

**BAG BALM SKIN PROTECTANT- petrolatum ointment  
VERMONT`S ORIGINAL, LLC**

*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.*

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**Drug Facts**

<b>Active ingredient</b>	<b>Purpose</b>
Petrolatum 84.3%	Skin Protectant

**Uses**

- temporarily protects minor: • cuts • scrapes • burns
- temporarily protects and helps relieve chapped or cracked skin and lips
- helps protect 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 ask a doctor if**

- condition worsens
- symptoms last more than 7 days or clear up and occur again within a few 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.

**Directions** Apply as needed

**Inactive Ingredients** Benzethonium Chloride, Lanolin, Paraffin Wax, Purified Water

**Questions?** 1-844-424-2256 • [www.bagbalm.com](http://www.bagbalm.com)

**PACKAGE LABEL**

**BAG BALM®**  
SKIN PROTECTANT

**FIRST AID**

For Minor Cuts, Burns, Scrapes, or Skin Abrasions

fragrance free





## BAG BALM SKIN PROTECTANT

petrolatum ointment

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:69845-031
<b>Route of Administration</b>	TOPICAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
Petrolatum (UNII: 4T6H12BN9U) (Petrolatum - UNII:4T6H12BN9U)	Petrolatum	84.3 g in 100 g

### Inactive Ingredients

Ingredient Name	Strength
Benzethonium Chloride (UNII: PH41D05744)	

<b>Lanolin</b> (UNII: 7EV65EAW6H)	
<b>Paraffin</b> (UNII: I9O0E3H2ZE)	
<b>Water</b> (UNII: 059QF0K00R)	

<b>Packaging</b>				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69845-031-04	113 g in 1 CAN; Type 0: Not a Combination Product	03/01/2018	
2	NDC:69845-031-02	57 g in 1 TUBE; Type 0: Not a Combination Product	03/01/2018	

<b>Marketing Information</b>			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part347	03/01/2018	

**Labeler** - VERMONT`S ORIGINAL, LLC (079593652)

<b>Establishment</b>			
Name	Address	ID/FEI	Business Operations
Denison Pharmaceuticals, LLC		001207208	manufacture(69845-031)

Revised: 7/2017

VERMONT`S ORIGINAL, LLC