LACTO BELLA Y- lactobacillus ferment, sophora flavescens root extract, lavandula angustifolia (lavender) oil, sodium chloride gel Bella Lab 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

Lactobacillus Ferment, Sophora Flavescens Root Extract, Lavandula Angustifolia (Lavender) Oil, Sodium Chloride

Water, etc.

It is one-time female cleaner contained subacidity fermentation extract and natural origin and, hygienic and easy to use.

It can help such as improvement odor, skin care, supply moisture and, pH balance.

keep out of reach of the children

After washing hands, opening individual packaging and removing the Y cap.

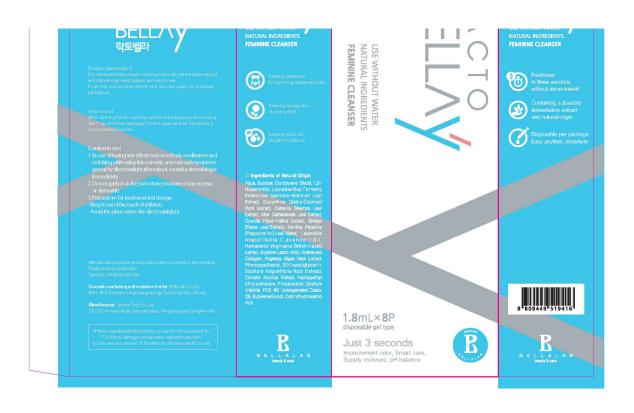
And then applying it to the required area.

Panty-liner is recommended outside.

- 1. In case of having side effects such as red rash, swollenness and /or itching while using this cosmetic, or in case such symptoms appear by direct sunlight after using it, consult a dermatologist immediately.
- 2. Do not apply it on the part where you have a scar, eczema or dermatitis.
- 3. Precautions for treatment and storage.
 - Keep it out of the reach of children.
 - Avoid the place where the direct sunlight is.

for external use only





LACTO BELLA Y

lactobacillus ferment, sophora flavescens root extract, lavandula angustifolia (lavender) oil, sodium chloride gel

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:72280-0001	
Route of Administration	VAGINAL			

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
LACTOBACILLUS REUTERI (UNII: 9913124QEE) (LACTOBACILLUS REUTERI - UNII:9913124QEE)	LACTOBACILLUS REUTERI	0.6 g in 100 mL	
SOPHORA FLAVESCENS ROOT (UNII: IYR6K8KQ5K) (SOPHORA FLAVESCENS ROOT - UNII:IYR6K8KQ5K)	SOPHORA FLAVESCENS ROOT	0.0015 g in 100 mL	
LAVENDER O IL (UNII: ZBP1YXW0H8) (LAVENDER O IL - UNII: ZBP1YXW0H8)	LAVENDER OIL	0.8 g in 100 mL	

SODIUM CHLORIDE (UNII: 451W47IQ8X) (CHLORIDE ION - UNII:Q32ZN48698, SODIUM SODIUM CHLORIDE CATION - UNII:LYR4M0NH37)

0.3 g in 100 mL

Inactive Ingredients

Strength **Ingredient Name**

WATER (UNII: 059QF0KO0R)

Packaging				
# Item Code	Package Description	Marketing Start Date	Marketing End Date	
NDC:72280- 0001-1	8 in 1 CARTON	0 4/27/20 18		
1	1.8 mL in 1 SYRINGE, 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		04/27/2018		

Labeler - Bella Lab Co., Ltd. (694921379)

Registrant - Bella Lab Co., Ltd. (694921379)

Establishment			
Name	Address	ID/FEI	Business Operations
Sphere Tech Co.,Ltd.		689907467	manufacture(72280-0001), pack(72280-0001), label(72280-0001)

Establishment				
Name	Address	ID/FEI	Business Operations	
Bella Lab Co., Ltd.		694921379	relabel(72280-001)	

Revised: 4/2018 Bella Lab Co., Ltd.