MORNING FRESH MINT TOOTHPASTE, WITH FLUORIDE- morning fresh mint toothpaste, with fluoride paste
Dynarex Corporation

4871-4873 Morning Fresh Mint Toothpaste, with Fluoride

Active Ingredient

Sodium Monofluorophosphate 0.76% (0.13% w/v fluoride ion)

Purpose

Anticavity

Use(s)

Helps protect against cavities

Warning

Do Not Swallow

Keep out of reach of children under 6 years of age

If you accidentally swallow more than used for brushing, get medical help or contact a Poison Control Center (1-800-222-1222) immediately.

Directions

• **Adults and children 2 years and older:** Brush teeth thoroughly after meals or at least twice a day or use as directed by a dentist or physician.

• **Children under 6 years:** to minimize swallowing, use a pea size amount and supervise brushing and rinsing until good habits are established.

• **Children under 2 years:** Use as directed by a dentist or physician.

Inactive Ingredients

Calcium Carbonate, Carboxymethylcellulose Sodium, Methylparaben, Peppermint Oil, Polyethylene Glycol 400, Propylparaben, Purified Water, Saccharin Sodium, Silicon Dioxide, Sodium Benzoate, Sodium Lauryl Sulfate, Sodium Pyrophosphate, Sodium Silicate, Sorbitol, Titanium Dioxide

Other Information

Store between 15º - 35ºC (59º - 95ºF)

Questions?

1-888-396-2739 Monday - Friday 9AM - 5PM EST

Label

4871 Label

Label

4872 Label

Label

4873 Label

MORNING FRESH MINT TOOTHPASTE, WITH FLUORIDE

morning fresh mint toothpaste, with fluoride paste

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:67777-190
Route of Administration	DENTAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
SODIUM MONOFLUOROPHOSPHATE (UNII: C810JCZ56Q) (FLUORIDE ION - UNII:Q80VPU408O)	FLUORIDE ION	1.3 mg in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**
SODIUM SILICATE (UNII: IJF18F77L3)	
WATER (UNII: 059QF0KO0R)	
CALCIUM CARBONATE (UNII: H0G9379FGK)	
SODIUM LAURYL SULFATE (UNII: 368GB5141J)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	

SODIUM BENZOATE (UNII: OJ245FE5EU)	
SACCHARIN SODIUM (UNII: SB8ZUX40TY)	
SORBITOL (UNII: 506T60A25R)	
CARBOXYMETHYLCELLULOSE SODIUM, UNSPECIFIED (UNII: K679OBS311)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
PEPPERMINT OIL (UNII: AV092KU4JH)	
POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ)	
SODIUM PYROPHOSPHATE (UNII: O352864B8Z)	

Product Characteristics

Color		**Score**	
Shape		**Size**	
Flavor	MINT	**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:67777-190-03	5 in 1 CASE	01/15/2025	
1	NDC:67777-190-02	144 in 1 BOX		
1	NDC:67777-190-01	25.14 mL in 1 TUBE; Type 0: Not a Combination Product		
2	NDC:67777-190-06	144 in 1 CASE	01/15/2025	
2	NDC:67777-190-05	1 in 1 BOX		
2	NDC:67777-190-04	44.36 mL in 1 TUBE; Type 0: Not a Combination Product		
3	NDC:67777-190-09	144 in 1 CASE	01/15/2025	
3	NDC:67777-190-08	1 in 1 BOX		
3	NDC:67777-190-07	81.33 mL in 1 TUBE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M021	01/15/2025	

Labeler - Dynarex Corporation (008124539)

Revised: 5/2026 Dynarex Corporation