MIMIBO PREMIUM TOOTH- silicon dioxide, tocopherol acetate, tetrasodium pyrophosphate paste, dentifrice
CONE MEDICAL 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.

Silicon Dioxide

Tocopherol Acetate

Tetrasodium Pyrophosphate

D-Sorbitol Solution

Concentrated Glycerin

Carboxymethylcellulose Sodium

Hydroxyapatite

Medicinal Carbon

Chitosan

Mica

Zeolite

Bamboo Salt

Xylitol

Steviol Glycoside

Papain

Grapefruit Seed Extract

L-Menthol

Mentha Oil

Propolis Extract

Chamomile Extract
Rosemary Extract

Sage Extract

Aloe Extract

Glycyrrhiza Extract

Lavender Oil

Sodium Cocoyl Glutamate

Lauroyl Amidopropyl Dimethyl Glycine Solution

Deionized Water

For dental care

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

100g 단상자 48 X 37 X165

100g 튜브

m100 은박

MIMIBO PREMIUM TOOTH

silicon dioxide, tocopherol acetate, tetrasodium pyrophosphate paste, dentifrice

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:73527-0003
Route of Administration	DENTAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
SILICON DIOXIDE (UNII: ETJ7Z6XBU4) (SILICON DIOXIDE - UNII:ETJ7Z6XBU4)	SILICON DIOXIDE	14 g in 100 g
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0) (.ALPHA.-TOCOPHEROL - UNII:H4N855PNZ1)	.ALPHA.-TOCOPHEROL ACETATE	0.1 g in 100 g
SODIUM PYROPHOSPHATE (UNII: O352864B8Z) (PYROPHOSPHORIC ACID - UNII:4E862E7GRQ)	SODIUM PYROPHOSPHATE	0.5 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
XYLITOL (UNII: VCQ006KQ1E)	
LEVOMENTHOL (UNII: BZ1R15MTK7)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:73527-0003-1	100 g in 1 TUBE; Type 0: Not a Combination Product	04/01/2021	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		04/01/2021	

Labeler - CONE MEDICAL Co., Ltd. (695811691)

Registrant - CONE MEDICAL Co., Ltd. (695811691)

Establishment

Name	Address	ID/FEI	Business Operations
HealingStory.,LTD.		688403124	manufacture(73527-0003)

Establishment

Name	Address	ID/FEI	Business Operations
CONE MEDICAL Co., Ltd.		695811691	label(73527-0003)

Revised: 11/2023

CONE MEDICAL Co., Ltd.