

**VICHY LABORATOIRES NORMADERM PHYTOACTION DAILY DEEP CLEANSING-
salicylic acid gel**

L'Oreal USA Products 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.

Drug Facts

Active ingredient

Salicylic acid...0.5%

Purpose

Acne treatment

Use

for the treatment of acne

Warnings

For external use only

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.

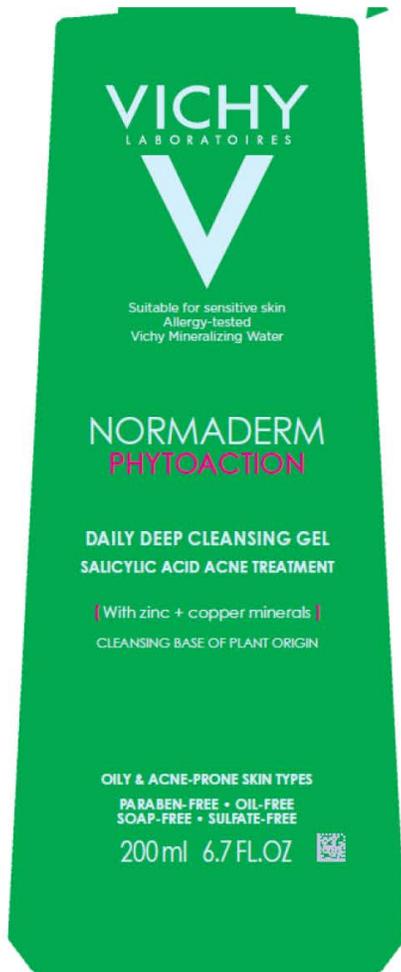
If swallowed, get medical help or contact a Poison Control Center right away.

Directions

- use twice daily
- wet face, then work product into a lather
- massage onto face avoiding the eyes
- rinse well

Inactive ingredients

water, coco-betaine, propanediol, PEG-120 methyl glyucose dioleate, sodium chloride, sodium cocoyl glycinate, dipropylene glycol, zinc gluconate, bifida ferment lysate, sodium hydroxide, sodium benzoate, phenoxyethanol, copper gluconate, caprylyl glycol, tetrasodium glutamate diacetate, fragrance



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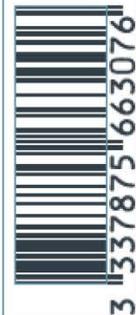
Questions or comments?

Call toll free
1-877-378-4249
 Monday – Friday
 (9 a.m. to 5 p.m. EST)

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200 ml/6.7 FL.OZ

Distribution reserved to agreed VICHY suppliers.
 Vichy LLC, New York, NY 10001
 VICHY CAI/CAF 03 Vichy France
 TSA 75000 93584 ST. QUIEN CEDEX FR
 Made In France www.vichyusa.com



LOT

EXP

C1 - Internal use

VICHY LABORATOIRES NORMADERM PHYTOACTION DAILY DEEP

CLEANSING

salicylic acid gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:49967-307
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
Salicylic Acid (UNII: O414PZ4LPZ) (Salicylic Acid - UNII:O414PZ4LPZ)	Salicylic Acid	5 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0K00R)	
COCO-BETAINE (UNII: 03DH2IZ3FY)	
PROPANEDIOL (UNII: 5965N8W85T)	
PEG-120 METHYL GLUCOSE DIOLEATE (UNII: YM0K64F20V)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
SODIUM COCOYL GLYCINATE (UNII: XLU9KH03XM)	
DIPROPYLENE GLYCOL (UNII: E107L85C40)	
ZINC GLUCONATE (UNII: U6WSN5SQ1Z)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
COPPER GLUCONATE (UNII: RV823G6G67)	
CAPRYLYL GLYCOL (UNII: 00YIU5438U)	
TETRASODIUM GLUTAMATE DIACETATE (UNII: 5EHL50I4MY)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:49967-307-01	200 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/15/2020	
2	NDC:49967-307-02	7 mL in 1 PACKET; Type 0: Not a Combination Product	04/29/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part333D	04/15/2020	

Labeler - L'Oreal USA Products Inc (002136794)

Establishment

Name	Address	ID/FEI	Business Operations
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Revised: 4/2020

L'Oreal USA Products Inc