

DIO WHITENING- sodium fluoride paste, 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

colloidal silicon dioxide, tocopherol acetate, 35% hydrogen peroxide solution

xylityl, sorbitol solution 70%, polyethylene glycol 1500, carboxymethylcellulose sodium, xanthan gum, poly vinyl pyrrolidone, sodium stannate, etc.

for dental care

keep out of reach of the children

brush teeth with your toothbrush

store at room temperature

for dental use only

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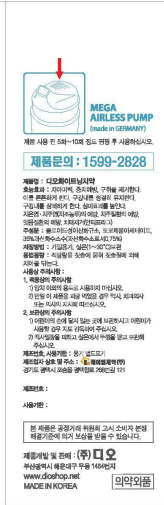
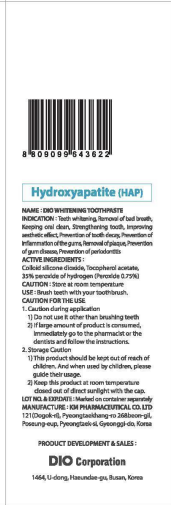


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DIO WHITENING

sodium fluoride paste, dentifrice

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:75902-400 1
Route of Administration	DENTAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
SILICON DIO XIDE (UNII: ETJ7Z6 XBU4) (SILICON DIO XIDE - UNII:ETJ7Z6 XBU4)	SILICON DIOXIDE	7 g in 100 g
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8 X80D2L0) (.ALPHA.-TOCOPHEROL - UNII:H4N8 55PNZ1)	.ALPHA.-TOCOPHEROL	0.2 g in 100 g
HYDROGEN PERO XIDE (UNII: BBX060 AN9 V) (HYDROGEN PERO XIDE - UNII:BBX060 AN9 V)	HYDROGEN PEROXIDE	2.14 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
XYLITOL (UNII: VCQ006KQ1E)	
WATER (UNII: 059QF0K00R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:75902-4001-1	90 g in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	01/22/2015	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		01/22/2015	

Labeler - Dio Corporation (631085206)**Registrant** - Dio Corporation (631085206)**Establishment**

Name	Address	ID/FEI	Business Operations
KMPPharmaceutical Co., Ltd.		688679158	manufacture(75902-4001)

Revised: 6/2017

Dio Corporation