CLEARCALM NON-DRYING ACNE TREATMENT- salicylic acid gel
Ren Ltd.

Clearcalm Non-Drying Acne Treatment Gel

Drug Facts

Active ingredient

Salicylic Acid 0.5%

Purpose

AcneTreatment

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

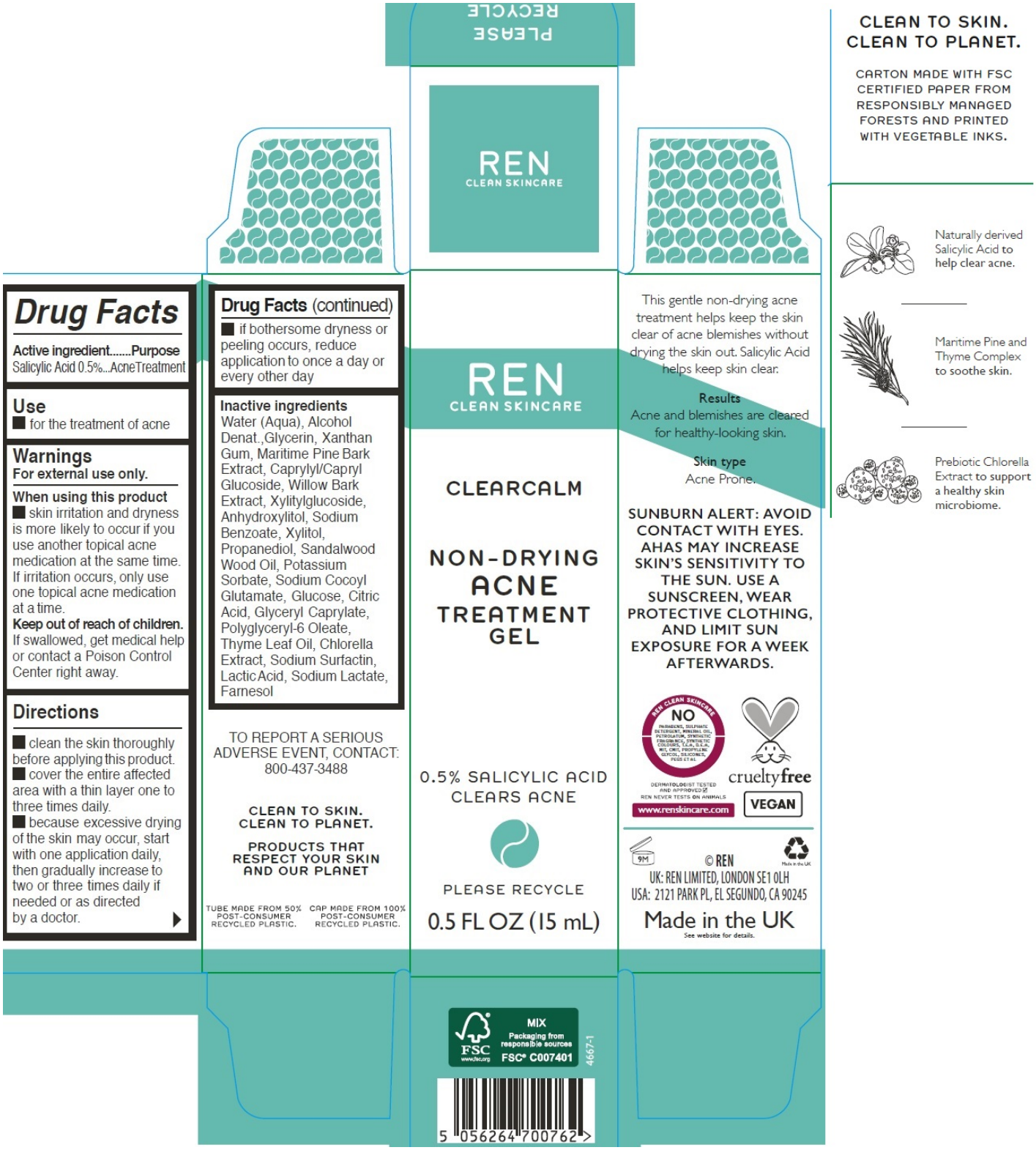
- clean the skin thoroughly before applying this product.
- cover the entire affected area with a thin layer one to three times daily.
- because excessive drying of the skin may occur, start with one application daily, then gradually increase to two or three times daily if needed or as directed by a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

Inactive ingredients

Water (Aqua), Alcohol Denat.,Glycerin, Xanthan Gum, Maritime Pine Bark Extract, Caprylyl/Capryl Glucoside, Willow Bark Extract, Xylitylglucoside, Anhydroxylitol, Sodium Benzoate, Xylitol, Propanediol, Sandalwood Wood Oil, Potassium Sorbate, Sodium Cocoyl Glutamate, Glucose, Citric Acid, Glyceryl Caprylate, Polyglyceryl-6

Oleate, Thyme Leaf Oil, Chlorella Extract, Sodium Surfactin, Lactic Acid, Sodium Lactate, Farnesol

Package Labeling:



CLEARCALM NON-DRYING ACNE TREATMENT
salicylic acid gel

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:51417-0005
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: 059QF0KO0R)	
ALCOHOL (UNII: 3K9958V90M)	
GLYCERIN (UNII: PDC6A3C0OX)	
XANTHAN GUM (UNII: TTV12P4NEE)	
CAPRYLYL/CAPRYL OLIGOGLUCOSIDE (UNII: E00JL9G9K0)	
WILLOW BARK (UNII: S883J9JDYX)	
XYLITYLGLUCOSIDE (UNII: O0IEZ166FB)	
ANHYDROXYLITOL (UNII: 8XWR7NN42F)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
XYLITOL (UNII: VCQ006KQ1E)	
PROPANEDIOL (UNII: 5965N8W85T)	
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)	
SODIUM COCOYL GLUTAMATE (UNII: BMT4RCZ3HG)	
ANHYDROUS DEXTROSE (UNII: 5SL0G7R0OK)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
GLYCERYL MONOCAPRYLATE (UNII: TM2TZD4G4A)	
LACTIC ACID, UNSPECIFIED FORM (UNII: 33X04XA5AT)	
SODIUM LACTATE (UNII: TU7HW0W0QT)	
FARNESOL (UNII: EB41QIU6JL)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:51417-0005-1	1 in 1 BOX	06/01/2019	
1		15 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	06/01/2019	

Labeler - Ren Ltd. (385613831)

Revised: 11/2023 Ren Ltd.