

SUPER SPOT REMOVER ACNE TREATMENT- salicylic acid gel
ORIGINS NATURAL RESOURCES 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.

SUPER SPOT REMOVER™ ACNE TREATMENT

Drug Facts

Active ingredient

Salicylic Acid 1.50%

Purpose

Acne treatment

Uses

- For the treatment of acne

Warnings

For external use only.

Avoid eye area.

Using other topical acne drugs at the same time or right after use of this product may increase dryness or irritation of the skin. If this occurs, only one medication should be used unless directed by a doctor.

Keep out of reach of children. If swallowed, call a doctor or contact a Poison Control Center right away.

Directions

- Cleanse skin.
- Apply a thin layer 1 to 3 times daily or as needed.
- If dryness or peeling occurs, reduce application to every other day.

Inactive ingredients

water\aquaeau • alcohol denat. • acrylates/c10-30 alkyl acrylate crosspolymer • hamamelis virginiana (witch hazel) • eugenia caryophyllus (clove) bud oil¹, organum heracleoticum flower oil¹, eugenol, limonene • cucumis sativus (cucumber) fruit extract • pyrus malus (apple) fruit extract • scutellaria baicalensis root extract • algae extract • poria cocos sclerotium extract • laminaria saccharina extract • lactobacillus ferment •

glycerin • caffeine • butylene glycol • tocopheryl acetate • magnesium ascorbyl phosphate • sodium hydroxide • phenoxyethanol <iln37864>

