LA ROCHE POSAY LABORATORIE DERMATOLOGIQUE EFFACLAR MEDICATED CLEANSER- salicylic acid gel
L'Oreal USA Products Inc

Drug Facts

Active ingredient

Salicylic acid 2%

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

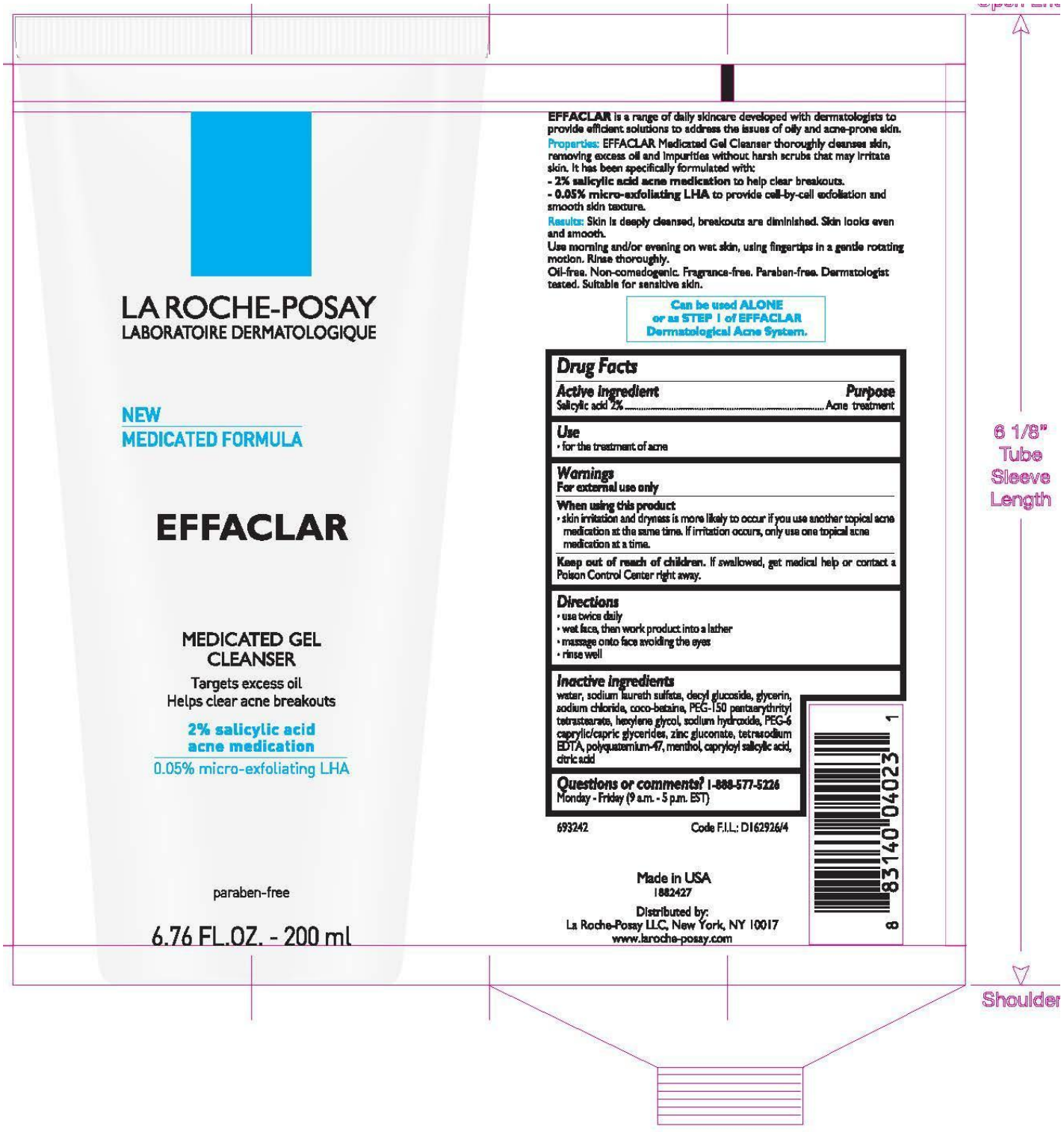
Inactive ingredients

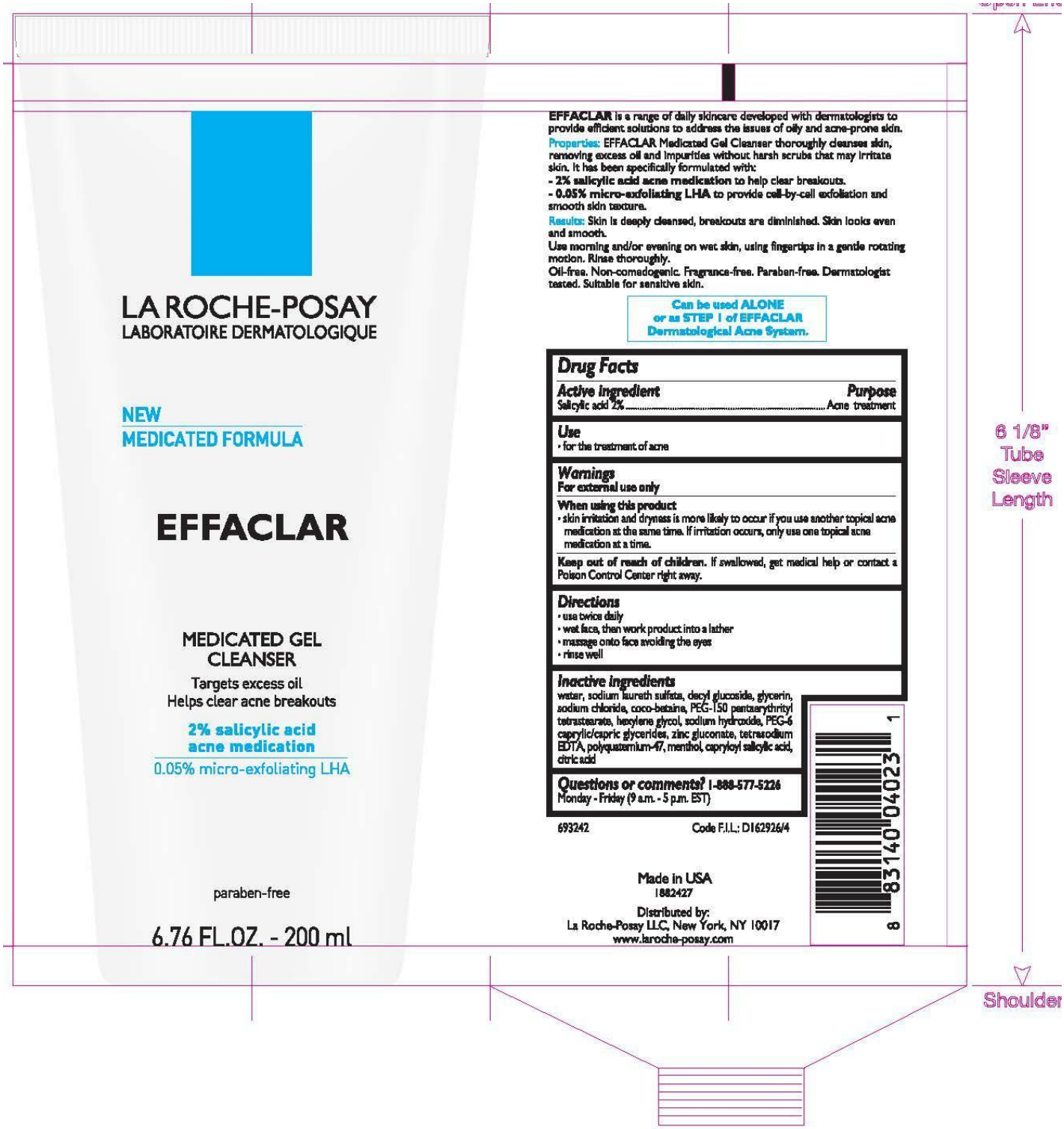
water, sodium laureth sulfate, decyl glucoside, glycerin, sodium chloride, coco-betaine, PEG-150 pentaerythrityl tetrastearate, hexylene glycol, sodium hydroxide, PEG-6 caprylic/capric glycerides, zinc gluconate, tetrasodium EDTA, polyquaternium-47, menthol, capryloyl salicylic acid, citric acid

Questions or comments?

1-888-577-5226

Monday - Friday (9 a.m. - 5 p.m. EST)





LA ROCHE POSAY LABORATORIE DERMATOLOGIQUE EFFACLAR MEDICATED CLEANSER

salicylic acid gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:49967-023

Route of Administration	TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
SALICYLIC ACID (UNII: O414PZ4LPZ) (SALICYLIC ACID - UNII:O414PZ4LPZ)	SALICYLIC ACID	20 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
SODIUM LAURETH SULFATE (UNII: BPV390UAP0)	
DECYL GLUCOSIDE (UNII: Z17H97EA6Y)	
