LESONEL FORMING FEMININE WASH- methylpropanediol 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

METHYLPROPANEDIOL

Water

Disodium Cocoamphodiacetate

Decyl Glucoside

Sodium Chloride

1,2-Hexanediol

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

Caprylyl Glycol

Ethylhexylglycerin

Steareth-30

The product, made from naturally deriveds afesurfactants, effectively maintains cleanliness and moisturein the external genital area,

providingarefreshing 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.

- 1. If you notice any unusual symptoms or side effects such as redness, swelling, or itching in the area where the cos metic product was applied, particularly after direct sunlight expo sure 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

for vaginal use only



LESONEL FORMING FEMININE WASH

methylpropanediol liquid

Product Information

Route of Administration VAGINAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
PANEDIOL (UNII: N8F53B3R4R) (METHYLPROPANEDIOL -	METHYL DDODANEDIOL	0.685 g

METHYLPROPANEDIOL (UNII: N8F53B3R4R) (METHYLPROPANEDIOL - UNII: N8F53B3R4R)

METHYLPROPANEDIOL 0.685 g in 100 mL

Inactive Ingredients

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

Packaging

ı	. acitaging			
	# Item Code	Package Description	Marketing Start Date	Marketing End Date
	1 NDC:74196-0007-1	300 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/18/2024	

Marketing Information

ranketing in					
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date		
unapproved drug other		01/18/2024			

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(74196-0007)	

Revised: 1/2024 ACE PHARMACEUTICAL CO LTD