STAIN REMOVER- stain remover(bamboo) liquid
Zhejiang Dibao Biotechnology 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.

C12-15 PARETH-10 5%

Sodium dodecyl sulphate 8%

OCTOXYNOL-11 4%

Bamboo extract 10.2%

Deionized water

Sodium phosphate, dibasic, dihydrate

Betaine

Coconut diethanolamide

HOW TO USE

Step 1-Wipe the fresh stains with a tissue or cloth
Step 2- Roll the ball until the liquid completely covers the stain
Step 3- After 10 seconds, gently wipe the stain area witha damp cloth

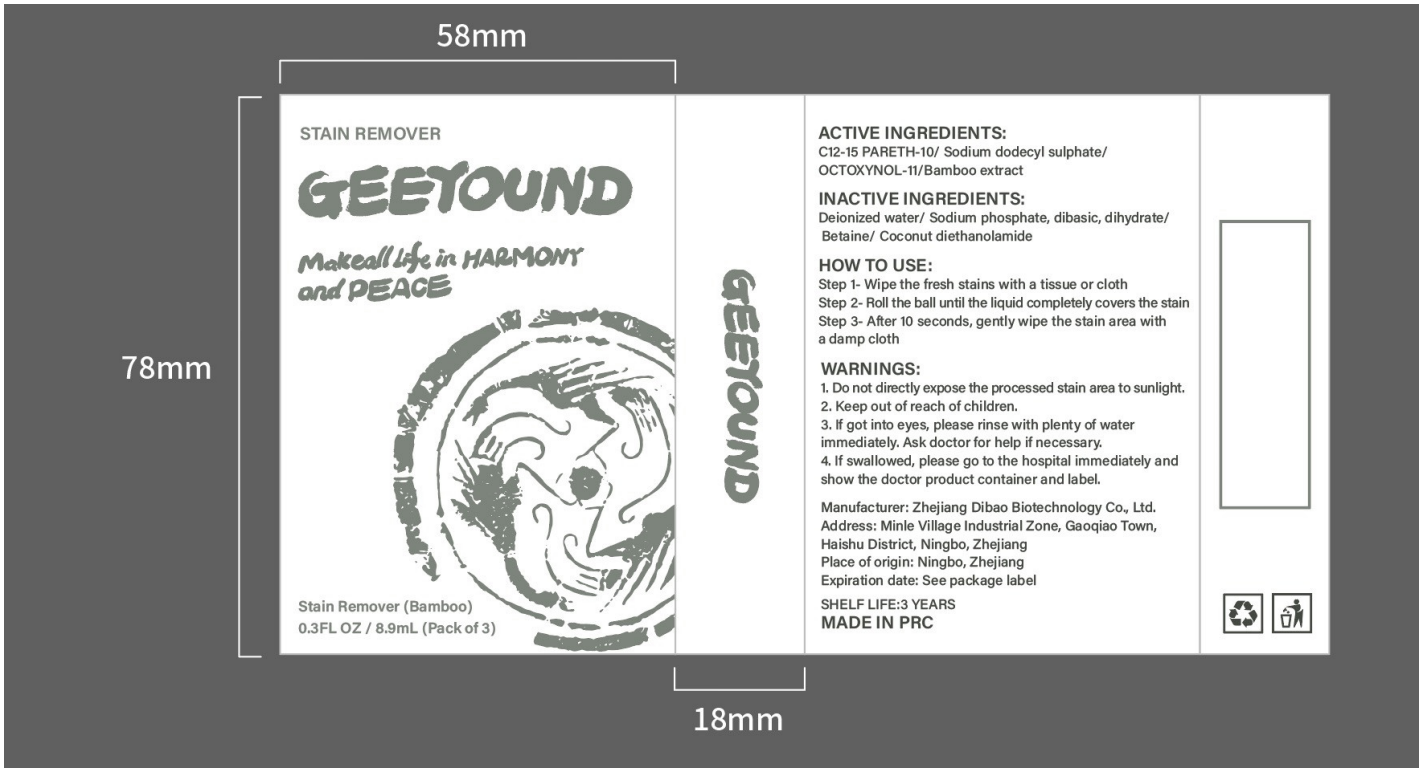
Spray enough product where you need to clean up to cover the dirt.

Keep out of reach of children.

In addition to stains

Keep in a cool and dry place

1.Do not directly expose the processed stain area to sunlight.
2.Keep out of reach of children.
3. If got into eyes, please rinse with plenty of water immediately. Ask doctor for help if necessary.
4. If swallowed, please go to the hospital immediately and show the doctor product container and label.



STAIN REMOVER

stain remover(bamboo) liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:82371-003
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
OCTOXYNOL-11 (UNII: SQL994V0M6) (OCTOXYNOL-11 - UNII:SQL994V0M6)	OCTOXYNOL-11	4 g in 100 mL
BAMBUSA VULGARIS TOP (UNII: FIW80T6P6V) (BAMBUSA VULGARIS TOP - UNII:FIW80T6P6V)	BAMBUSA VULGARIS TOP	10.2 g in 100 mL
SODIUM LAURYL SULFATE (UNII: 368GB5141J) (LAURYL SULFATE - UNII:DIQ16UC154)	SODIUM LAURYL SULFATE	8 g in 100 mL
C12-15 PARETH-10 (UNII: Z0QJT9586T) (C12-15 PARETH-10 - UNII:Z0QJT9586T)	C12-15 PARETH-10	5 g in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**
SODIUM PHOSPHATE, DIBASIC, DIHYDRATE (UNII: 94255I6E2T)	
BETAINE (UNII: 3SCV180C9W)	
WATER (UNII: 059QF0KO0R)	
COCO DIETHANOLAMIDE (UNII: 92005F972D)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:82371-003-01	8.9 mL in 1 BOTTLE; Type 0: Not a Combination Product	12/28/2021	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		12/28/2021	

Labeler - Zhejiang Dibao Biotechnology Co., Ltd. (603023294)

Establishment

Name	Address	ID/FEI	Business Operations
Zhejiang Dibao Biotechnology Co., Ltd.		603023294	manufacture(82371-003)

Revised: 12/2021 Zhejiang Dibao Biotechnology Co., Ltd.