1 essential oil

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New York, NY 10022

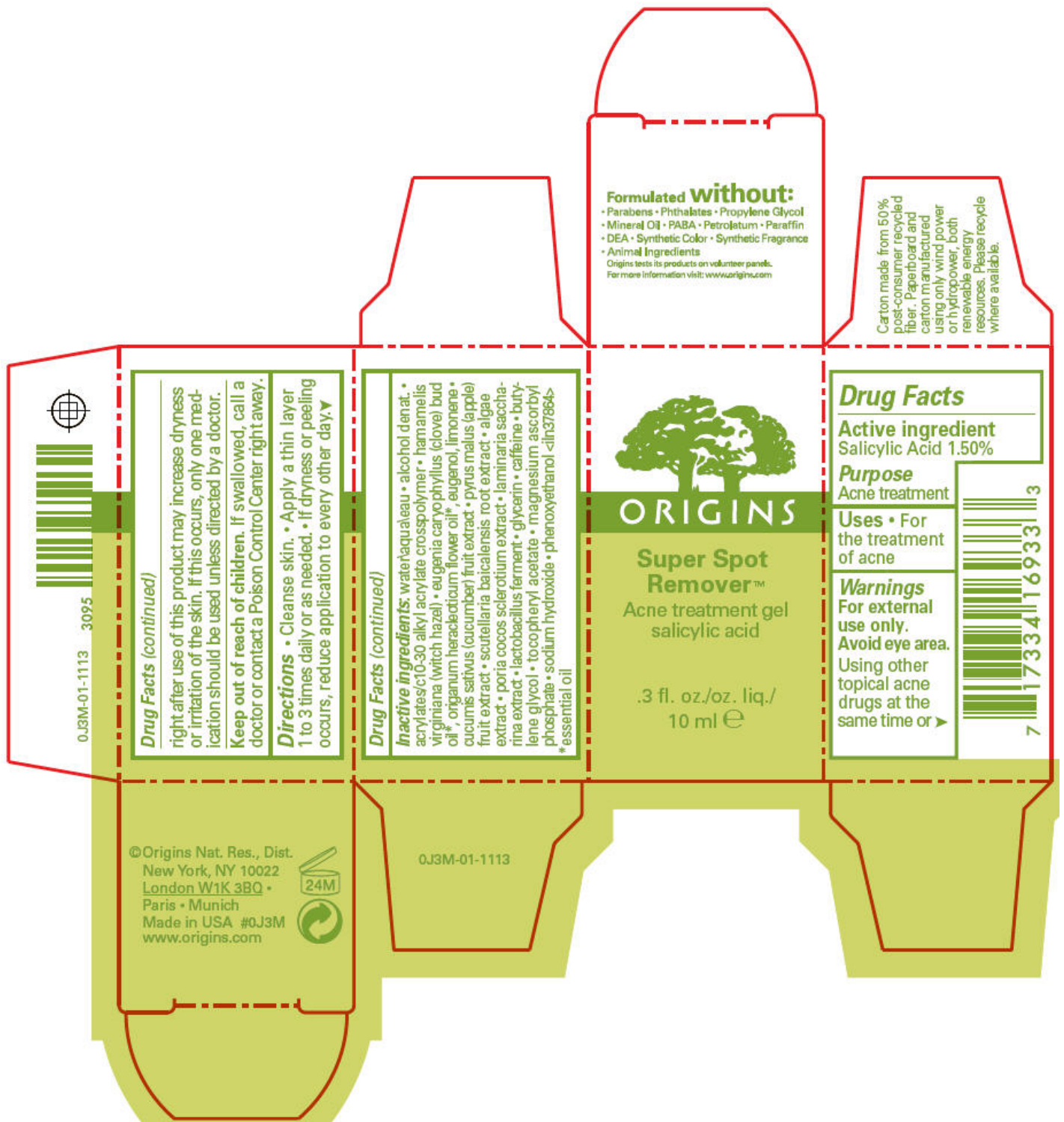
PRINCIPAL DISPLAY PANEL - 10 ml Carton

ORIGINS

**Super Spot
Remover**™

Acne treatment gel
salicylic acid

.3 fl. oz./oz. liq./
10 ml e



SUPER SPOT REMOVER ACNE TREATMENT

salicylic acid gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:59427-005
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
SALICYLIC ACID (UNII: 0414PZ4LPZ) (SALICYLIC ACID - UNII:0414PZ4LPZ)	SALICYLIC ACID	15 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
ORIGANUM VULGARE SUBSP. HIRTUM FLOWER (UNII: 3799SP11NY)	
WATER (UNII: 059QF0KO0R)	
WITCH HAZEL (UNII: 101I4J0U34)	
CLOVE OIL (UNII: 578389D6D0)	
EUGENOL (UNII: 3T8H1794QW)	
CUCUMBER (UNII: YY7C30VXJT)	
APPLE (UNII: B423VGH5S9)	
SCUTELLARIA BAICALENSIS ROOT (UNII: 7J95K7ID2S)	
SACCHARINA LATISSIMA (UNII: 68CMP2MB55)	
GLYCERIN (UNII: PDC6A3C0OX)	
CAFFEINE (UNII: 3G6A5W338E)	
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
MAGNESIUM ASCORBYL PHOSPHATE (UNII: 0R822556M5)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
ALCOHOL (UNII: 3K9958V90M)	
CARBOMER 1342 (UNII: 809Y72KV36)	
FU LING (UNII: XH37TWY5O4)	
LIMOSILACTOBACILLUS FERMENTUM (UNII: 2C1F12C6AP)	
PORPHYRIDIUM PURPUREUM (UNII: K2P8K2558N)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:59427-005-01	1 in 1 CARTON	11/24/2020	
1		10 mL in 1 CONTAINER; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	M006	04/01/2012	

Labeler - ORIGINS NATURAL RESOURCES INC. (611716283)**Registrant** - Estee Lauder Companies Inc. (790802086)

Establishment			
Name	Address	ID/FEI	Business Operations
Bentley Laboratories, LLC		068351753	manufacture(59427-005)

Establishment			
Name	Address	ID/FEI	Business Operations
The Estee Lauder Inc		802599436	manufacture(59427-005)

Revised: 9/2023

ORIGINS NATURAL RESOURCES INC.