SALTRAIN LTOOTHPASTE- sodium chloride, precipitated calcium carbonate paste, dentifrice
K.Boeun Pharmaceutical 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.

Sodium Chloride, Precipitated Calcium Carbonate

D-Sorbitol Solution
Concentrated Glycerin
Polyethylene Glycol 1500
Xanthan gum
Titanium Oxide
Silicon Dioxide
Sodium Cocoyl Glutamate
Enzymatically Modified Stevia
Xylitol
Ascorbic Acid
Tocopherol Acetate
Propolis Extract
Green Tea Extract

Aloe Extract
l-Menthol
Mentha Oil
Lemon Oil
Water

aids in the prevention of dental cavities

plaque removal (anti-plaque)

prevention of periodontal disease, prevention of gum disease

Keep out of reach of children

Apply an appropriate amount to your toothbrush and brush your teeth by brushing.

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 right away.

Other Information

■ Store in an airtight container at room temperature

■ Date of use : 36 months from the date of manufacture

For dental use only

SALTRAIN LTOOTHPASTE

sodium chloride, precipitated calcium carbonate paste, dentifrice

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:74724-0031
Route of Administration	DENTAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
SODIUM CHLORIDE (UNII: 451W47IQ8X) (CHLORIDE ION - UNII:Q32ZN48698)	SODIUM CHLORIDE	3 g in 100 g
CALCIUM CARBONATE (UNII: H0G9379FGK) (CALCIUM CATION - UNII:2M83C4R6ZB)	CALCIUM CARBONATE	33 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
XYLITOL (UNII: VCQ006KQ1E)	
WATER (UNII: 059QF0KO0R)	

Packaging

		Marketing Start	Marketing End

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:74724-0031-1	180 g in 1 TUBE; Type 0: Not a Combination Product	01/01/2023	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		01/01/2023	

Labeler - K.Boeun Pharmaceutical Co.,Ltd. (695674074)

Registrant - K.Boeun Pharmaceutical Co.,Ltd. (695674074)

Establishment

Name	Address	ID/FEI	Business Operations
K.Boeun Pharmaceutical Co.,Ltd.		695674074	manufacture(74724-0031)

Revised: 12/2022 K.Boeun Pharmaceutical Co.,Ltd.