#### HARMON FACE VALUES ADVANCED SEVERE DRY SKIN THERAPYpetrolatum ointment Delta Pharma Inc.

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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# Harmon Face Values Advanced Skin Ointment

#### **Active Ingredient**

Petrolatum (41%)

#### Purpose

Skin Protectant (ointment)

# 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

# Warning

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 the reach of children.

If swallowed, seek medical help or contact a Poison Control Center right away.

# Directions

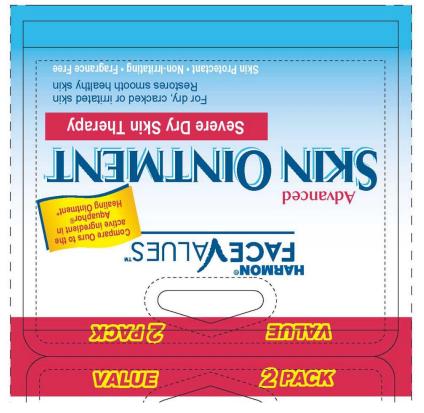
Apply as needed

Inactive Ingredients

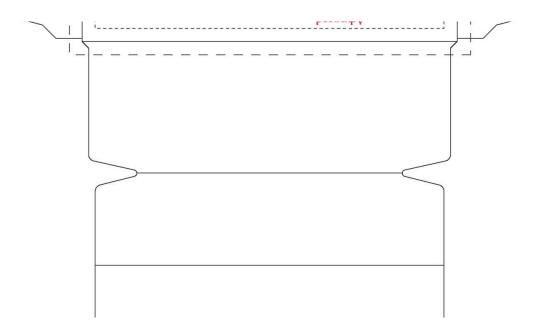
Mineral Oil, Ceresin, Lanolin Alcohol, Panthenol, Glycerin, Bisabolol











# HARMON FACE VALUES ADVANCED SEVERE DRY SKIN THERAPY

petrolatum ointment

P	roduct Informat	tion					
Product Type HUMAN OTC DRUG Item Code (Source)						NDC:66353-100	
	oute of Administra	tion	TOPICAL				
I	Jute of Automistic	uon					
A	ctive Ingredient	/Active Moi	ety				
Ingredient Name Basis of Strer							Strength
Pe	trolatum (UNII: 4T6)	H12BN9U) (Petro		Pe tro la tum		41 g in 100 g	
In	active Ingredie	nts					
			Ingredient Name			Strength	
Mi	neral Oil (UNII: T5L	.8 T28 FGP)					
Ce	resin (UNII: Q1LS2U	JO3A)					
LA	NOLIN ALCOHOL	<b>S</b> (UNII: 884C3F	A9HE)				
Pa	nthenol (UNII: WV9	CM0067Z)					
Gl	ycerin (UNII: PDC6A	3C0OX)					
LE	EVOMENOL (UNII: 2	4WE03BX2T)					
Pa	ackaging						
#	Item Code		Package Description	Marke	ting Start Date	Mark	eting End Date
1	NDC:66353-100-10	2 in 1 CARTON		11/21/20	11		
		10 g in 1 TUBE	; Type 0: Not a Combination Prod	uct			
1			; Type 0: Not a Combination Prod		11		

Marketing Information									
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date						
OTC monograph final	part347	11/21/2011							

# Labeler - Delta Pharma Inc. (200161730)

Establishment										
Name	Address	ID/FEI	<b>Business Operations</b>							
Delta Pharma Inc.		200161730	manufacture(66353-100)							

Revised: 12/2018

Delta Pharma Inc.