# UP AND UP SENSITIVE TOOTHPASTE ENAMEL STRENGTHENING FORMULA PLUS WHITENING - potassium nitrate and sodium fluoride paste Team Technologies, 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.

-----

#### UP & Up Sensitive Toothpaste Enamel Strengthening formula plus whitening

#### **Active Ingredients**

Potassium nitrate 5%......Antihypersensitivity Sodium Fluoride 0.15% w/v fluoride ion .....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**

**If pain/sensitivity still persists** after 4 weeks of use, please visit your dentist. **Stop use and ask a dentist** if the problem 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 swallow 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 thoroughly at least 1 minute after each meal or at least twice a day or as directed by your dentist. Be sure to brush sensitive areas of the teeth.

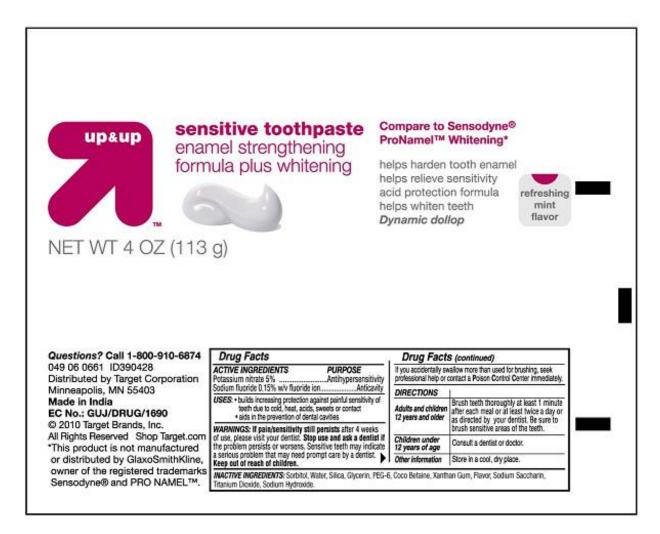
**Children under 12 years of age**Consult a dentist or doctor

**Other information** Store in a cool, dry place.

#### **Inactive ingredients:**

Sorbitol, Water, Silica, Glycerin, PEG-6, COCAMIDOPROPYL BETAINE, Xanthan Gum, Flavor, Sodium Saccharin, Titanium Dioxide, Sodium Hydroxide.

#### **Principal Display Panel**



Antihypersensitivity, Anticavity

KEEP OUT OF REACH OF CHILDREN

## UP AND UP SENSITIVE TOOTHPASTE ENAMEL STRENGTHENING FORMULA PLUS WHITENING

potassium nitrate and sodium fluoride paste

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:67659-081
Route of Administration	DENTAL		

Active Ingredient/Active Moiety				
Ingredient Name	Basis of Strength	Strength		
POTASSIUM NITRATE (UNII: RU45X2JN0Z) (POTASSIUM CATION - UNII:295O53K152)	POTASSIUM NITRATE	5 mg in 1 g		
SODIUM FLUORIDE (UNII: 8ZYQ1474W7) (FLUORIDE ION - UNII:Q80VPU408O)	SODIUM FLUORIDE	1.5 mg in 1 g		

Inactive Ingredients	
Ingredient Name	Strength
SORBITOL (UNII: 506T60A25R)	

WATER (UNII: 059QF0KO0R)	
SILICON DIO XIDE (UNII: ETJ7Z6 XBU4)	
GLYCERIN (UNII: PDC6A3C0OX)	
POLYETHYLENE GLYCOL 300 (UNII: 5655G9 Y8 AQ)	
COCAMIDO PRO PYL BETAINE (UNII: 50CF3011KX)	
XANTHAN GUM (UNII: TTV12P4NEE)	
SACCHARIN SO DIUM (UNII: SB8 ZUX40 TY)	
TITANIUM DIO XIDE (UNII: 15FIX9 V2JP)	
SO DIUM HYDRO XIDE (UNII: 55X0 4QC32I)	

Product Characteristics			
Color	white	Score	
Shape		Size	
Flavor	MINT (refreshing mint flavor)	Imprint Code	
Contains			

l	Packaging			
l	# Item Code	Package Description	Marketing Start Date	Marketing End Date
l	1 NDC:67659-081-02	1 in 1 CARTON		
ı	1 NDC:67659-081-01	113 g in 1 TUBE		

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph not final	part356	08/25/2010		

### Labeler - Team Technologies, Inc (192339703)

Revised: 8/2010 Team Technologies, Inc