IDODENT FLOURIDE MINT- fluoride paste
United Exchange Corp.

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

Active Ingredient **Purpose**

Sodium monofluorophosphate 0.76%---------- Anticavity

(Total Fluoride Content - 1000 ppm Approx.)

Uses

Regular brushing with fluoride toothpaste helps protect teeth and roots against cavities

Warnings

Warnings

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

In case of accidental overdose, seek professional assistance or contact a Poison Control Center immediately.

Directions

Adults and children 2 years and older	Brush Teeth thoroughly after meals, 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 until good habits are established.
Children under 2 years	Ask a dentist or physician.

Other Information

Store at room temperature 20-25°C (68-77°F)

Inactive ingredients

calcium carbonate, water, sorbitol, hydrated silica, sodium lauryl sulfate, flavor, cellulose gum, sodium silicate, tetrasodium, propphosphate, sodium saccharin, titanium dioxide, methylparaben, PEG-8, propylparaben

Distributed by:

United Exchange Corp.

17211 Valley View Ave.

Cerritos, CA 90703 USA

IODENT FLOURIDE MINT

fluoride paste

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:65923-181
Route of Administration	DENTAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
SODIUM MONOFLUOROPHOSPHATE (UNII: C810JCZ56Q) (FLUORIDE ION - UNII:Q80VPU408O)	FLUORIDE ION	0.76 g in 100 g

Inactive Ingredients	
Ingredient Name	**Strength**
SACCHARIN SODIUM (UNII: SB8ZUX40TY)	
SODIUM PYROPHOSPHATE (UNII: O352864B8Z)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
SODIUM LAURYL SULFATE (UNII: 368GB5141J)	
CALCIUM CARBONATE (UNII: H0G9379FGK)	
WATER (UNII: 059QF0KO0R)	

CARBOXYMETHYLCELLULOSE SODIUM, UNSPECIFIED FORM (UNII: K679OBS311)	
SORBITOL (UNII: 506T60A25R)	
HYDRATED SILICA (UNII: Y6O7T4G8P9)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
SODIUM SILICATE (UNII: IJF18F77L3)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:65923-181-18	1 in 1 CARTON	12/21/2016	
1		181 g in 1 TUBE; Type 1: Convenience Kit of Co-Package		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part355	10/23/2014	

Labeler - United Exchange Corp. (840130579)

Revised: 4/2018 United Exchange Corp.