CLEAR PROOF BLEMISH CONTROL TONER ACNE MEDICATION- salicylic acid liquid
Mary Kay 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.

Clear Proof Blemish Control Toner

Drug Facts

Active ingredient

Salicylic Acid (2% W/W)

Purpose

Acne Medication

Uses

- for the management of acne
- helps prevent new acne pimples

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.
- avoid contact with the eyes

Stop use and ask a doctor

if irritation or sensitivity develops or increases

Keep out of reach of children.

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

Directions

- clean the skin thoroughly before applying this product
- cover the entire affected area with a thin layer one to three times daily
- because excessive 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

alcohol denat., citric acid, diethylhexyl sodium sulfosuccinate, eucalyptus globulus leaf oil, menthol, propylene glycol, simethicone, water

Principal Display Panel - 147 mL bottle

clearproof

blemish control toner

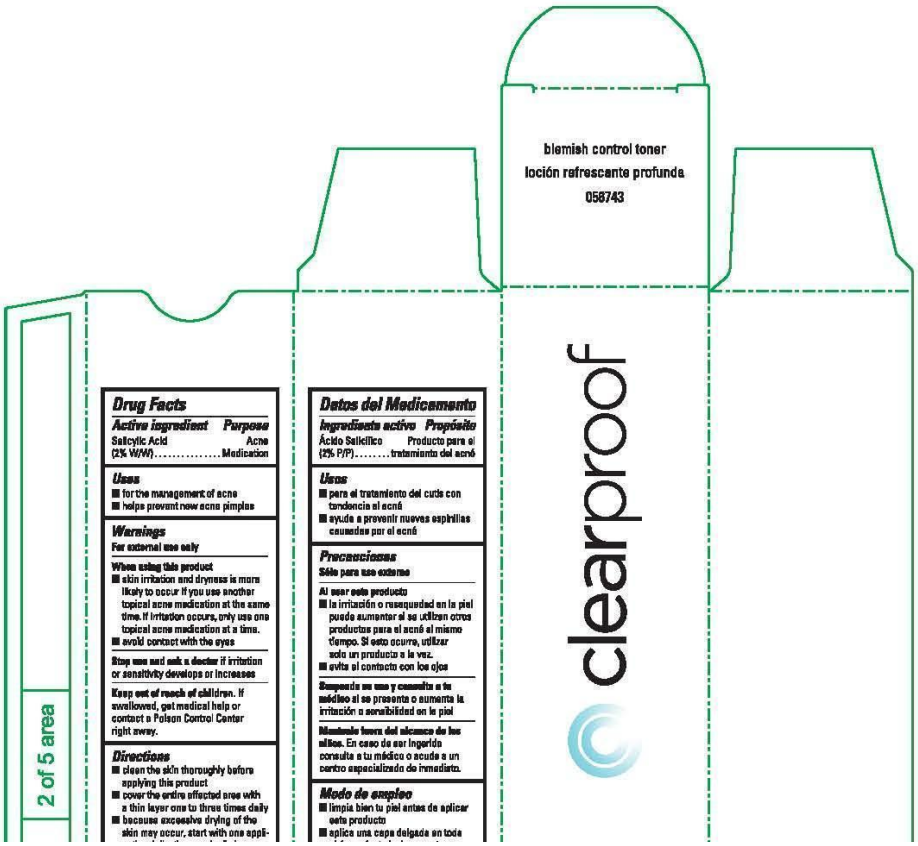
acne medication

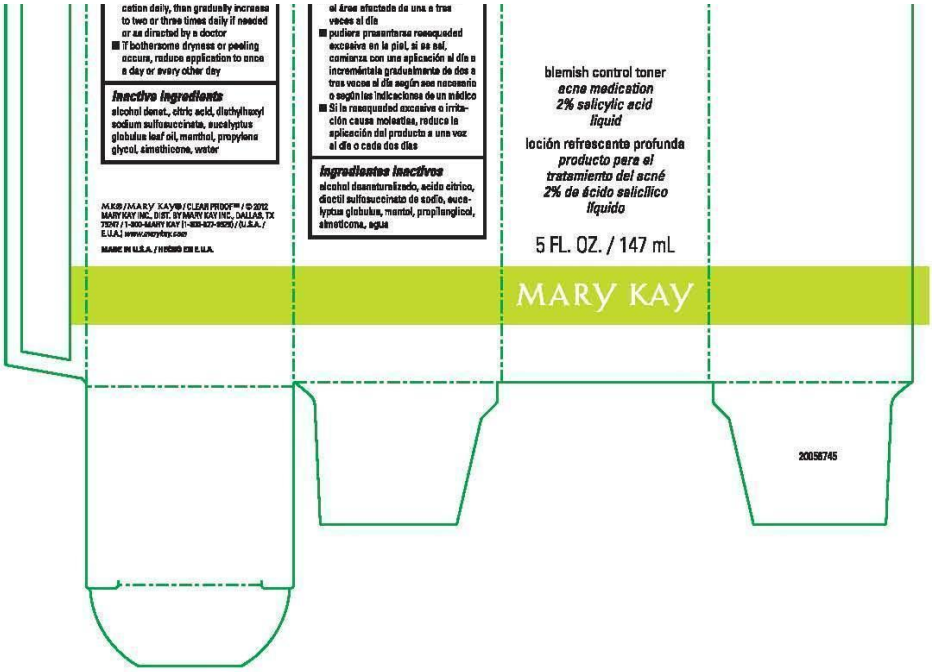
2% salicylic acid

liquid

5 FL. OZ. / 147 mL

Mary Kay





CLEAR PROOF BLEMISH CONTROL TONER ACNE MEDICATION

salicylic acid liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:51531-6743
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
SALICYLIC ACID (UNII: O414PZ4LPZ) (SALICYLIC ACID - UNII:O414PZ4LPZ)	SALICYLIC ACID	2 g in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
ALCOHOL (UNII: 3K9958V90M)	
DENATONIUM BENZOATE (UNII: 4YK5Z54AT2)	
TERT-BUTYL ALCOHOL (UNII: MD83SFE959)	

PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
DOCUSATE SODIUM (UNII: F05Q2T2JA0)	
EUCALYPTUS OIL (UNII: 2R04ONI662)	
MENTHOL (UNII: L7T10EIP3A)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:51531-6743-5	1 in 1 CARTON	08/15/2013	
1		147 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		
2	NDC:51531-6743-9	26 mL in 1 BOTTLE; Type 0: Not a Combination Product	08/15/2013	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	M006	08/15/2013	

Labeler - Mary Kay Inc. (049994452)

Establishment

Name	Address	ID/FEI	Business Operations
Port Jervis Laboratories Inc.		001535103	manufacture(51531-6743)

Establishment

Name	Address	ID/FEI	Business Operations
Mary Kay Inc.		103978839	manufacture(51531-6743)

Revised: 9/2022 Mary Kay Inc.