# LUCKY SUPER SOFT SENSITIVE- sodium fluoride potassium nitrate paste, dentifrice Delta Brands Inc

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.

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#### **Drug Facts**

#### **Active Ingredient**

Potassium nitrate - 5%

Sodium fluoride - 0.24%

### **Purpose**

Antihypersensitivity

Anticavity

#### Uses

■ builds increasing protection against painful sensitivity of teeth due to cold, heat, acids, sweets or contact ■ aids in the prevention of dental cavities

#### Warnings

**When using this product** ■ if pain/sensitivity still persists after 4 weeks of use please visit your dentist

#### Stop and ask a dentist

■ if problem the persists or worsens. Sensitive teeth may indicate a serious problem that may need prompt care by a dentist.

#### Keep out of reach of children

If you accidentally swallowed more than used for brushing, seek professional help or contact a Poison Control Center immediately

#### **Directions**

adults and children 12 years and older brush teeth for at least one minute, preferably after each meal, or at least twice a day, or as directed by your dentist. Be sure to brush sensitive areas children under 12 years of age consult a dentist or doctor.

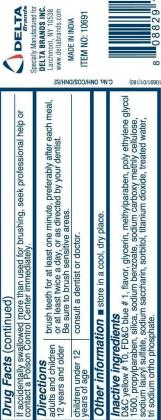
#### **Inactive ingredients**

DC Yellow10, FDC Blue 1, flavor, glycerin, methylparaben, poly ethylene glycol 1500, propylparaben, silica, sodium benzoate, sodium carboxy methyl cellulose, sodium lauryl sulfate, sodium saccharin, sorbitol, titanium dioxide, treated water, tri sodium ortho phosphate









Other information store in a cool, dry place.

Warnings
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problem that may need prompt care by a dentist.

Keep out of reach of children.

consult a dentist or doctor.

children under 12 years of age

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**Directions** adults and children 12 years and older

Drug Facts (continued)

Purpose .Antihypersensitivity .....Anticavity

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**Drug Facts** 













## LUCKY SUPER SOFT SENSITIVE

sodium fluoride potassium nitrate paste, dentifrice

<b>Product Information</b>			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:20276-550
Route of Administration	DENTAL		

Active Ingredient/Active Moiety		
Ingredient Name	Basis of Strength	Strength
POTASSIUM NITRATE (UNII: RU45X2JN0Z) (NITRATE ION - UNII:T93E9Y2844)	POTASSIUM NITRATE	50 mg in 1 g
SODIUM FLUORIDE (UNII: 8ZYQ1474W7) (FLUORIDE ION - UNII:Q80 VPU408O)	FLUORIDE ION	2.4 mg in 1 g

Inactive Ingredients	
Ingredient Name	Strength
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
GLYCERIN (UNII: PDC6A3C0OX)	
METHYLPARABEN (UNII: A2I8 C7HI9 T)	
POLYETHYLENE GLYCOL 1500 (UNII: 1212Z7S33A)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
SILICON DIO XIDE (UNII: ETJ7Z6 XBU4)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
CARBOXYMETHYLCELLULOSE SODIUM (UNII: K679 OBS 311)	
SODIUM LAURYL SULFATE (UNII: 368GB5141J)	
SACCHARIN SODIUM (UNII: SB8ZUX40TY)	
SORBITOL (UNII: 506T60A25R)	
TITANIUM DIO XIDE (UNII: 15FIX9 V2JP)	
D&C YELLOW NO. 10 (UNII: 35SW5USQ3G)	
SO DIUM PHO SPHATE, TRIBASIC, DO DECAHYDRATE (UNII: B70850 QPHR)	
WATER (UNII: 059QF0KO0R)	

Product Characteristics			
Color	green	Score	

Shape			Size		
Flavor		MINT	Imprint Code		
Contains					
Packaging					
# Item Code		Package Description	n	Marketing Start Date	Marketing End Date
1 NDC:20276-550-43	122 g in 1 TUBI	E; Type 0: Not a Combina	ation Product	02/27/2014	
Marketing Information					
Marketing Categor	y Applicat	tion Number or Mono	graph Citation	Marketing Start Date	Marketing End Date
OTC monograph not fin	al part356			02/27/2014	

## Labeler - Delta Brands Inc (102672008)

Revised: 7/2017 Delta Brands Inc