TEEVODAR- allantoin cream
PEER PHARM LTD.

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

Active Ingredient

Allantoin 0.5%

Purpose

Skin Protectant

Uses

Temporarily protects minor cuts scrapes burns

Warnings

- **For external use only**
- **When using this product** do not get into eyes
- **Stop use and ask a doctor if**
- contition worsens
- symptoms last more than 7 days or clear up and occur again within a few days
- **Keep out of reach of children.** If swallowed, get medical help or contact a Poison Control Center right away.

Directions

- Apply as needed.
- Children under 12 years of age: consult a doctor

Inactive Ingredients

Ceteareth-20, Cetearyl alcohol, Cetyl alcohol, ethylhexylglycerin, Hamamelis virginiana (witch hazel) leaf extract, Isopropyl myristate, Lanolin, Mineral oil, Origanum vulgare (oregano) leaf extract, Petrolatum, Phenoxyethanol, Purified water, Rosmarinus officinalis (rosemary) leaf extract, Zea mayes (corn) oil

Product label

TEEVODAR

allantoin cream

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:69435-1301

Route of Administration	TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALLANTOIN (UNII: 344S277G0Z) (ALLANTOIN - UNII:344S277G0Z)	ALLANTOIN	0.5 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
POLYOXYL 20 CETOSTEARYL ETHER (UNII: YRC528SWUY)	
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)	
CETYL ALCOHOL (UNII: 936JST6JCN)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
HAMAMELIS VIRGINIANA LEAF (UNII: T07U1161SV)	
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)	
LANOLIN (UNII: 7EV65EAW6H)	
MINERAL OIL (UNII: T5L8T28FGP)	
OREGANO (UNII: 0E5AT8T16U)	
PETROLATUM (UNII: 4T6H12BN9U)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
WATER (UNII: 059QF0KO0R)	
ROSEMARY (UNII: IJ67X351P9)	
CORN OIL (UNII: 8470G57WFM)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69435-1301-1	1 in 1 CARTON	05/15/2023	
1		30 g in 1 TUBE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part347	05/15/2023	

Labeler - PEER PHARM LTD. (514678390)

Registrant - PEER PHARM LTD. (514678390)

Revised: 5/2023 PEER PHARM LTD.