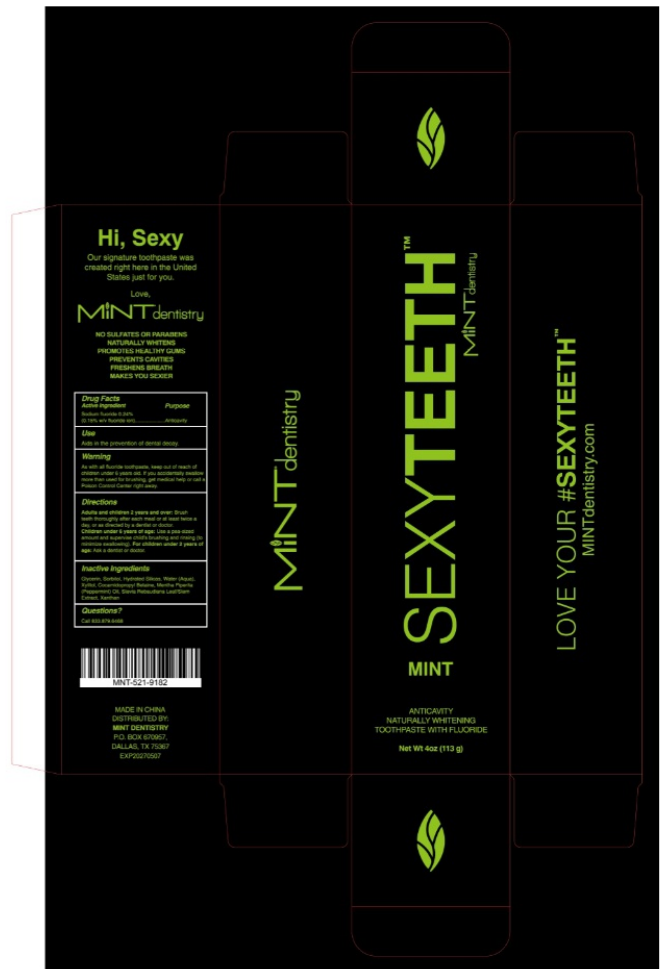


SEXYTEETHMINTTOOTHPASTE- toothpaste paste, dentifrice SHANTOU S.E.Z BAOJIE INDUSTRY CO., LTD

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.

Warning

As with all fluoride toothpaste, keep out of reach of children under 6 years old. If you accidentally swallow more than used for brushing, get medical help or call a Poison Control Center right away.



Inactive Ingredients

Glycerin, Sorbitol, Hydrated Silicas, Water (Aqua), Xylitol, Cocamidopropyl Betaine, Mentha Piperita (Peppermint) Oil, Stevia Rebaudiana Leaf/Stem Extract, Xanthan

Use

Aids in the prevention of dental decay.

Warning

As with all fluoride toothpaste, keep out of reach of children under 6 years old. If you accidentally swallow more than used for brushing, get medical help or call a Poison Control Center right away.

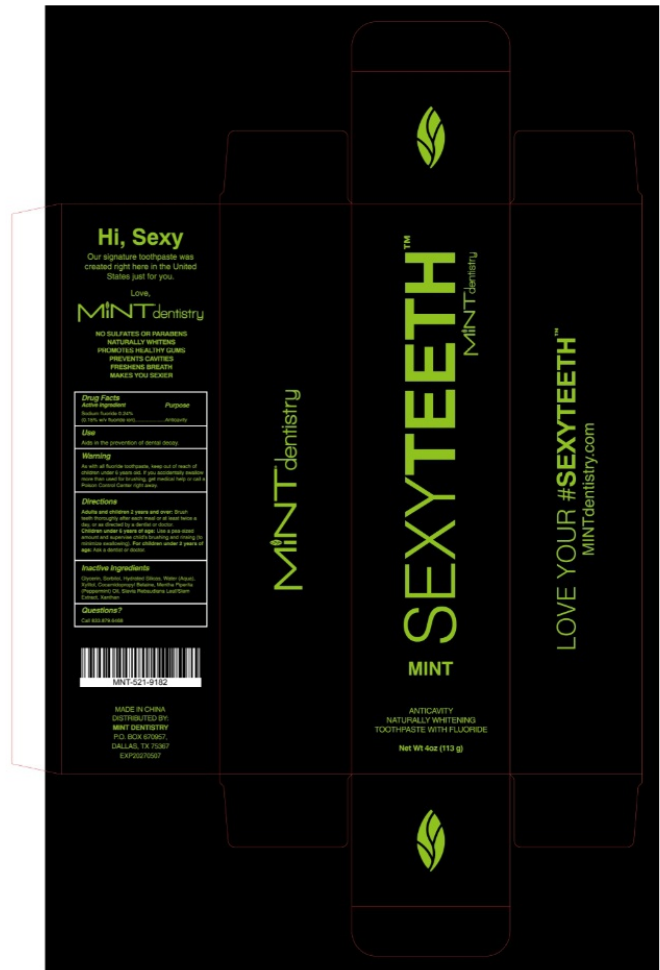
Drug Facts

Active Ingredient

Sodium fluoride 0.24%
(0.15% w/v fluoride ion).....Anticavity

Purpose

Use





SEXYTEETHMINTTOOTHPASTE

toothpaste paste, dentifrice

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:74913-703
Route of Administration	DENTAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
SODIUM FLUORIDE (UNII: 8ZYQ1474W7) (FLUORIDE ION - UNII:Q80VPU4080)	SODIUM FLUORIDE	0.24 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
SORBITOL (UNII: 506T60A25R)	60 g in 100 g
HYDRATED SILICA (UNII: Y607T4G8P9)	12 g in 100 g
SPEARMINT OIL (UNII: C3M81465G5)	0.8 g in 100 g
WATER (UNII: 059QF0KO0R)	16.16 g in 100 g
COCAMIDOPROPYL BETAINE (UNII: 5OCF3011KX)	1.5 g in 100 g
GLYCERIN (UNII: PDC6A3C0OX)	5 g in 100 g

XYLITOL (UNII: VCQ006KQ1E)	3 g in 100 g
STEVIA REBAUDIANA LEAF (UNII: 6TC6NN0876)	0.5 g in 100 g
XANTHAN GUM (UNII: TTV12P4NEE)	0.8 g in 100 g

Product Characteristics

Color		Score	
Shape		Size	
Flavor	MINT	Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:74913-703-01	113 g in 1 TUBE; Type 0: Not a Combination Product	06/15/2024	



Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		06/15/2024	

Labeler - SHANTOU S.E.Z BAOJIE INDUSTRY CO., LTD (546345856)

Registrant - SHANTOU S.E.Z BAOJIE INDUSTRY CO., LTD (546345856)

Establishment

Name	Address	ID/FEI	Business Operations
SHANTOU S.E.Z BAOJIE INDUSTRY CO., LTD		546345856	manufacture(74913-703)

Revised: 6/2024

SHANTOU S.E.Z BAOJIE INDUSTRY CO., LTD