UREA 40 PLUS HA- urea gel SSG Ventures Inc

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.

Urea 40 Plus HA



ACTIVE INGREDIENTS:

UREA

SOOTHES ROUGH & DRY SKIN

HEALS CORNS & CALLOUSES

SOFTENS NAILS

SOOTHES ROUGH & DRY SKIN

FOR EXTERNAL USE ONLY. AVOID CONTACT TO EYES.

KEEP OUT OF REACH OF CHILDREN. STOP USE AND CONSULT DOCTOR IF CONDITION WORSENS OR CLEARS UP AND REOCCURS.

KEEP LID FIRMLY CLOSED. STORE IN COOL DRY PLACE.

KEEP OUT OF REACH OF CHILDREN.

APPLY TO AFFECTED AREA AT LEAST TWICE DAILY OR AS NEEDED.

ALOE VERA, CARBOMER, COCONUT OIL, EMULSIFIERS, GREEN TEA, MINERAL OIL, PRESERVED WATER, POPYLENE GLYCOL, SODIUM HYALURONATE, TEA TREE OIL, TRIETHANOLOAMINE, XANTHAM GUM

KEEP LID FIRMLY CLOSED. STORE IN COOL DRY PLACE.

ENHANCED WITH TEA TREE OIL & ALOE VERA

APPLY TO AFFECTED AREA AT LEAST TWICE DAILY OR AS NEEDED.USE CONTINUOUSLY FOR 2-3 WEEKS FOR OPTIMAL RESULTS.

FOR EXTERNAL USE ONLY. AVOID CONTACT TO EYES. STOP USE AND CONSULT DOCTOR IF CONDITION WORSENS OR CLEARS UP AND REOCCURS.

UREA 40 PLUS HA			
urea gel			
Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:81376-213
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
UREA (UNII: 8W8T17847W) (UREA - UNII:8W8T17847W)	UREA	40 g in 113 g	

Inactive Ingredients		
Ingredient Name	Strength	
CARBO XYPO LYMETHYLENE (UNII: 0 A5MM30 7FC)		
XANTHAN GUM (UNII: TTV12P4NEE)		
MINERAL OIL (UNII: T5L8T28FGP)		
COCONUT OIL (UNII: Q9L0O73W7L)		
WATER (UNII: 059QF0KO0R)		
TRIDECETH-10 (UNII: G624N6MSBA)		
TEA TREE OIL (UNII: VIF565UC2G)		
PROPYLENE GLYCOL (UNII: 6 DC9 Q16 7 V3)		
ALOE VERA LEAF (UNII: ZY81Z83H0 X)		
HYALURONATE SODIUM (UNII: YSE9 PPT4TH)		

Product Characteristics			
Color	white (Opaque White)	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

Packaging				
ı	# Item Code	Package Description	Marketing Start Date	Marketing End Date
ı	1 NDC:81376-213-01	113 g in 1 JAR; Type 0: Not a Combination Product	09/20/2019	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		09/20/2019	

Labeler - SSG Ventures Inc (047626115)

Establishment				
Name	Address	ID/FEI	Business Operations	
SSG Ventures Inc		047626115	manufacture(81376-213)	

Revised: 2/2021 SSG Ventures Inc