

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
DIO
TOOTHPASTE
화이트닝 거품치약



Allantoin / Sodium Fluoride
/ Hydroxyapatite | 50ml

제품문의 : 1599-2828
소비자가격 : 13,000원

품명 : 화이트닝 거품치약 효능효과 : 이를 희고 튼튼/구강청결/구강상쾌/충치예방/구취제거/치은염·치주염(치조농루)의 예방/치주질환 예방/잇몸질환 예방/심미효과 성분 : 하이드록시아파타이트, 쑥추출물, 천궁추출물, 건강틴크, 알란토인, 플루오르화 나트륨 등 용법용량 : 1일 1~수회 적당량을 칫솔 또는 입에 분무 후 칫솔질 또는 가글링을 한다. 저장방법 : 기밀용기, 실온(1~30℃)보관
사용기간 : 제조일로부터 36개월 제조원 :  한국콜마
제품개발 및 판매 : (주)디오 부산광역시 해운대구 우동 1464번지

본 제품은 공정거래 위원회 고시 소비자 분쟁해결기준에 의거 보상을 받을 수 있습니다.

Name : Whitening Foaming-Toothpaste **Indication :** Oral cleanness, Anti-halitosis, Prophylaxis of oral diseases(carries,gingiva disease,periodontal disease, periodontoclasia,gingivitis) **Active ingredients :** Sodium Fluoride, Allantoin **Also contains :** Hydroxy apatite, Cnidium Rhizome Extract, Ginger Tincture, Mugwort Extract etc. **Use :** Brush teeth with your tooth brush or gargle after spraying it into your mouth.
Caution : store at room temperature **LOT NO. & MFG. Date :** see crimp of container **Manufacture :** Korea Kolmar 618-3, Sinjung-ri, Jeonui-myeon, Yeongi-gun, Chungcheongnam-do, Korea
Product Development & Sales : **DIO Corporation** 1464, U-dong, Haeundae-gu, Busan, Korea
www.dioshop.net | MADE IN KOREA K100910

CLEAN TOOTH & MOUTHWASH
FOR SENSITIVE TEETH 50ml 1.76 oz.



다른배출
제품 : PP
캡 : PP
의약외품

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: FV98Z8G1TP)	
METHYL PARABEN (UNII: A2I8C7HI9T)	
WATER (UNII: 059QF0K00R)	

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