CLEAR 40 PLUS SA- 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.

Clear 40 Plus SA



ACTIVE INGREDIENTS:

UREA 40%, SALICYLIC ACID 2%, TEA TREE OIL, ALOE VERA

SOOTHES ROUGH & DRY SKIN

HEALS CORNS & CALLOUSES

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.USE CONTINUOUSLY FOR 2-3 WEEKS FOR OPTIMAL RESULTS.

CARBOMER, XANTHAN GUM, MINERAL OIL, PROPYLENE GLYCOL, EMULSIFIERS, TRIETHANOLAMINE

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.

CLEAR 40 PLUS SA

urea gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:81376-232
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	
SALICYLIC ACID (UNII: O414PZ4LPZ) (SALICYLIC ACID - UNII:O414PZ4LPZ)	SALICYLIC ACID	6.5 g in 113 g	

Inactive Ingredients				
Ingredient Name	Strength			
ALOE VERA LEAF (UNII: ZY81Z83H0X)				
CARBOXYPOLYMETHYLENE (UNII: 0A5MM307FC)				
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: 6DC9Q167V3)				

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

F	Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:81376- 232-01	113 g in 1 JAR; Type 0: Not a Combination Product	09/20/2019		
2	NDC:81376- 232-02	226 g in 1 BOTTLE, DISPENSING; 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(81376-232)	

Revised: 4/2021 SSG Ventures Inc