

**SALT AND STONE SUNSCREEN LIP BALM SPF 30- zinc oxide stick**  
**Salt and Stone LLC**

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**Salt & Stone Sunscreen Lip Balm SPF 30, 0.15 Oz.(4.3 g)**

**Drug Facts**

**Active Ingredients**

Zinc Oxide (Non-Nano) 8 %

**Purpose**

Sunscreen

**Uses**

• Helps prevent sunburn. • If used as directed with other sun protection measures (see directions), decreases the risk of skin cancer and early skin aging caused by the sun.

**Warnings**

For external use only.

Do not use on damaged or broken skin

When using this product keep out of eyes. Rinse with water to remove

Stop use and ask a doctor if rash occurs.

**Keep out of reach of children.**

If product is swallowed, get medical help or contact a Poison Control Center immediately.

**Directions**

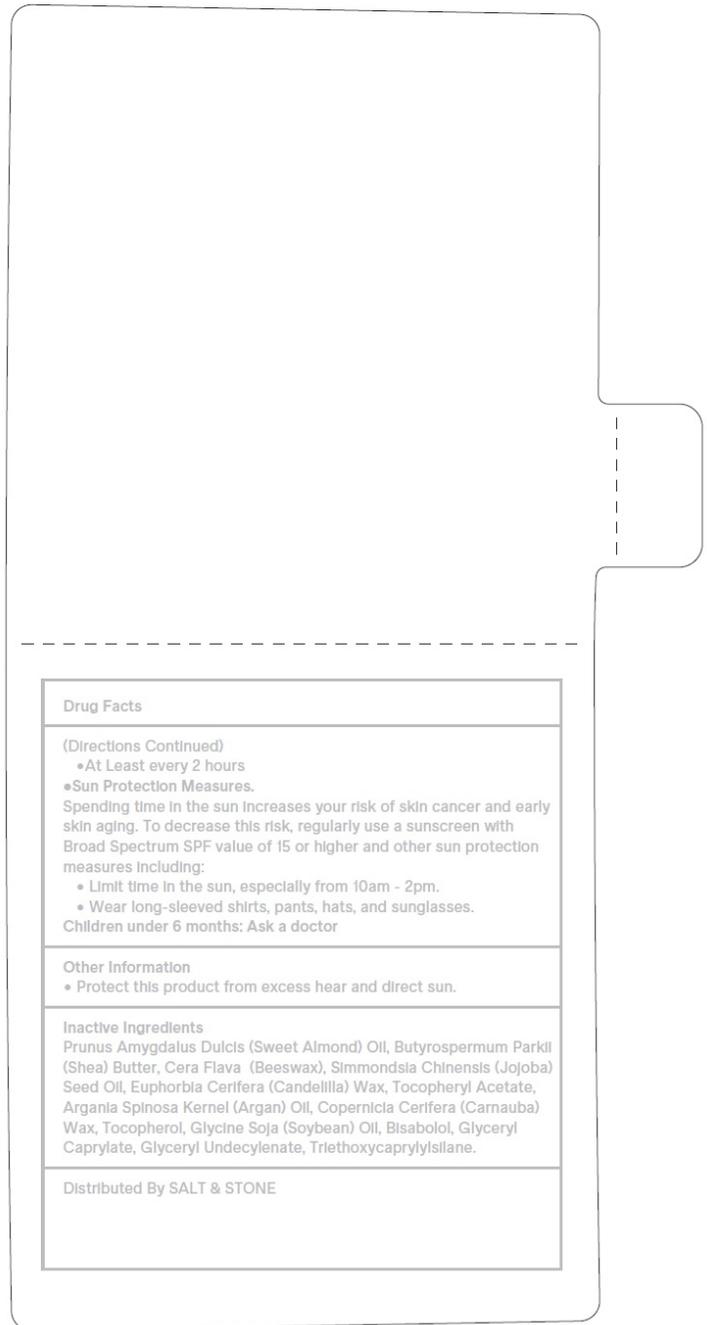
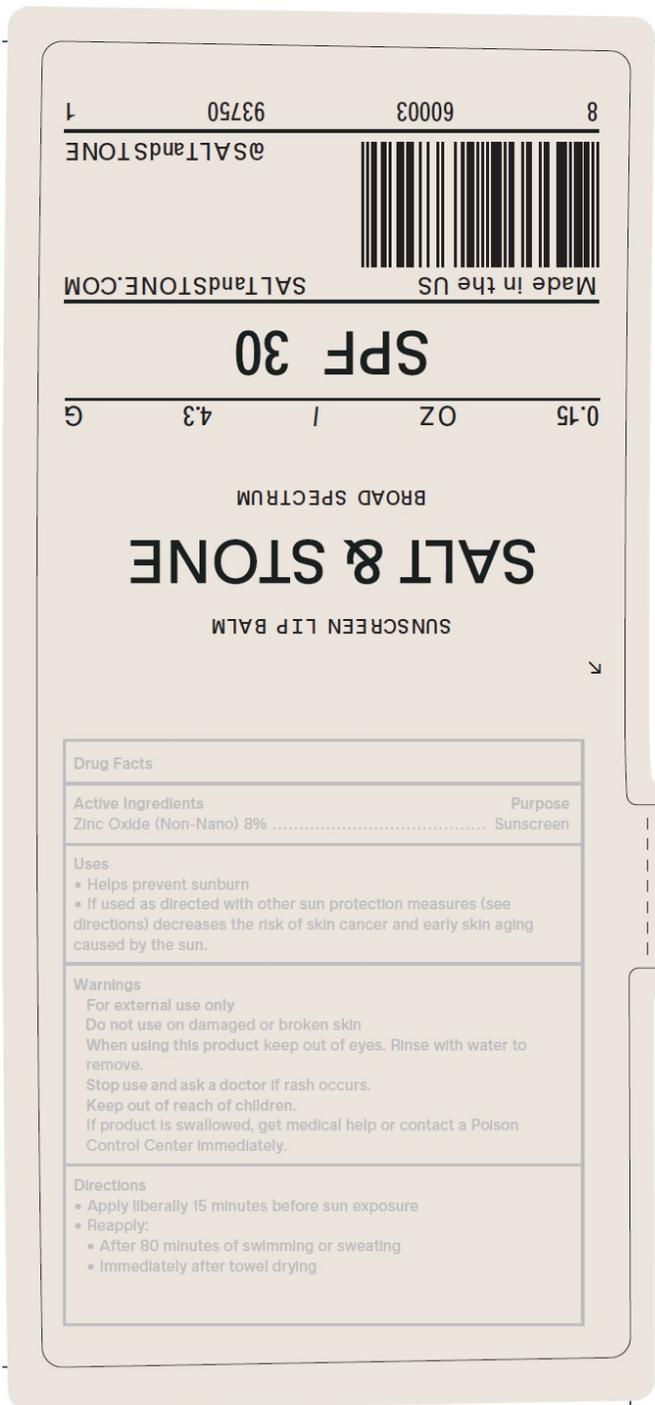
- Apply liberally 15 minutes before sun exposure
- Reapply:
  - After 80 minutes of swimming or sweating
  - Immediately after towel drying
  - At Least every 2 hours
- Spending time in the sun increases your risk of skin cancer and early skin aging. To decrease this risk, regularly use a sunscreen with Broad Spectrum SPF value of 15 or higher and other sun protection measures including:
  - Limit time in the sun, especially from 10am - 2pm.
  - Wear long-sleeved shirts, pants, hats, and sunglasses. **Sun Protection Measures.**
- **Children under 6 months: Ask a doctor**

