#### VICHY LABORATOIRES NORMADERM BHA EXFOLIATING ACNE TREATMENTsalicylic acid lotion COSMETIQUE ACTIVE PRODUCTION

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## **Drug Facts**

### **Active ingredient**

Salicylic acid 1%

### Purpose

Acne treatment

#### Uses

- for the treatment of acne
- clears acne blemishes and allows skin to heal
- helps prevent new acne blemishes

### Warnings

#### For external use only

## Flammable until dry.

Keep away from flames and heat.

## When using this product

skin irritation and dryness is more likely to occur if you use another topical acne medication at the same time. If irritation occurs, only use one topical acne medication at a time.

## Keep out of reach of children.

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

#### Directions

- cleanse the skin thoroughly before applying this product
- cover the entire affected area with a thin layer one to three times daily
- because too much drying of the skin may occur, start with one application daily, then gradually increase to two or three times daily if needed or as directed by a doctor
- if bothersome dryness or peeling occurs, reduce application to once a day or every other day

## Inactive ingredients

water, alcohol denat., hydroxyethylpiperazine ethane sulfonic acid, glycolic acid, glycerin, propanediol, sodium hydroxide, PEG/PPG/polybutylene glycol-8/5/3 glycerin, sodium polyacrylate, adenosine, ammonium polyacryloyldimethyl taurate, ascorbyl glucoside, vitreoscilla ferment, trisodium ethylenediamine disuccinate, xanthan gum, panthenol, menthol

## **Questions or comments?**

Call toll free **1-877-37-VICHY (84249)** 

Monday – Friday (9 a.m. to 5 p.m. EST)





C1 - Internal use

# VICHY LABORATOIRES NORMADERM BHA EXFOLIATING ACNE

salicylic acid lotion							
Product Information							
Product Type	HUMAN OTC DRUG	Item Code (	Source)	NDC:69	9625-231		
Route of Administration	TOPICAL						
Active Increations / Active	Maiatra						
Active Ingredient/Active	-			-			
	dient Name		Basis of Streng	-	Strength		
SALICYLIC ACID (UNII: O414PZ4L	PZ) (SALICYLIC ACID - UNII:C	)414PZ4LPZ)	SALICYLIC ACID	1	.0 mg in 1 m		
Inactive Ingredients							
	Ingredient Name	9			Strength		
WATER (UNII: 059QF0K00R)							
ALCOHOL (UNII: 3K9958V90M)							
HYDROXYETHYLPIPERAZINE ET	HANE SULFONIC ACID (UN	II: RWW266YE9I)					
GLYCOLIC ACID (UNII: 0WT12SX3	85)						
GLYCERIN (UNII: PDC6A3C0OX)							
PROPANEDIOL (UNII: 5965N8W85T)							
PROPANEDIOL (UNII: 5965N8W85	Γ)		SODIUM HYDROXIDE (UNII: 55X04QC32I)				
SODIUM HYDROXIDE (UNII: 55X0	4QC32I)						
SODIUM HYDROXIDE (UNII: 55X0 SODIUM POLYACRYLATE (2500	4QC32I)						
SODIUM HYDROXIDE (UNII: 55X0 SODIUM POLYACRYLATE (2500 ADENOSINE (UNII: K72T3FS567)	4QC32I) <b>DOO MW)</b> (UNII: 05I15JNI2J)	MPA.S) (UNII: F	01RIY4371)				
SODIUM HYDROXIDE (UNII: 55X0 SODIUM POLYACRYLATE (2500) ADENOSINE (UNII: K72T3FS567) AMMONIUM POLYACRYLOYLDIM	4QC32I) DOO MW) (UNII: 05I15JNI2J) IETHYL TAURATE (55000	MPA.S) (UNII: F	01RIY4371)				
SODIUM HYDROXIDE (UNII: 55X0 SODIUM POLYACRYLATE (2500 ADENOSINE (UNII: K72T3FS567) AMMONIUM POLYACRYLOYLDIM ASCORBYL GLUCOSIDE (UNII: 2V	4QC32I) 000 MW) (UNII: 05I15JNI2J) IETHYL TAURATE (55000 752R0NHXW)	<b>MPA.S)</b> (UNII: F	01RIY4371)				
SODIUM HYDROXIDE (UNII: 55X0 SODIUM POLYACRYLATE (2500 ADENOSINE (UNII: K72T3FS567) AMMONIUM POLYACRYLOYLDIM ASCORBYL GLUCOSIDE (UNII: 2V VITREOSCILLA LYSATE (UNII: 681 TRISODIUM ETHYLENEDIAMINE	4QC32I) DOO MW) (UNII: 05I15JNI2J) IETHYL TAURATE (55000 /52R0NHXW) DV8T89Y2) DISUCCINATE (UNII: YA22H		01RIY4371)				
SODIUM HYDROXIDE (UNII: 55X0 SODIUM POLYACRYLATE (2500 ADENOSINE (UNII: K72T3FS567) AMMONIUM POLYACRYLOYLDIM ASCORBYL GLUCOSIDE (UNII: 2V VITREOSCILLA LYSATE (UNII: 680 TRISODIUM ETHYLENEDIAMINE XANTHAN GUM (UNII: TTV12P4NE	4QC32I) DOO MW) (UNII: 05I15JNI2J) IETHYL TAURATE (55000 52R0NHXW) DV8T89Y2) DISUCCINATE (UNII: YA22H E)		01RIY4371)				
	4QC32I) DOO MW) (UNII: 05I15JNI2J) IETHYL TAURATE (55000 52R0NHXW) DV8T89Y2) DISUCCINATE (UNII: YA22H E)		01RIY4371)				

# Packaging

#	ltem Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69625-231- 01	1 in 1 CARTON	12/12/2022	
1		30 mL in 1 BOTTLE; Type 0: Not a Combination Product		
2	NDC:69625-231- 02	1 in 1 CARTON	12/12/2022	06/01/2023
2		3 mL in 1 TUBE; Type 0: Not a Combination Product		
3	NDC:69625-231- 03	1 in 1 CARTON	12/12/2022	
3		5 mL 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 Drug	M006	12/12/2022		

Labeler - COSMETIQUE ACTIVE PRODUCTION (282658798)

Establishment				
Name	Address	ID/FEI	<b>Business Operations</b>	
COSMETIQUE ACTIVE PRODUCTION		282658798	manufacture(69625-231)	

Establishment				
Name	Address	ID/FEI	<b>Business Operations</b>	
Interspray		364829903	pack(69625-231)	

Revised: 12/2023

COSMETIQUE ACTIVE PRODUCTION