LESONEL FEMININE FOAMING WASH- allantoin liquid
ACE PHARMACEUTICAL 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

ALLANTOIN

Water
Disodium Cocoamphodiacetate
Sodium Chloride
1,2-Hexanediol

Decyl Glucoside
Butylene Glycol
Hexylene Glycol
Lactobacillus/Soybean Ferment Extract
Streptococcus Thermophilus Ferment
Bifida Ferment Lysate

Lactobacillus Ferment Lysate
Hydrogenated Lecithin

Ceramide NP
Allantoin
Trehalose
Betaine
Sodium PCA
Citric Acid
Disodium EDTA

Methylpropanediol
Caprylyl Glycol
Ethylhexylglycerin
Steareth-30

The product, made from naturally derived safe surfactants, effectively maintains cleanliness and moisture

in the external genital area, providing a refreshing cleansing sensation without causing irritation.

KEEP OUT OF REACH OF THE CHILDREN

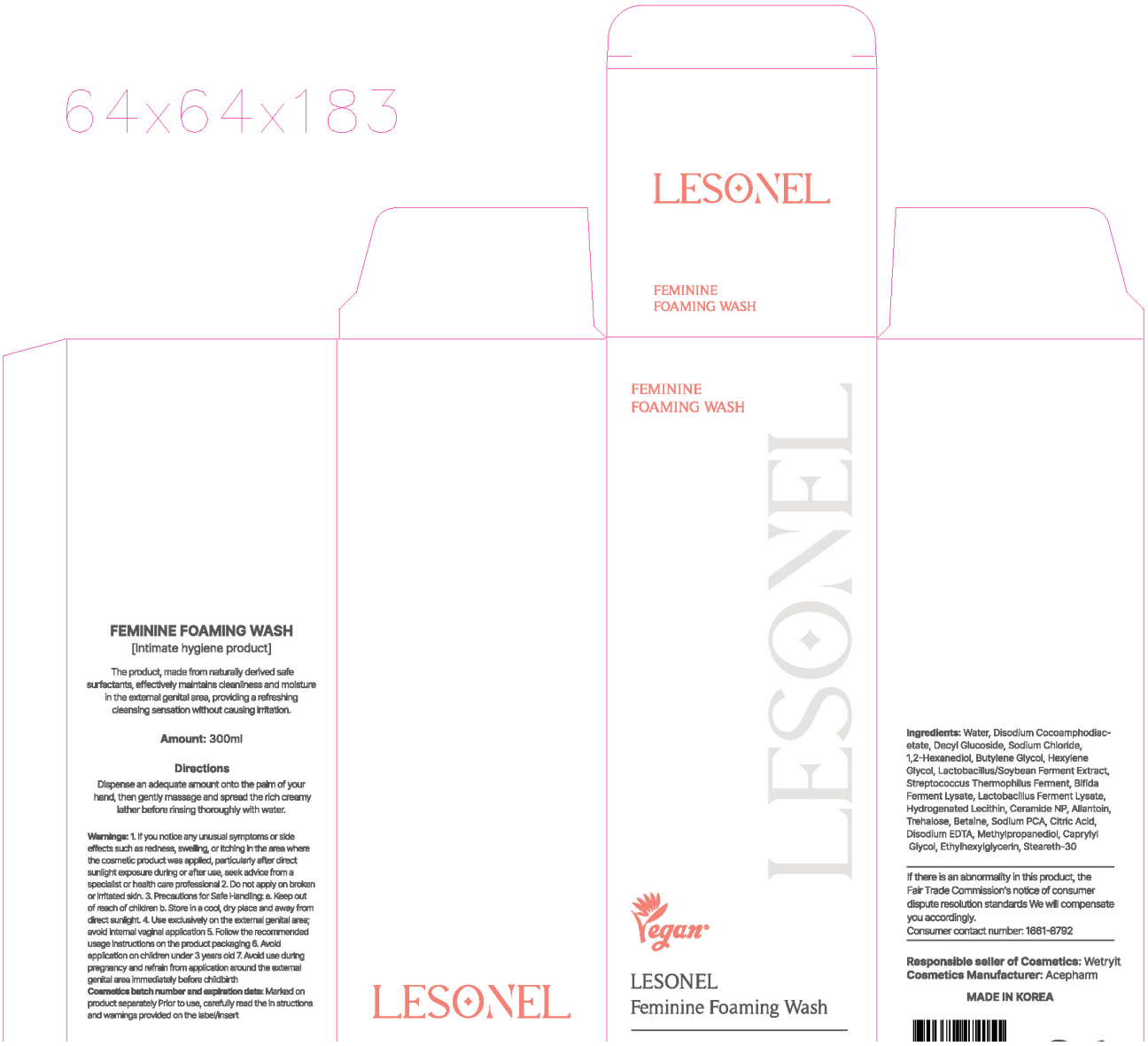
Dispense an adequate amount onto the palm of your hand, then gently massage and spread the rich creamy

