**WHITENING FOAMING-TOOTHPASTE - sodium fluoride gel, dentifrice**
**Dio Corporation**

*Disclaimer: This drug has not been found by FDA to be safe and effective, and this labeling has not been approved by FDA. For further information about unapproved drugs, click here.*

---

## Drug Facts

active ingredients: sodium fluoride, allantoin

inactive ingredients: ethanol sodium lauryl sulfact, xylitol, sodium saccharin, ginger, mugwort extract, cnidium extract, mint, methylparaben, water

for dental care

keep out of reach of the children

brush teeth with your tooth brush or gargle after spraying it into your mouth

store at room temperature

use when needed
dental use only


## WHITENING FOAMING-TOOTHPASTE

sodium fluoride gel, dentifrice

| Product Information | | | |
|---|---|---|---|
| **Product Type** | HUMAN OTC DRUG | **Item Code (Source)** | NDC:75902-2001 |

| Route of Administration | DENTAL |
|---|---|

| Active Ingredient/Active Moiety | | |
|---|---|---|
| **Ingredient Name** | **Basis of Strength** | **Strength** |
| **SODIUM FLUORIDE** (UNII: 8ZYQ1474W7) (FLUORIDE ION - UNII:Q80VPU408O) | SODIUM FLUORIDE | 0.001 mL in 1 mL |
| **ALLANTOIN** (UNII: 344S277G0Z) (ALLANTOIN - UNII:344S277G0Z) | ALLANTOIN | 0.0005 mL in 1 mL |

| Inactive Ingredients | |
|---|---|
| **Ingredient Name** | **Strength** |
| **ALCOHOL** (UNII: 3K9958V90M) | |
| **SODIUM LAURYL SULFATE** (UNII: 368GB5141J) | |
| **XYLITOL** (UNII: VCQ006KQ1E) | |
| **SACCHARIN SODIUM** (UNII: SB8ZUX40TY) | |
| **GINGER** (UNII: C5529G5JPQ) | |
| **ARTEMISIA PRINCEPS LEAF** (UNII: SY077EW02G) | |
| **CNIDIUM OFFICINALE ROOT** (UNII: 8S3OZD358J) | |
| **MINT** (UNII: FV98Z8GITP) | |
| **METHYLPARABEN** (UNII: A2I8C7HI9T) | |
| **WATER** (UNII: 059QF0KO0R) | |

| Packaging | | | | |
|---|---|---|---|---|
| **#** | **Item Code** | **Package Description** | **Marketing Start Date** | **Marketing End Date** |
| **1** | NDC:75902-2001-1 | 50 mL in 1 BOTTLE, PUMP | | |

## Marketing Information

| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|---|---|---|---|
| unapproved drug other | | 10/04/2011 | |

## Labeler - Dio Corporation (631085206)

## Registrant - Dio Corporation (631085206)

## Establishment

| Name | Address | ID/FEI | Business Operations |
|---|---|---|---|
| Dio Corporation | | 631085206 | manufacture |

Revised: 10/2011 Dio Corporation