits (to minimize swallowing).
- Supervise children as necessary until capable of using without supervision.
- Children under 6 years of age: Consult a dentist or doctor.

Other information

- Cold weather may cloud this product, but will not affect its anticavity properties.
- Store at room temperature.

Inactive ingredients

Water (Aqua), Sorbitol, Propylene Glycol, Poloxamer 407, Sodium Lauryl Sulfate, Flavor, Phosphoric Acid, Eucalyptol, Methyl Salicylate, Thymol, Menthol, Disodium Phosphate, Sucralose, Red 40 (CI 16035), Blue 1 (CI 42090).

Label Copy



Drug Facts

Active Ingredient

Sodium Fluoride 0.02% (0.01% w/v Fluoride ion)

PurposeAnticavity

Uses aids in the prevention of dental cavities or decay

Warnings Keep out of reach of children, it more than used for rinsing is accidentally swallbured, get medical help or contact a Poison Control Center immediately.

Directions • Adults and children 6 years of age and older: use twice a day after brushing your teeth with a toothpaste, Vigorousty swish 10 mL of rinse between you teeth for 1 minute then spit out, Do not swallow the rinse, Do not eat or drink for provided the safter rinsing, • Instruct children 12 years of age in good rinsing habits (to minimize swallowing), • Supervise children as necessary until capable of using without supervision. • Onlidren under 6 years of age: Consult a demister or doctor.

Other Information Cold weather may cloud this product, but will not affect its anticavity properties, Store at room temperature,

Inactive Ingredients Water (AQUA), SORBITOL, PROPYLENI GLYCOL, POLOXAWER 407, SCOIUM LAURYL SULFATE, FLAVOR, PHOSPHORIC ACID, EUCAL YPTOL, METHYL SALLCYLATE, THYMOL, MENTHOL, DISCOIUM PHOSPHATE, SUCRALOSE, RED 40 ICI 16085I, BLUE 1 ICI 420801.

DISTRIBUTED BY GIANT EAGLE, INC., PITTSBURGH, PA 15238-2809 © 2019 GIANT EAGLE, INC., QUESTIONS? COMMENTS? CALL 1-800-553-2324 GiantEagle.com

MADE IN CANADA

100% satisfaction or double your money back.

06-24340

TO OPEN: squeeze cap and turn. To close, turn cap until it clicks. Do not use if printed band around cap is broken or missing.

COOLBOOST

Fight plaque and kill odor causing germs while experiencing the cool freshness of CoolBoost™ technology! CoolBoost™ technology Provides a consistent and long lasting cooling effect that continues to work long after use — for up to 3 hours, without affecting the taste of food or drink. The best part? Your cool fresh breath will become cleaner as you reduce plaque and kill germs with every use!

CoolBoost™ is a Registered Trademark of Apollo Health & Beauty Care, Inc.

This product is not manufactured or distributed by Johnson & Johnson Healthcare products, distributor of Listerine. No animal testing



GIANT EAGLE WINTERMINT ZERO ALCOHOL

sodium fluoride mouthwash

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:63148-022

Route of Administration ORAL

Active Ingredient/Active Moiety

Ingredient Name

Basis of Strength

SODIUM FLUORIDE (UNII: 8 ZYQ 1474W7) (FLUORIDE ION - UNII:Q80 VPU408O)

FLUORIDE ION

0.2 mg in 1 mL

Inactive Ingredients Ingredient Name Strength WATER (UNII: 059QF0KO0R) SORBITOL (UNII: 506T60A25R) PROPYLENE GLYCOL (UNII: 6DC9Q167V3) POLOXAMER 407 (UNII: TUF2IVW3M2) **SODIUM LAURYL SULFATE** (UNII: 368GB5141J) PHO SPHO RIC ACID (UNII: E4GA8884NN) EUCALYPTOL (UNII: RV6J6604TK) METHYL SALICYLATE (UNII: LAV5U5022Y) THYMOL (UNII: 3J50 XA376E) MENTHOL (UNII: L7T10EIP3A) SODIUM PHO SPHATE DIBASIC DIHYDRATE (UNII: 9425516 E2T) SUCRALOSE (UNII: 96K6UQ3ZD4) FD&C RED NO. 40 (UNII: WZB9127XOA) FD&C BLUE NO. 1 (UNII: H3R47K3TBD)

Packaging								
	#	Item Code	Package Description	Marketing Start Date	Marketing End Date			
	1	NDC:63148-022- 33	1000 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	10/21/2019				

Marketing Information								
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date					
OTC monograph final	part355	10/21/2019						

Labeler - Apollo Health and Beauty Care Inc. (201901209)

Registrant - Apollo Health and Beauty Care Inc. (201901209)

Establishment								
Name	Address	ID/FEI	Business Operations					
Apollo Health and Beauty Care Inc.		201901209	manufacture(63148-022)					

Revised: 10/2019 Apollo Health and Beauty Care Inc.