## **Other Information**

- Protect this product from excess heat and direct sun.

## **Inactive Ingredients**

Prunus Amygdalus Dulcis (Sweet Almond) Oil, Butyrospermum Parkii (Shea) Butter, Cera Flava (Beeswax), Simmondsia Chinensis (Jojoba) Seed Oil, Euphorbia Cerifera (Candelilla) Wax, Tocopheryl Acetate, Argania Spinosa Kernel (Argan) Oil, Copernicia Cerifera (Carnauba) Wax, Tocopherol, Glycine Soja (Soybean) Oil, Bisabolol, GlycerylCaprylate, Glyceryl Undecylenate, Triethoxycaprylylsilane



SALT AND STONE SUNSCREEN LIP BALM SPF 30			
zinc oxide stick			
Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:71585-139
Route of Administration	TOPICAL		
Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	

<b>ZINC OXIDE</b> (UNII: SOI2LOH54Z) (ZINC CATION - UNII:13S1S8SF37)	ZINC CATION	0.08 g in 1 g
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### Inactive Ingredients

Ingredient Name	Strength
<b>ALMOND OIL</b> (UNII: 66YXD4DKO9)	
<b>SHEA BUTTER</b> (UNII: K49155WL9Y)	
<b>CANDELILLA WAX</b> (UNII: WL0328HX19)	
<b>.ALPHA.-TOCOPHEROL ACETATE</b> (UNII: 9E8X80D2L0)	
<b>CARNAUBA WAX</b> (UNII: R12CBM0EIZ)	
<b>TOCOPHEROL</b> (UNII: ROZB2556P8)	
<b>SOYBEAN OIL</b> (UNII: 241ATL177A)	
<b>LEVOMENOL</b> (UNII: 24WE03BX2T)	
<b>GLYCERYL MONOCAPRYLATE</b> (UNII: TM2TZD4G4A)	
<b>GLYCERYL 1-UNDECYLENATE</b> (UNII: B68LJT9544)	
<b>TRIETHOXYCAPRYLYLSILANE</b> (UNII: LDC331P08E)	

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:71585-139-00	4.3 g in 1 APPLICATOR; Type 0: Not a Combination Product	11/23/2020	

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M020	11/23/2020	

**Labeler** - Salt and Stone LLC (080683697)

Revised: 1/2025

Salt and Stone LLC