

KENAF WET WIPE- allantoin liquid
MEDICELL BIO 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 0.5%

Water, etc

Wet tissue

keep out of reach of the children

Apply to skin and wipe

1. For external use only.

1) Do not use over the wound and irritated skin with eczema or infections

2) Discontinue using the product if signs of irritations and/or rashes occur and seek professional medical help

2. Keep the cover closed tight after use

3. Do not put the left over contents back into the container to avoid deterioration

1) Avoid contact with eyes

2) Keep out of reach of children

3) Store in a cool and dry place, away from direct sun light

for external use only



KENAF WET WIPE

allantoin liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:72944-0001
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALLANTOIN (UNII: 344S277G0Z) (ALLANTOIN - UNII:344S277G0Z)	ALLANTOIN	0.05 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0K00R)	
1,2-HEXANEDIOL (UNII: TR046Y3K1G)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date

1	NDC:72944-0001-1	190 g in 1 POUCH; Type 0: Not a Combination Product	04/09/2017	
Marketing Information				
Marketing Category	Application Number or Monograph Citation		Marketing Start Date	Marketing End Date
unapproved drug other			04/09/2017	

Labeler - MEDICELL BIO CO., Ltd. (689843703)

Registrant - MEDICELL BIO CO., Ltd. (689843703)

Establishment			
Name	Address	ID/FEI	Business Operations
MEDICELL BIO CO., Ltd.		689843703	manufacture(72944-0001)

Revised: 2/2023

MEDICELL BIO CO., Ltd.