KUNDAL PURE REFRESHING WHITENING TOOTH (SPEARMINT)- hydrogen peroxide, silicon dioxide paste THESKINFACTORY Co., Ltd.

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

Hydrogen Peroxide, Konasil (Fumed Silica = Silicon Dioxide)

Glycerin, PEG-32, Cellulose Gum, Xanthan Gum, Disodium Pyrophosphate, Salvia Officinalis (Sage) Leaf Extract, Eucalyptus Globulus Leaf Extract, Chamomilla Recutita (Matricaria) Flower Extact, Camellia Sinensis Leaf Extract, Disodium EDTA, Xylitol, Stevioside, Sodium Benzotate, Sodium Cocoyl Glumate, Menthol, Cooling Fragrance, Spearmint Oil, Spearmint Flavor, Water

for dental care

KEEP OUT OF REACH OF THE CHILDREN

Adults and children 2 years of age and older: Brush teeth thoroughly, preferably after each meal or at least twice a day, or as directed by a dentist or doctor.

Children 2 to 6 years of age: Instruct children under 6 years of age in good brushing and rinsing habits (to minimize swallowing). Supervise children as necessary until capable of using without supervision.

Children under 2 years of age: Consult a dentist or doctor.

Do not swallow.

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

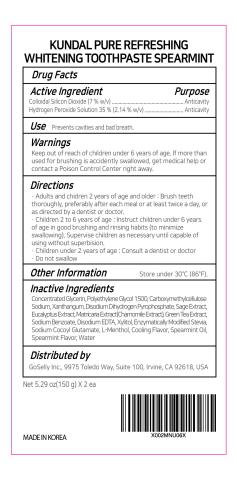
For dental use only

주식회사 더 스킨팩토리 <u>IMESKIN</u>		인쇄사양	
품목명	슈피겐_미백치약_2구단상자_OTC_덧방라벨	칼선	
수정날짜	2020 / 08 / 31	K100	
작업자	컨텐츠팀 최민용 010-2950-4855		

작업지시사항 HISTORY				
2020-08-31	슈피겐_미백치약_2구단상자_OTC_덧방라벨_신규			

규격: 70 X 140(mm)

※ 담당은 본 이미지를 사내 폴더에 올렸음을 확인합니다.※ 본 이미지는 사내 폴더에 올린 최종 데이타 이미지와 동일합니다.



1

KUNDAL PURE REFRESHING WHITENING TOOTH (SPEARMINT)

hydrogen peroxide, silicon dioxide paste

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:74773-0041	
Route of Administration	DENTAL			

Active Ingredient/Active Moiety		
Ingredient Name	Basis of Strength	Strength
HYDROGEN PERO XIDE (UNII: BBX060AN9V) (HYDROGEN PERO XIDE - UNII:BBX060AN9V)	HYDROGEN PEROXIDE	2.14 g in 100 g

Inactive Ingredients		
Ingredient Name	Strength	
MENTHO L (UNII: L7T10 EIP3A)		
XANTHAN GUM (UNII: TTV12P4NEE)		
XYLITOL (UNII: VCQ006KQ1E)		

]	Packaging			
7	# Item Code	Package Description	Marketing Start Date	Marketing End Date
:	NDC:74773-0041-1	2 in 1 BOX	09/01/2020	
:	1	150 g in 1 TUBE; Type 0: Not a Combination Product		

Marketing Information					
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date		
unapproved drug other		09/01/2020			

Labeler - THESKINFACTORY Co., Ltd. (694804099)

Registrant - THESKINFACTORY Co., Ltd. (694804099)

Establishment				
Name	Address	ID/FEI	Business Operations	
DONG IL PHARMS CO., LTD.		557810721	manufacture(74773-0041)	

Establishment				
Name	Address	ID/FEI	Business Operations	
THESKINFACTORY Co., Ltd.		694804099	label(74773-0041)	

Revised: 9/2020 THESKINFACTORY Co., Ltd.