FLAVON WHITE MINT TOOTH- calcium carbonate, allantoin powder, dentifrice
BS and 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

Calcium Carbonate 45.0%
Allantoin 1.0%

PURPOSE

Anti-plaque
Anti-gingivitis, Anti-Periodontitis

Uses

Aids in the removal of plaque and prevention gingivitis and periodontitis.

WARNINGS

Keep out of the reach of children under 6 years of age.
If you accidentally swallow more than used for brushing, seek professional help or contact a poison control center immediately.

KEEP OUT OF REACH OF CHILDREN

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

Directions

Adults and children 2 yrs. older: Brush teeth thoroughly after meals or at least twice a day, or use as directed by a dentist or physician. Do not swallow.
Children under 6 yrs.: To minimize swallowing, use a pea-sized amount and supervise brushing until good habits are established.
Children under 2 yrs.: Ask a dentist or physician.

Other Information

Store at room temperature

QUESTIONS

Visit www.beauti-science.com/whitelabs/en

INACTIVE INGREDIENTS

Sorbitol, Sodium Bicarbonate, Sodium Lauryl Sulfate, Xylitol, Xanthan Gum, Mint Flavor, Menthol, Mentha Piperita Oil, Stevioside, Ascorbic Acid

PACKAGE LABEL.PRINCIPAL DISPLAY PANEL

FLAVON WHITE MINT TOOTH

calcium carbonate, allantoin powder, dentifrice

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:73660-003
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
Calcium Carbonate (UNII: H0G9379FGK) (CALCIUM CATION - UNII:2M83C4R6ZB)	CALCIUM CATION	45.0 g in 100 g
Allantoin (UNII: 344S277G0Z) (ALLANTOIN - UNII:344S277G0Z)	Allantoin	1.0 g in 100 g

Inactive Ingredients	
Ingredient Name	**Strength**
Sorbitol (UNII: 506T60A25R)	
Sodium Bicarbonate (UNII: 8MDF5V39QO)	

Sodium Lauryl Sulfate (UNII: 368GB5141J)	
Xylitol (UNII: VCQ006KQ1E)	
Xanthan Gum (UNII: TTV12P4NEE)	
Menthol (UNII: L7T10EIP3A)	
PEPPERMINT OIL (UNII: AV092KU4JH)	
Stevioside (UNII: 0YON5MXJ9P)	
ASCORBIC ACID (UNII: PQ6CK8PD0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:73660-003-02	1 in 1 CARTON	07/01/2020	
1	NDC:73660-003-01	50 g in 1 BOTTLE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		07/01/2020	

Labeler - BS and Co. Ltd (689513606)

Registrant - BS and Co. Ltd (689513606)

Establishment

Name	Address	ID/FEI	Business Operations
BIOSTECH CO., LTD.		687294330	manufacture(73660-003)

Revised: 7/2020 BS and Co. Ltd