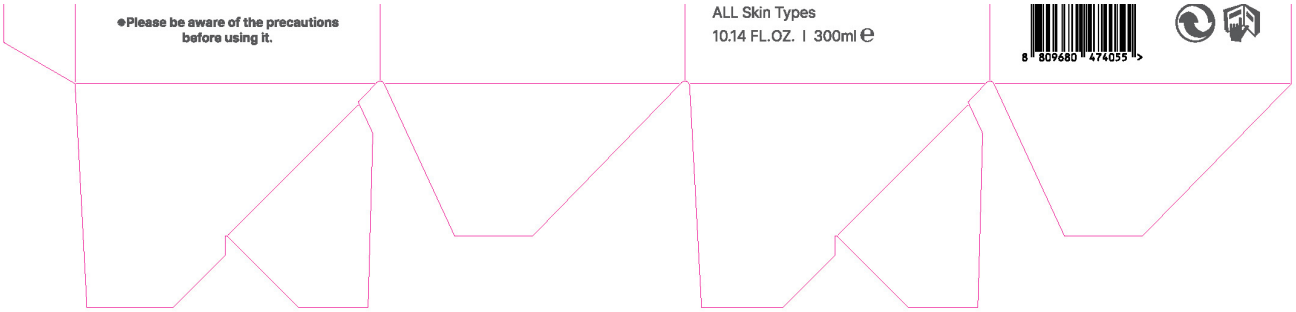
lather before rinsing thoroughly with water.

Warnings: 1. If you notice any unusual symptoms or side effects such as redness, swelling, or itching in the area where the cosmetic product was applied, particularly after direct sunlight exposure during or after use, seek advice from a specialist or health care

professional 2. Do not apply on broken or irritated skin. 3. Precautions for Safe Handling: a. Keep out of reach of children b. Store in a cool, dry place and away from direct sunlight. 4. Use exclusively on the external genital area; avoid internal vaginal application 5. Follow the recommended usage instructions on the product packaging 6. Avoid application on children under 3 years old 7. Avoid use during pregnancy and refrain from application around the external genital area immediately before childbirthPrecautions for Safe Handling: a. Keep out of reach of children

for vaginal use only





LESONEL FEMININE FOAMING WASH

allantoin liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:74196-0011
Route of Administration	VAGINAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALLANTOIN (UNII: 344S277G0Z) (ALLANTOIN - UNII:344S277G0Z)	ALLANTOIN	0.1 g in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:74196-0011-1	300 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	01/01/2025	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		01/01/2025	

Labeler - ACE PHARMACEUTICAL CO.,LTD. (689058677)

Registrant - ACE PHARMACEUTICAL CO.,LTD. (689058677)

Establishment

Name	Address	ID/FEI	Business Operations
ACE PHARMACEUTICAL CO LTD		689058677	manufacture(74196-0011)

Revised: 3/2025

ACE PHARMACEUTICAL CO.,LTD.