

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

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**4871 Morning Fresh Mint Toothpaste, with Fluoride NDC # 67777-190-02**

***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 15° - 35°C (59° - 95°F)

## Questions?

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

## Label



4871 Label

## Label



4872 Label

## Label



4873 Label

## MORNING FRESH MINT TOOTH PASTE, WITH FLUORIDE

morning fresh mint toothpaste, with fluoride paste

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:67777-190
<b>Route of Administration</b>	DENTAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>SODIUM MONOFLUOROPHOSPHATE</b> (UNII: C810JCZ56Q) (FLUORIDE ION - UNII:Q80VPU4080)	FLUORIDE ION	1.3 mg in 1 mL

### Inactive Ingredients

Ingredient Name	Strength
<b>SODIUM SILICATE</b> (UNII: IJF18F77L3)	
<b>WATER</b> (UNII: 059QF0KO0R)	
<b>CALCIUM CARBONATE</b> (UNII: H0G9379FGK)	
<b>SODIUM LAURYL SULFATE</b> (UNII: 368GB5141J)	
<b>METHYLPARABEN</b> (UNII: A2I8C7HI9T)	
<b>PROPYLPARABEN</b> (UNII: Z8IX2SC1OH)	

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

### Product Characteristics

<b>Color</b>		<b>Score</b>	
<b>Shape</b>		<b>Size</b>	
<b>Flavor</b>	MINT	<b>Imprint Code</b>	
<b>Contains</b>			

### 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)