DR DENTAL CARE 750ML-sodium fluoride liquid Jangin Pharm Co., Ltd.

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.

ACTIVE INGREDIENT

Active ingredient:

Sodium fluoride 0.02% (0.009% w/v fluoride ion)

INACTIVE INGREDIENT

Inactive ingredients:

Water, Alcohol (3%), Xylitol, Glycerin, Allantoin, Citric Acid, Sodium Citrate, Catechins, Sodium Benzoate, Menthol, Red Ginseng, Sophora Angustifolia Root, Swertia, Licorice, Caramel

PURPOSE

Purpose: anticavity

WARNINGS

Warning:

- 1. Keep out of reach of children under 6 years old.
- 2. Avoid direct sunlight, store in room temperature.
- 3. Close lid after use.
- 4. Consult doctor before use if you are allergic to any of the ingredients.
- 5. If significant amount is accidentally swallowed, drink large amount of milk and get medical help.

KEEP OUT OF REACH OF CHILDREN

Keep out of reach of children:

If significant amount is accidentally swallowed, drink large amount of milk and get medical help.

Use

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Mouthwash with outstanding oral disease prevention effect

Directions

Directions:

Rinse mouth with 10-15ml for 30 seconds.

PACKAGE LABEL.PRINCIPAL DISPLAY PANEL

Drodental care liquid





[Drug Facts]

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Dr. dental care liquid

식물성분 함유











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Drodental care liquid





치주염·치조농루·잇몸질환·치주질환 예방 구강내 악취제거·충치·치은염 예방, 이를 희고 튼튼하게!

Embodied a home oral care

The oral care composition using green tea as you can see outstanding results in tooth care product.



(750ml)

Dr. dental care liquid





DR DENTAL CARE 750ML

sodium fluoride liquid

Product Type HUMAN OTC DRUG Item Code (Source) NDC:69255-010

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name Basis of Strength Strength

SODIUM FLUORIDE (UNII: 8ZYQ1474W7) (FLUORIDE ION - UNII:Q80VPU408O)	FLUORIDE ION	0.15 mg in 750 mL
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Inactive Ingredients	
Ingredient Name	Strength
Water (UNII: 059QF0KO0R)	
Alcohol (UNII: 3K9958V90M)	

	Packaging			
ı	# Item Code	Package Description	Marketing Start Date	Marketing End Date
	1 NDC:69255-010-01	750 mL in 1 CARTON		

Marketing Information					
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date		
OTC monograph final	part355	08/01/2014			

Labeler - Jangin Pharm Co., Ltd. (688733680)

Registrant - Jangin Pharm Co., Ltd. (688733680)

Establishment					
Name	Address	ID/FEI	Business Operations		
Jangin Pharm Co., Ltd.		688733680	manufacture(69255-010)		

Revised: 9/2014 Jangin Pharm Co., Ltd.