

CRUEX PETROLEUM JELLY- white petrolatum jelly
MKJ BRANDS LLC

Cruex Petroleum Jelly

Drug Facts

Active Ingredient

White Petrolatum USP 30%

Purpose

Skin protectant

Uses

temporarily protects ■ minor cuts, scrapes, burns ■ and help relieve chapped or cracked skin and lips ■ from the drying effects of wind and cold weather

Warnings

For external use only

When using this product

do not get into eyes

Stop use and consult a doctor

if the condition lasts more than 7 days.

Do not use on

■ deep or puncture wounds
■ animal bites ■ serious burns

Keep out of reach of children

If swallowed, get medical help or contact a Poison Control Center right away. (1-800-222-1222)

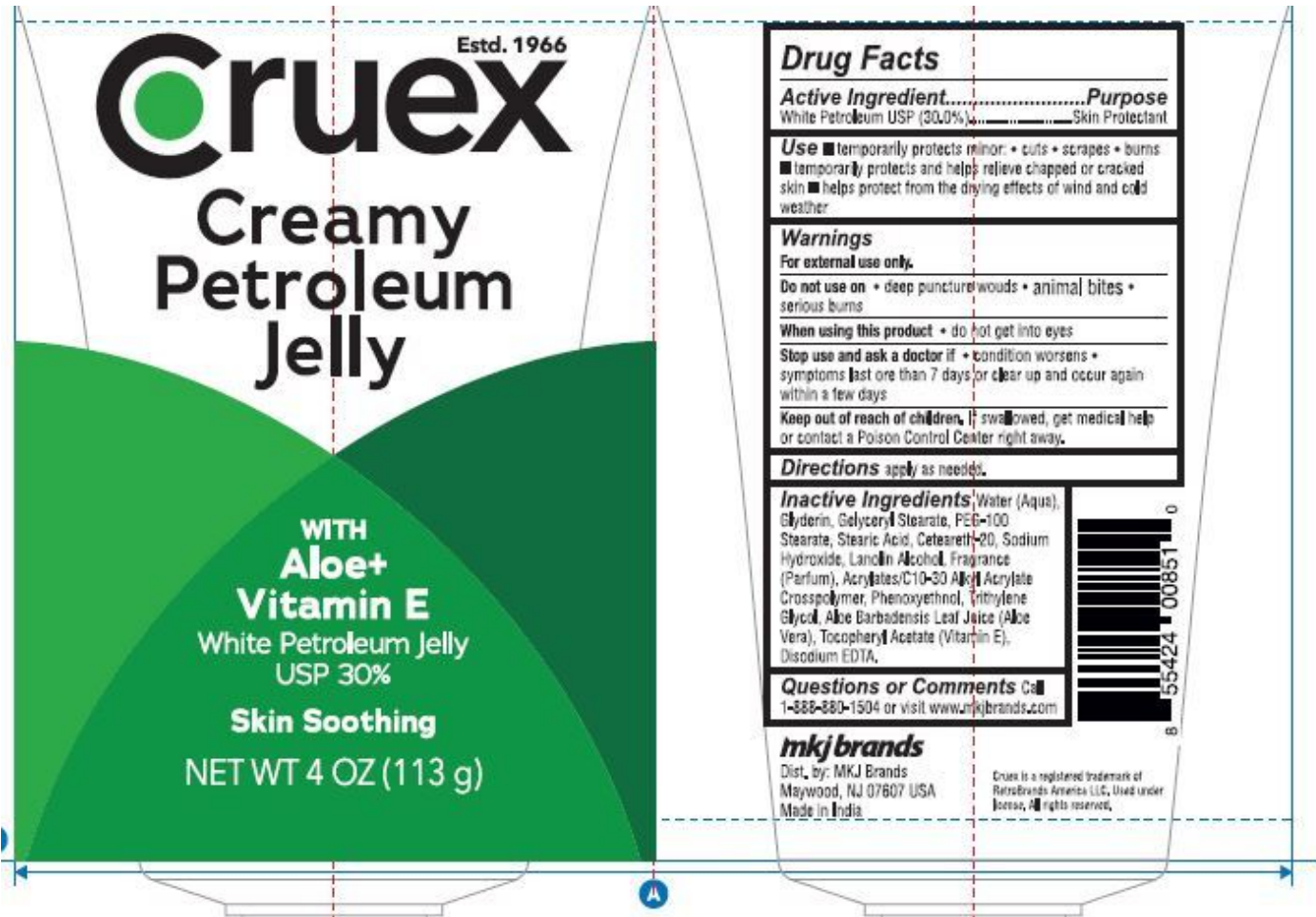
Directions

apply as needed

Aloe barbadensis leaf extract, frafrance, tocopheryl acetate, water, glycerin, gelyceryl stearate, PEG-100 Stearate, stearic acid, sodium hydroxide, Lanolin alcohol, Fragrance,

Acrylates 10-30 alkyl acrylate crosspolymer, pheonxyethanol, triethylene glycol, dospdoium EDTA

Petrolatum Jelly



CRUEX PETROLEUM JELLY

white petrolatum jelly

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:77782-004
Route of Administration	TOPICAL		
Active Ingredient/Active Moiety			
Ingredient Name		Basis of Strength	Strength
PETROLATUM (UNII: 4T6H12BN9U) (PETROLATUM - UNII:4T6H12BN9U)		PETROLATUM	30 g in 100 g
Inactive Ingredients			
Ingredient Name			Strength
LANOLIN ALCOHOL (UNII: 884C3FA9HE)			
PHENOXYETHANOL (UNII: HIE492ZZ3T)			

ACRYLATES/C10-30 ALKYL ACRYLATE CROSSPOLYMER (60000 MPA.S) (UNII: 8Z5ZAL5H3V)				
STEARIC ACID (UNII: 4ELV7Z65AP)				
AQUA (UNII: 059QF0KO0R)				
GLYCERIN (UNII: PDC6A3C0OX)				
CETEARETH-20 (UNII: YRC528SWJY)				
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)				
ALOE BARBADENSIS LEAF (UNII: ZY81Z83H0X)				
EDETATE DISODIUM (UNII: 7FLD91C86K)				
PEG-100 MONOSTEARATE (UNII: YD01N1999R)				
TRIETHYLENE GLYCOL (UNII: 3P5SU53360)				
GLYCERYL STEARATE (UNII: 230OU9XXE4)				
SODIUM HYDROXIDE (UNII: 55X04QC32I)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:77782-004-13	113 g in 1 JAR; Type 0: Not a Combination Product	12/10/2025	
Marketing Information				
Marketing Category		Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug		M016	09/09/2025	

Labeler - MKJ BRANDS LLC (827648101)

Revised: 12/2025

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