

THERABREATH WHITENING- sodium fluoride paste, dentifrice
Church & Dwight Co., Inc.

Therabreath™ Whitening Dentist Formulated Toothpaste

Drug Facts

Active Ingredient

Sodium fluoride 0.24% (0.17% w/v fluoride ion)

Purpose

Anticavity

Use

aids in the prevention of dental decay

Warning

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

If more than used for brushing is accidentally swallowed, get medical help or contact a Poison Control Center right away.

Directions

• do not swallow • supervise children as necessary until capable of using without supervision

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 doctor
children under 6 years of age	instruct in good brushing and rinsing habits (to minimize swallowing)
children under 2 years of age	consult a dentist or doctor

Inactive Ingredients

water, sodium bicarbonate, sorbitol, hydrated silica, glycerin, tetrasodium pyrophosphate, sodium tripolyphosphate, flavor, sodium lauryl sulfate, sodium lauroyl sarcosinate, poloxamer 407, xanthan gum, titanium dioxide, sodium saccharin, sucralose

Questions or Comments?

TheraBreath is manufactured by

Church & Dwight Co., Inc. Ewing, NJ 08628 U.S.A.

Phone: 1-800-973-7374 WEB: WWW.THERABREATH.COM

Principal Display Panel

LONG LASTING FOR EPIC FRESHNESS

TheraBreath[™]

WHITENING

DENTIST FORMULATED TOOTHPASTE

Anticavity

SIGNIFICANTLY WHITENS AFTER 1 WEEK*

TWIN PACK *with twice daily brushing

Neutralizes the source of bad breath[†]

[†]in laboratory tests

Gently whitens

Dr. Harold Katz

Dazzling Mint

2 TUBES of 4.0 oz (113 g)

TOTAL NET WT. 8.0 oz (226 g)

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DAZZLING MINT

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Tamper Evident: Do not use if inner foil is torn, cut or missing.

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TheraBreath™ is manufactured by Church & Dwight Co., Inc., Ewing, NJ 08628 U.S.A.

Made in the USA with US and internationally-sourced ingredients.

TBTU-99621-01 72032836

THERABREATH WHITENING

sodium fluoride paste, dentifrice

Product Information

Product Type		HUMAN OTC DRUG	Item Code (Source)	NDC:10237-272
Route of Administration		ORAL		
Active Ingredient/Active Moiety				
Ingredient Name			Basis of Strength	Strength
SODIUM FLUORIDE (UNII: 8ZYQ1474W7) (FLUORIDE ION - UNII:Q80VPU408O)			SODIUM FLUORIDE	0.243 g in 100 g
Inactive Ingredients				
Ingredient Name				Strength
WATER (UNII: 059QF0KO0R)				
SODIUM BICARBONATE (UNII: 8MDF5V39QO)				
SODIUM TRIPOLYPHOSPHATE (UNII: 5HK03SA80J)				
SODIUM LAUROYL SARCOSINATE (UNII: 632GS99618)				
SUCRALOSE (UNII: 96K6UQ3ZD4)				
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)				
SORBITOL (UNII: 506T60A25R)				
HYDRATED SILICA (UNII: Y6O7T4G8P9)				
SODIUM LAURYL SULFATE (UNII: 368GB5141J)				
POLOXAMER 407 (UNII: TUF2IVW3M2)				
GLYCERIN (UNII: PDC6A3C0OX)				
SODIUM SACCHARIN (UNII: SB8ZUX40TY)				
TETRASODIUM PYROPHOSPHATE (UNII: O352864B8Z)				
XANTHAN GUM (UNII: TTV12P4NEE)				
Product Characteristics				
Color		white	Score	
Shape			Size	
Flavor		MINT	Imprint Code	
Contains				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:10237-272-24	2 in 1 CARTON	08/01/2025	
1		113 g in 1 TUBE; Type 0: Not a Combination Product		
2	NDC:10237-272-04	1 in 1 CARTON	12/15/2025	
2		113 g in 1 TUBE; Type 0: Not a Combination Product		
Marketing Information				
Marketing		Application Number or Monograph	Marketing Start	Marketing End

Category	Citation	Date	Date
OTC Monograph Drug	M021	08/01/2025	

Labeler - Church & Dwight Co., Inc. (001211952)

Establishment

Name	Address	ID/FEI	Business Operations
Church & Dwight Co., Inc.		043690812	manufacture(10237-272)

Revised: 8/2025

Church & Dwight Co., Inc.