ALEGRIA NEO- silicon dioxide, allantoin paste, dentifrice
Seowangmo Corporation

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

Active Ingredient: Dental Type Silica 14.0%, Aluminium Chlorohydroxy Allantoinate 0.04%

INACTIVE INGREDIENT

Inactive Ingredients: D-Sorbitol Solution, Aqua, Polyethylene Glycol 1500, Mica, Concentrated Glycerin, Sodium Cocoyl Glutamate, Carboxymethylcellulose Sodium, Peppermint Oil, Anetol, Sophora Extract, Grapefruit Seed Extract, Xylitol, Glucosyl Stevia, L-Menthol, Hydroxyapatite, Eucalyptus Extract, Sage Extract, Chamomile Extract, Aloe Extract, Glycyrrhiza Extract Powder, Argentum

PURPOSE

Purpose: Gum Care

WARNINGS

Warnings: If more than used for brushing is accidentally swallowed, get medical help or contact a Poison Control Center right away.

KEEP OUT OF REACH OF CHILDREN

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

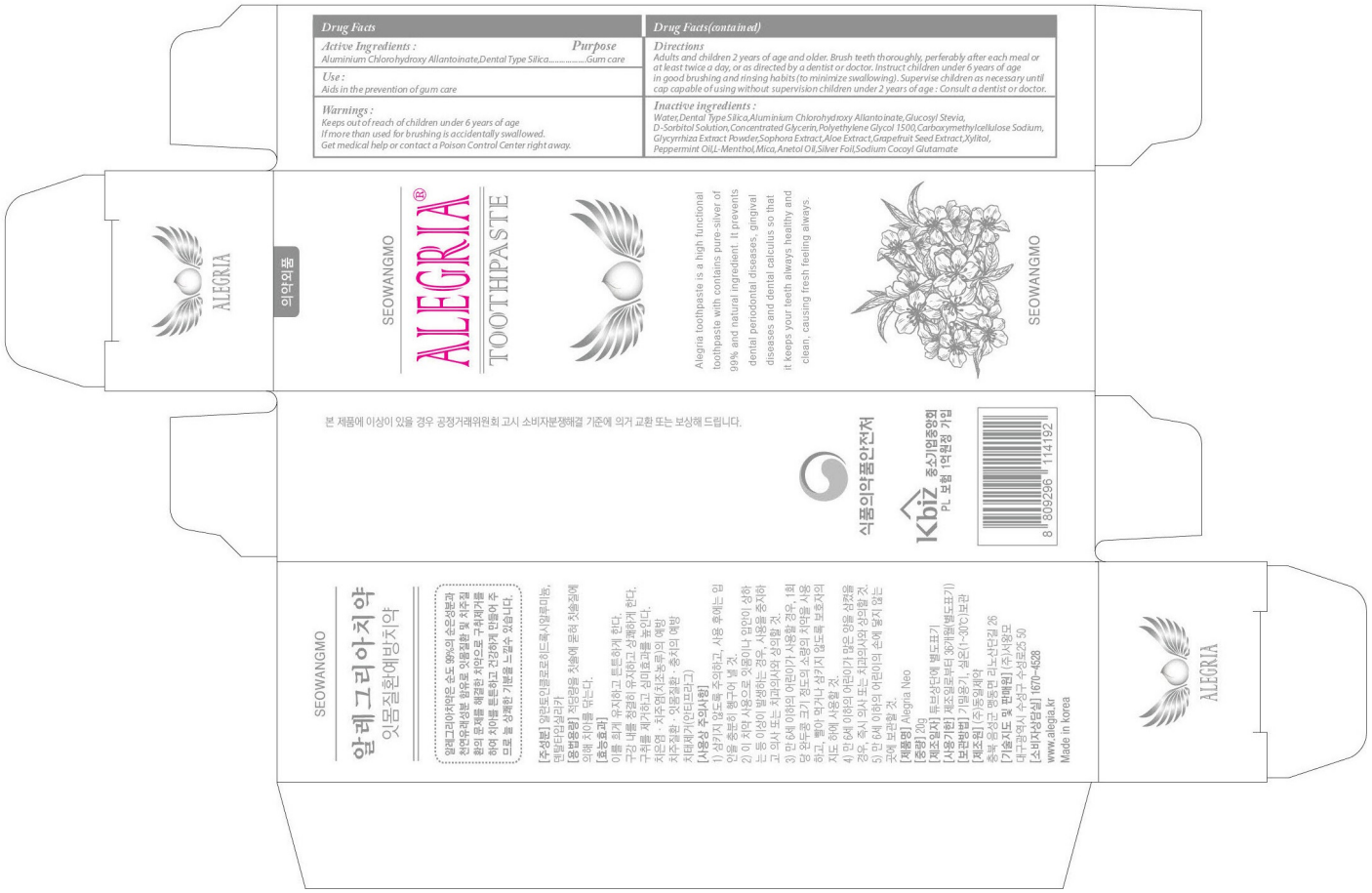
Use

Use: Aids in the prevention of gum care

Directions

Directions: 1) Adults and children 2 years of age and older: Brush teeth throughly preferably after each meal or at least twice a day or as directed by a dentist or doctor. 2) Children under 6 years: To minimize swallowing, use a pea-sized amount and supervise brushing until good habits are established. 3) Children under 2 years: Consult a dentist or doctor.

PACKAGE LABEL.PRINCIPAL DISPLAY PANEL



ALEGRIA NEO

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Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:70406-020
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
SILICON DIOXIDE (UNII: ETJ7Z6XBU4) (SILICON DIOXIDE - UNII:ETJ7Z6XBU4)	SILICON DIOXIDE	2.80 g in 20 g
Allantoin (UNII: 344S277G0Z) (ALLANTOIN - UNII:344S277G0Z)	Allantoin	0.008 g in 20 g

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
Xylitol (UNII: VCQ006KQ1E)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:70406-020-02	1 in 1 CARTON	10/01/2017	

1	NDC:70406-020-01	20 g in 1 TUBE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		10/01/2017	

Labeler - Seowangmo Corporation (689605288)

Registrant - Seowangmo Corporation (689605288)

Establishment

Name	Address	ID/FEI	Business Operations
Seowangmo Corporation		689605288	manufacture(70406-020)

Revised: 11/2017 Seowangmo Corporation