# UP AND UP SENSITIVE TOOTHPASTE 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.

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### UP AND UP SENSITIVE TOOTHPASTE WHITENING

#### Active Ingredients

Potassium nitrate 5%.....Antihypersensitivity Sodium Fluoride 0.243% (0.15% w/v fluoride ion)....Anticavity toothpaste

#### **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** do not use longer than 4 weeks unless recommended by a dentist or doctor. **Stop and ask a dentist** if problems persists or worsens. Sensitive teeth may indicate a serious problem that may require prompt care by a dentist. **Keep out of reach of children**. If you accidentally swallow more that 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.

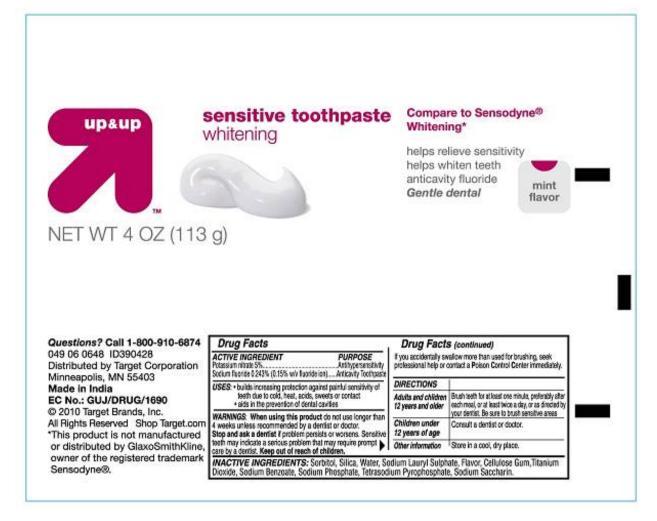
Children under 12 years of age Consult a dentist or doctor

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

#### **Inactive ingredients:**

Sorbitol, Silica, Water, Sodium Lauryl Sulphate, Flavor, Cellulose Gum, Titanium Dioxide, Sodium Benzoate, Sodium Phosphate, Tetrasodium Pyrophosphate, Sodium Saccharin

**Principal Display Panel** 



## Antihypersensitivity, Anticavity toothpaste KEEP OUT OF REACH OF CHILDREN

UP AND UP SENSITIVE TOOTHPASTE WHITENING									
potassium nitrate and sodium fluo	oride paste								
Product Information									
Product T ype	HUMAN OTC DRUG	Item Code (Source)		NDC:67659-079					
Route of Administration	DENTAL								
Active Ingredient/Active Mo	biety								
I	<b>Basis of Strength</b>		Strength						
POTASSIUM NITRATE (UNII: RU45X2JN0Z) (POTASSIUM CATION - UNII:295053K152) POTASSIUM					5 mg in 1 g				
SODIUM FLUORIDE (UNII: 8ZYQ1474W7) (FLUORIDE ION - UNII:Q80VPU408O) SODIUM FL					2.43 mg in 1 g				
Inactive Ingredients									
Ingredient Name					Strength				

SOR	<b>BITOL</b> (UNII: 506T60	) A25R)					
SILIC	CON DIO XIDE (UNII:	ETJ7Z6XBU4)					
WAT	<b>ER</b> (UNII: 059QF0KO	0 R)					
SOD	UM LAURYL SULFA	<b>ATE</b> (UNII: 368GB5141J)					
CRO	SCARMELLOSE SO	DIUM (UNII: M28OL1HH48)					
TITA	NIUM DIO XIDE (UNI	II: 15FIX9V2JP)					
	<b>UM BENZOATE</b> (UN						
SOD	<b>UM PHO SPHATE</b> (U	NII: SE337SVY37)					
		<b>ATE</b> (UNII: O352864B8Z)					
SACO	CHARIN SODIUM (UN	NII: SB8ZUX40TY)					
Pro	duct Characteris	stics					
Colo	r	white		Score			
Shap	e			Size			
Flavo	or	MINT (Mint Flavor)		Imprint Code			
Cont	ains						
Pac	kaging						
#	Item Code	Package Description	Marketing	g Start Date Mark		rketing End Date	
1 ND	C:67659-079-02	1 in 1 CARTON					
1 ND	C:67659-079-01	113 g in 1 TUBE					
Ma	rketing Infor	mation					
Ma	rketing Category	Application Number or Monog	raph Citation	Marketing Start	t Date	Marketing End Date	
OTC	monograph not final	part356		08/25/2010			

Labeler - Team Technologies, Inc (192339703)

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Team Technologies, Inc