# DR. SAYMAN SALVE- salve ointment Sheffield Pharmaceuticals 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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#### **Dr Sayman Salve**

#### Active ingredient

Petrolatum - 87.3 %

Zinc Oxide-8.8%

#### Purpose

Skin protectant

Sunscreen

#### Uses

- Temporarily protects minor cuts, scrapes and burns.
- Helps releive chapped or cracked skin and lips.
- Helps protectfrom the drying effects of windand cold weather.
- Providesminimal portection agaisnt sunburn.
- Helps prevent sunburn.

#### Warnings

#### For external use only

**Sun Alert:** limiting sun exposure, wearing protectvie clothing, and using sunscreens may reduce the risks of skin aging, skin cancer, and other harmful effects of the sun.

- When using this product keep out of eyes. Rinse with water to remove.
- Stop use and ask a doctor if condition worsens or symptoms last more than seven days or clear up and occur again within a few days.
- Don't use on deep/ puncture wounds, animal bites or serious burns.

# Keep out of reach of children if swallowed, get medical help or contact a Poison Control Center right away.

#### **Directions**

- Apply liberally before exposureand as needed
- Children under 6 months of age:ask a doctor
- Reapply as needed or after towel drying, swimming

#### Other information

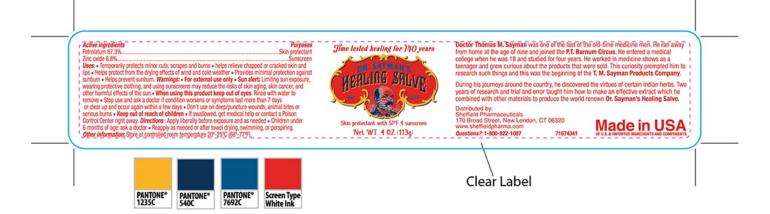
Store at controlled room temperature 20°C -25°C (68°F - 77°F)

#### **Inactive ingredients**

### Lanolin, Propylene Glycol, Fragrance, Quaternium-15

# **Principal Display Panel - Jar**

Dr Sayman's Salve NDC 11527-152-68
Healing Salve
Ideal for Dry, Chapped or Irritated skin
Also for minor cut,scrapes and burns
NET WT 4 oz (113g)



### **Principal Display Panel - Carton**

Dr Sayman's Salve NDC 11527-152-68
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# DR. SAYMAN SALVE

salve ointment

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:11527-152
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
PETROLATUM (UNII: 4T6 H12BN9 U) (PETROLATUM - UNII:4T6 H12BN9 U)	PETROLATUM	87.3 mg in 1 g	
ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC CATION - UNII:13S1S8SF37)	ZINC CATION	8.8 mg in 1 g	

Inactive Ingredients		
Ingredient Name	Strength	
LANOLIN (UNII: 7EV65EAW6H)		
PROPYLENE GLYCOL (UNII: 6 DC9 Q167V3)		
QUATERNIUM-15 (UNII: E40 U0 3LEM0)		

Product Characteristics			
Color	white	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

ı	Packaging			
l	# Item Code	Package Description	Marketing Start Date	Marketing End Date
l	1 NDC:11527-152-68	1 in 1 CARTON	0 1/16/20 15	
1	1	113 g in 1 JAR; 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	0 1/16/20 15	

# Labeler - Sheffield Pharmaceuticals LLC (151177797)

Establishment			
Name	Address	ID/FEI	Business Operations
Sheffield Pharmaceuticals LLC		151177797	manufacture(11527-152)

Revised: 4/2020 Sheffield Pharmaceuticals LLC