

MORNING FRESH- sodium fluoride paste
Dynarex Corporation

4872 Morning Fresh Toothpaste NDC 67777-175-01
4873 Morning Fresh Toothpaste NDC 67777-175-02

Active Ingredient

Sodium Monofluorophosphate 0.76 %

Purpose

Anticavity

Use(s)

- Helps protect against cavities

Warnings

- **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 sized amount and supervise brushing and rinsing until good habits are established.
- **Children under 2 years:** Use as Directed by a dentist or physician.

Inactive Ingredient

Calcium Carbonate, Carboxymethylcellulose Sodium, Flavor, Methylparaben, Polyethylene Glycol 400, Propylparaben, Saccharin Sodium, Silicon Sioxide, Sodium Benzoate, Sodium Lauryl Sulphate, Sodium Pyrophosphate, Sodium Silicate, Sorbitol, Titanium Dioxide, Water

Other Information

Store between 15° - 35°C (59° - 95°F)

Questions?

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

Label



4872 Morning Fresh Toothpaste, with Fluoride

Label



4873 Morning Fresh Toothpaste, With Fluoride

MORNING FRESH			
sodium fluoride paste			
Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:6777-175
Route of Administration	ORAL		
Active Ingredient/Active Moiety			
Ingredient Name		Basis of Strength	Strength
SODIUM MONOFLUOROPHOSPHATE (UNII: C810JCZ56Q) (FLUORIDE ION - UNII:Q80VPU4080)		FLUORIDE ION	76 mg in 1 g
Inactive Ingredients			
Ingredient Name			Strength
SODIUM BENZOATE (UNII: OJ245FE5EU)			
SORBITOL (UNII: 506T60A25R)			
SACCHARIN SODIUM (UNII: SB8ZUX40TY)			
WATER (UNII: 059QF0KO0R)			
CARBOXYMETHYLCELLULOSE SODIUM (UNII: K679OBS311)			
SODIUM LAURYL SULFATE (UNII: 368GB5141J)			

SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
SODIUM SILICATE (UNII: IJF18F77L3)	
SODIUM PYROPHOSPHATE (UNII: O352864B8Z)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
CALCIUM CARBONATE (UNII: H0G9379FGK)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:67777-175-11	144 in 1 CASE	10/14/2014	
1	NDC:67777-175-10	1 in 1 BOX		
1	NDC:67777-175-01	43 g in 1 TUBE; Type 0: Not a Combination Product		
2	NDC:67777-175-13	144 in 1 CASE	10/14/2014	
2	NDC:67777-175-12	1 in 1 BOX		
2	NDC:67777-175-02	78 g 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	10/14/2014	

Labeler - Dynarex Corporation (008124539)

Registrant - Dynarex Corporation (008124539)

Revised: 12/2024

Dynarex Corporation