L'ECLAT BLANC TOOTHPASTE- silicon dioxide, xylitol, tocopherol acetate paste, dentifrice Toaon Co., Ltd.

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 ingredients: Silicon dioxide 12.0%, Xylitol 0.5%, Tocopherol Acetate 0.1%

INACTIVE INGREDIENT

Inactive ingredients:

Sorbitol, Purified Water, Glycerin, Sodium Cocoyl Glutamate, Sodium Cellulose Glycolate, Aroma (Fruity Floral), Menthol, Sodium Benzoate, Stevioside, Aloe Vera Leaf Extract, Sodium Hydrogen Carbonate, Eucalyptus Extract, Green Tea Extract, Sage Extract, Chamomile Extract, Rosemary Extract, Silver, Certified Organic Ingredients

PURPOSE

Purpose: Anti-plaque, Anti-cavity, Anti-gingivitis

WARNINGS

Warnings:

Keep out of the reach of children under 2 years of age. If you accidentally swallow more than used for brushing, seek professional assistance immediately.

KEEP OUT OF REACH OF CHILDREN

Keep out of the reach of children under 2 years of age.

Uses

Uses:

Keep your teeth white and strong.

Keep your mouth clean.

Refresh the inside of the mouth.

To prevent tooth decay and remove bad breath.

Increase esthetic effect.

Removal of plaque (anti-plaque).

Prevention of gingivitis.

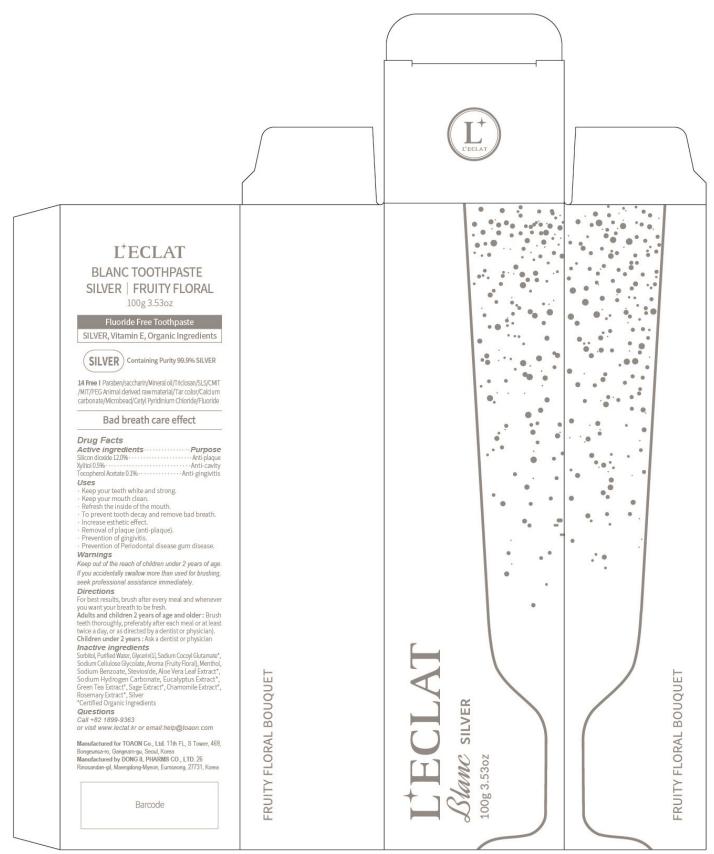
Prevention of Periodontal disease gum disease.

Directions

Directions:

For best results, brush after every meal and whenever you want your breath to be fresh. Adults and children 2 years of age and older: Brush teeth thoroughly, preferably after each meal or at least twice a day, or as directed by a dentist or physician).

PACKAGE LABEL.PRINCIPAL DISPLAY PANEL



silicon dioxide, xylitol, tocopherol acetate paste, dentifrice

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:73175-030
Route of Administration	ORAL		

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
Silicon dioxide (UNII: ETJ7Z6XBU4) (SILICON DIOXIDE - UNII:ETJ7Z6XBU4)	Silicon dioxide	12.0 g in 100 g	
Xylitol (UNII: VCQ006KQ1E) (XYLITOL - UNII:VCQ006KQ1E)	Xylitol	0.5 g in 100 g	
.ALPHATO COPHERO L ACETATE (UNII: 9E8 X80 D2L0) (.ALPHATOCOPHERO L - UNII: H4N855PNZ1)	.ALPHATOCOPHEROL ACETATE	0.1 g in 100 g	

Inactive Ingredients	
Ingredient Name	Strength
Sorbitol (UNII: 506T60A25R)	
Water (UNII: 059QF0KO0R)	

Packaging			
# Item Code	Package Description	Marketing Start Date	Marketing End Date
1 NDC:73175-030-02	1 in 1 CARTON	06/01/2019	
1 NDC:73175-030-01	100 g in 1 TUBE; Type 0: Not a Combination Product		

Marketing Info	rmation		
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		06/01/2019	

Labeler - Toaon Co., Ltd. (694977241)

Registrant - Toaon Co., Ltd. (694977241)

Establishment				
Name	Address	ID/FEI	Business Operations	
DONG IL PHARMS CO., LTD.		557810721	manufacture(73175-030)	

Revised: 7/2019 Toaon Co., Ltd.