

FOOT CLEAN GOLD- propolis wax powder KONEL

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.

ACTIVE INGREDIENT

Active Ingredient: Propolis Extract 12.0%

INACTIVE INGREDIENT

Inactive Ingredients: ERIOBOTRYA JAPONICA LEAF EXTRACT, SAURURUS CHINENSIS LEAF EXTRACT, PERILLA FRUTESCENS LEAF EXTRACT, ACORUS GRAMINEUS ROOT EXTRACT, ALOE BARBADENSIS LEAF POWDER, SOPHORA FLAVESCENS ROOT POWDER, ATRACTYLODES JAPONICA EXTRACT

PURPOSE

Purpose: Skin Protectant

WARNINGS

Warnings: For external use only When using this product do not get into eyes, Stop use and ask a doctor if rash occurs Do not use on - deep puncture wounds - animal bites - serious burns Keep out of reach of children

KEEP OUT OF REACH OF CHILDREN

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Use

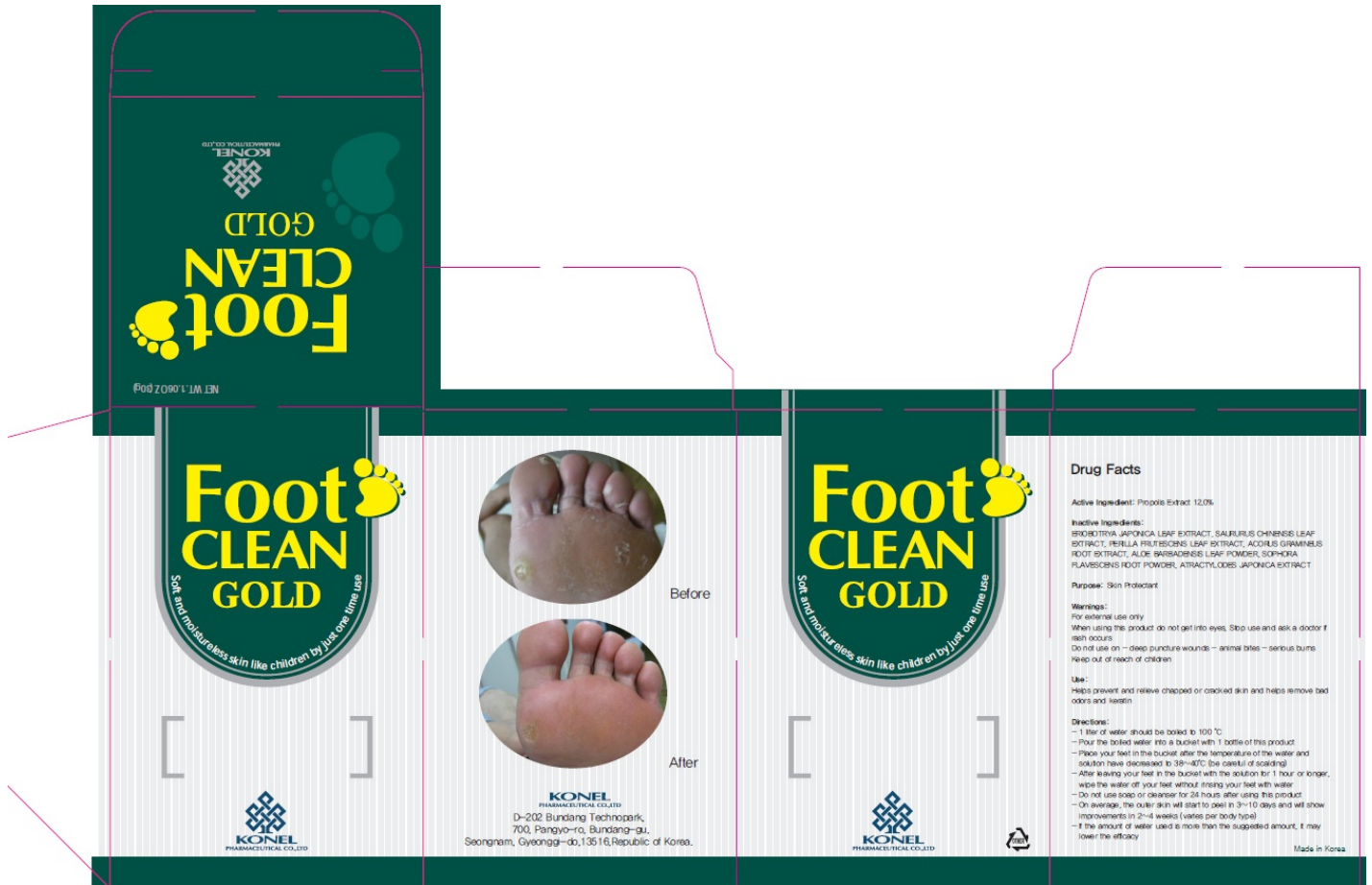
Use: Helps prevent and relieve chapped or cracked skin and helps remove bad odors and keratin

Directions

Directions: - 1 liter of water should be boiled to 100 °C - Pour the boiled water into a bucket with 1 bottle of this product - Place your feet in the bucket after the temperature of the water and solution have decreased to 38~40°C (be careful of scalding) - After leaving your feet in the bucket with the solution for 1 hour or longer, wipe the water off your feet without rinsing your feet with water - Do not use soap or cleanser for 24 hours after using this product - On average, the outer skin will start to peel in 3~10 days and will show improvements in 2~4 weeks (varies per body type) - If the amount

of water used is more than the suggested amount, it may lower the efficacy

PACKAGE LABEL.PRINCIPAL DISPLAY PANEL





FOOT CLEAN GOLD

propolis wax powder

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:71059-010
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
PROPOLIS WAX (UNII: 6Y8XYV2NOF) (PROPOLIS WAX - UNII:6Y8XYV2NOF)	PROPOLIS WAX	3.6 g in 30 g

Inactive Ingredients

Ingredient Name	Strength
ERIOBOTRYA JAPONICA LEAF (UNII: Z02066SV11)	
PERILLA FRUTESCENS LEAF (UNII: T4L5881Y68)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
	NDC:71059			

1	NDC: 71059-010-02	1 in 1 CARTON	11/01/2016	
1	NDC: 71059-010-01	30 g in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		
Marketing Information				
Marketing Category	Application Number or Monograph Citation		Marketing Start Date	Marketing End Date
unapproved drug other			11/01/2016	

Labeler - KONEL (689605036)

Registrant - KONEL (689605036)

Establishment			
Name	Address	ID/FEI	Business Operations
KONEL		689605036	manufacture(71059-010)

Revised: 10/2022

KONEL