GLYCERIN (UNII: PDC6A3C0OX)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
COCO-BETAINE (UNII: 03DH2IZ3FY)	
PEG-150 PENTAERYTHRITYL TETRASTEARATE (UNII: 8L4OOQ76AM)	
HEXYLENE GLYCOL (UNII: KEH0A3F75J)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
PEG-6 CAPRYLIC/CAPRIC GLYCERIDES (UNII: GO50W2HWO8)	
ZINC GLUCONATE (UNII: U6WSN5SQ1Z)	
MENTHOL (UNII: L7T10EIP3A)	
CAPRYLOYL SALICYLIC ACID (UNII: 5F7PJF6AA4)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:49967-023-01	200 mL in 1 TUBE; Type 0: Not a Combination Product	08/01/2014	
2	NDC:49967-023-02	100 mL in 1 TUBE; Type 0: Not a Combination Product	08/01/2014	
3	NDC:49967-023-03	15 mL in 1 TUBE; Type 0: Not a Combination Product	10/01/2022	
4	NDC:49967-023-04	3 mL in 1 PACKET; Type 0: Not a Combination Product	10/01/2022	
5	NDC:49967-023-05	200 mL in 1 TUBE; Type 0: Not a Combination Product	10/01/2022	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M006	08/01/2014	

Labeler - L'Oreal USA Products Inc (002136794)

Establishment

Name	Address	ID/FEI	Business Operations
L'Oreal USA, Inc.		185931458	manufacture(49967-023) , pack(49967-023)

Revised: 12/2023

L'Oreal USA Products Inc