

SPRY DENTIFRICE- toothpaste paste, dentifrice
Xlear Inc.

Spry Fluoride Toothpaste

Warning: keep out of reach of children under 6 years of age.

Warnings: If more than used for brushing is accidentally swallowed, get medical help or contact poison control center right away.

Brush teeth thoroughly, preferably after each meal or at least twice a day, or as directed by a dentist or doctor.

Directions: Adults and children two years and older:

Brush teeth thoroughly, preferably after each meal or at least twice a day, or as directed by a dentist or doctor.

Instruct children under 6 years of age in good brushing and rinsing habits (to minimize swallowing).

Supervise children as necessary until capable of using without supervision.

Children under 2 years of age: consult a dentist or doctor.

Warnings: 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 poison control center right away

Sodium fluoride 0.243% (w/w)

Anticavity toothpaste

Aids in the prevention of dental cavities

Anticavity Toothpaste

Directions: adults and children two years and older:

brush teeth thoroughly, preferably after each meal or at least twice a day, or as directed by a dentist or doctor.

Water, silica, vegetable glycerin, xylitol, erythritol, lauryl glucoside, L-arginine, sodium methyl cocoyl taurate, natural spearmint flavor, cranberry extract, xanthan gum, titanium dioxide, stevia, cellulose gum, sodium trimetaphosphate, sodium benzoate, zinc citrate, aloe vera



SPRY DENTIFRICE

toothpaste paste, dentifrice

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:27017-142
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
SODIUM FLUORIDE (UNII: 8ZYQ1474W7) (FLUORIDE ION - UNII:Q80VPU4080)	FLUORIDE ION	2.43 mg in 1 g

Inactive Ingredients

Ingredient Name	Strength
ZINC CITRATE (UNII: K72I3DEX9B)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
CELLULOSE GUM (UNII: K679OBS311)	
GLYCERIN (UNII: PDC6A3C0OX)	
STEVIA LEAF (UNII: 6TC6NN0876)	
ERYTHRITOL (UNII: RA96B954X6)	
LAURYL GLUCOSIDE (UNII: 76LN7P7UCU)	
SODIUM METHYL COCOYL TAURATE (UNII: JVL98CG53G)	
WATER (UNII: 059QF0KO0R)	

SODIUM BENZOATE (UNII: OJ245FE5EU)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
N2-(4-FLUOROBENZOYL)-L-ARGININE (UNII: NN7Q1D14KA)	
SPEARMINT OIL (UNII: C3M81465G5)	
XANTHAN GUM (UNII: TTV12P4NEE)	
SODIUM TRIMETAPHOSPHATE (UNII: 3IH6169RL0)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
XYLITOL (UNII: VCQ006KQ1E)	
CRANBERRY (UNII: 0MVO31Q3QS)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:27017-142-01	1 g in 1 TUBE; Type 0: Not a Combination Product	06/11/2017	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M021	06/11/2017	

Labeler - Xlear Inc. (839884058)

Registrant - Xlear Inc. (839884058)

Establishment

Name	Address	ID/FEI	Business Operations
Xlear Inc.		839884058	manufacture(27017-142)

Revised: 11/2025

Xlear Inc.