GREENFISH, BODY-S- potassium carbonate liquid KOTUKU INC.

Disclaimer: This drug h	as not been found by F	DA to be safe and	l effective, and	this labeling l	has not been
approved by FDA. For	further information abo	ut unapproved dr	ugs, click here	2.	

Greenfish, Body-S

Potassium carbonate (0.1%)

antibacterial

Keep out of reach of children

Sanitizer to help decrease bacteria on skin

Warnings

For external use only

Do not use when liquid comes in contact with eyes, rinse immediately with water and consult the doctor.

When using this product

Avoid contact with eyes. If contact occurs, rinse eyes thoroughly with water.

Store at room temperature

Directions

Spray on skin as needed.

Inactive Ingredients

Calcium Hydroxide, Citric Acid, Tea Catechin, Hydroxypropylmethyl Cellulose, Distilled water

Greenfish, Body-S



ALL Natural

Body-Se

Natural Skin Care Solution Skin · Hair · Nails



- The best body sanitizer ever made.
- No harmful effects. Completely safe.
- Kills 99.999% of harmful bacteria that can cause Food Poisoning and Sickness.
- Can be used in any setting, on any part of the external human body.

MIST & FIX

80ml / 3.5fl.oz

Drug Facts Active Ingredient
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Sanitizer to help decrease bacteria on skin

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water
Manufactured by
KOTUKU INC.
3130 Wilshire Blvd. #402 Los Angeles CA 90010
Question or Comments
Call +1-213-805-6596

www.greenfishinc.com





GREENFISH, BODY-S

potassium carbonate liquid

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:70610-102

Route of Administration TOPICAL

Active Ingredient/Active Moiety

reave ingredient with the				
Ingredient Name	Basis of Strength	Strength		
POTASSIUM CARBONATE (UNII: BQN1B9B9HA) (CARBONATE ION - UNII:7UJQ5OPE7D)	POTASSIUM CARBONATE	0.001 in 80 mL		

Inactive Ingredients			
Ingredient Name	Strength		
CITRIC ACID MONO HYDRATE (UNII: 2968 PHW8 QP)			
CALCIUM HYDRO XIDE (UNII: PF5DZW74VN)			
HYPROMELLOSES (UNII: 3NXW29V3WO)			
water (UNII: 059QF0KO0R)			

]	Packaging				
#	# Item Code Package Description		Marketing Start Date	Marketing End Date	
1	NDC:70610-102- 01	80 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	04/18/2016		

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
unapproved drug other		04/07/2016		

Labeler - KOTUKU INC. (050192473)

Registrant - KOTUKU INC. (050192473)

Establishment				
Name	Address	ID/FEI	Business Operations	
KOTUKU INC.		050192473	manufacture(70610-102)	

Revised: 4/2016 KOTUKU INC.