VERADEX-E- petrolatum and lanolin ointment
ABBE Laboratories, Inc.

Drug Facts

Active Ingredients
Petrolatum USP 58.95%
Anhydrous Lanolin USP 35%

Purpose
Petrolatum USP 58.95% ..... Protectant
Anhydrous Lanolin USP 35% ..... Protectant

Other ingredients: Tocopheryl Acetate, Cetyl Esters, Ethyl Methylphenylglycidate, Benzyl Cinnamate, Salicylic Acid.

Warnings: For external use only.
Discontinue use if signs of irritation appear. If condition worsens consult a doctor.
Do not use over deep, infected, or puncture wounds.

Keep from children.

Indications: Temporarily relieves minor burns, dry, chapped, fissured and macerated skin and lips, also relieves Psoriais, Eczema, mild sunburn, abrasions and contusions, diaper dermatitis, hemorrhoids, and prickly heat.

Directions: Apply a thin film to clean, dry skin three times daily. May be covered with sterile gauze, if desired.

NDC 68605-3700-5

VERADEX-E® OINTMENT

Super Emollient, Protectant
Healing Ointment

Net Wt. 100g.

U.S. Pat. # 5,645,826

ABBE LABORATORIES, INC.™
Farmingdale, NY 11735

VERADEX-E

petrolatum and lanolin ointment

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:68605-3700
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
PETROLATUM (UNII: 4T6H12BN9U) (PETROLATUM - UNII:4T6H12BN9U)	PETROLATUM	589.5 mg in 1 g
LANOLIN (UNII: 7EV65EAW6H) (LANOLIN - UNII:7EV65EAW6H)	LANOLIN	350 mg in 1 g

Inactive Ingredients

Ingredient Name	Strength
ALPHA-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
CETYL ESTERS WAX (UNII: D072FFP9GU)	
ETHYL METHYLPHENYLGLYCIDATE (UNII: UD51D5KR4A)	
BENZYL CINNAMATE (UNII: V67O3RO97U)	
SALICYLIC ACID (UNII: O414PZ4LPZ)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:68605-3700-2	15 g in 1 JAR; Type 0: Not a Combination Product	01/01/2008	
2	NDC:68605-3700-3	28 g in 1 JAR; Type 0: Not a Combination Product	01/01/2008	
3	NDC:68605-3700-4	56 g in 1 JAR; Type 0: Not a Combination Product	01/01/2008	
4	NDC:68605-3700-5	100 g in 1 JAR; Type 0: Not a Combination Product	01/01/2008	10/31/2023
5	NDC:68605-3700-6	400 g in 1 JAR; Type 0: Not a Combination Product	01/01/2008	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M016	01/01/2008	

Labeler - ABBE Laboratories, Inc. (781745286)

Establishment

Name	Address	ID/FEI	Business Operations
ABBE Laboratories, Inc.		781745286	manufacture(68605-3700)

Revised: 11/2023

ABBE Laboratories, Inc.