

**CREAM- zinc oxide, titanium dioxide lotion**  
**Oxygen Development LLC**

*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.*

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**SUGAR RUSH SKIN TREAT PORELESS TINTED MOISTURIZER BROAD SPECTRUM  
SPF 20 SUNSCREEN**

**Active Ingredients**

Titanium Dioxide 4.55%

Zinc Oxide 3.5%

**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 aging caused by the sun.

**Warnings**

For external use only. Stop use and ask a doctor if rash occurs. Do not use on damage or broken skin.

**When using this product**

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

**Keep out of the reach of children.**

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

**Directions**

Shake before use. Apply liberally 15 minutes before sun exposure. Use a water resistant sunscreen if swimming or sweating. Reapply at least every 2 hours. Children under 6 months: Ask a doctor. Sun protection measures. Spending time in the sun increases your risk of skin cancer and early skin aging. To decrease the risk, regularly use a sunscreen with a Broad Spectrum SPF value of 15 or higher and other sun protection measures including:

- Limit time in the sun, especially from 10 a.m. - 2 p.m.
- Wear long-sleeved shirts, pants, hats, and sunglasses

### Other information

Protect the product in this container from excessive heat and direct sunlight.

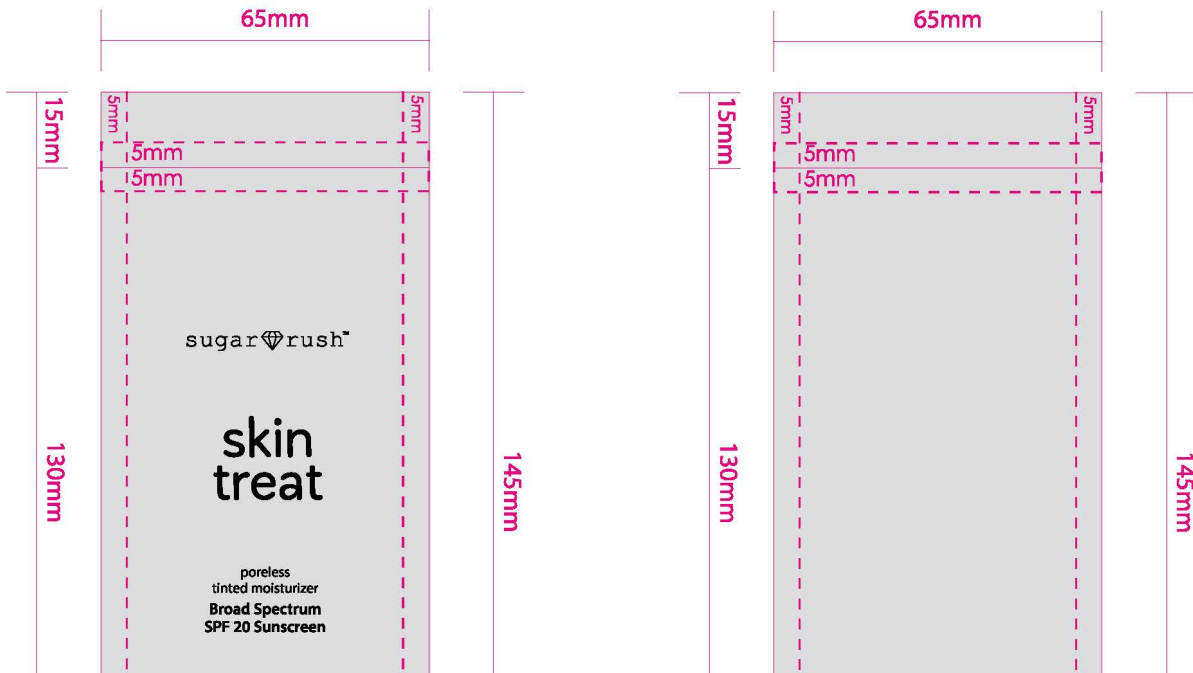
You may report a serious adverse reaction to: tarte c/o Report Reaction, LLC, P.O. Box 22, Plainsboro, NJ 08536-0222

### Inactive Ingredients

Cyclopentasiloxane, Isododecane, Polysilicone-11, Polymethylsilsesquioxane, Hexyl Laurate, PEG-10 Dimethicone, Polyglyceryl-4 Isostearate, Stearic Acid, Cetyl PEG/PPG-10/1 Dimethicone, Alumina, Triethoxycaprylylsilane, Dipalmitoyl Hydroxyproline, Diamond Powder

### Package Label - Principal Display Panel

sugar  rush	
SUB-BRAND: SUGAR RUSH	FILE NAME: 01541CO-2 GG SKIN TREAT SPF
COMPONENT: ZIP BAG	SHADE: DEEP
DATE OPENED: 9/1/20	DATE REVISED:
REASON FOR CHANGE:	
<b>COLORS</b>	
<input checked="" type="checkbox"/> 9120C <input type="checkbox"/> WHITE <input type="checkbox"/> BLACK <input type="checkbox"/> CLEAR	
FINISH: GLOSSY	







F: front panel width      B: back panel width

CW: circumferential width

L: tube length (from shoulder to bottom)

Eyemark (2×4 mm): mark for end sealing

Upper edge(7mm): space for end sealing

Bottom edge(5mm): non-printing space

Dia 13xL50  
shadow area:

F/B  
20mm

printing area

CW  
40mm

L(changeable)  
50mm

## CREAM

zinc oxide, titanium dioxide lotion

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:61354-066
<b>Route of Administration</b>	TOPICAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>ZINC OXIDE</b> (UNII: SOI2LOH54Z) (ZINC OXIDE - UNII:SOI2LOH54Z)	ZINC OXIDE	3.49 mg in 100 mg
<b>TITANIUM DIOXIDE</b> (UNII: 15FIX9V2JP) (TITANIUM DIOXIDE - UNII:15FIX9V2JP)	TITANIUM DIOXIDE	4.54 mg in 100 mg

### Inactive Ingredients

Ingredient Name	Strength
<b>BEC 10 DIMETHICONE (600 CST)</b> (UNII: 8DD7V16V40)	1.65 mg

<b>PEG-10 DIMETHICONE (800 CST)</b> (UNII: 8FR7V13VMU)	in 100 mg
<b>DIMETHICONE CROSSPOLYMER (450000 MPAS AT 12% IN CYCLOPENTASILOXANE)</b> (UNII: UF7620L1W6)	55.74 mg in 100 mg
<b>ISODODECANE</b> (UNII: A8289P68Y2)	10.69 mg in 100 mg
<b>POLYMETHYLSILSESQUIOXANE (4.5 MICRONS)</b> (UNII: 59Z907ZB69)	3.19 mg in 100 mg
<b>DIMETHICONE/VINYL DIMETHICONE CROSSPOLYMER (SOFT PARTICLE)</b> (UNII: 9E4CO0W6C5)	4.68 mg in 100 mg
<b>HEXYL LAURATE</b> (UNII: 4CG9F9W01Q)	1.76 mg in 100 mg
<b>POLYGLYCERYL-4 ISOSTEARATE</b> (UNII: 820DPX33S7)	1.5 mg in 100 mg

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:61354-066-01	1 in 1 CARTON	05/06/2022	
1		100 mg 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 final	part352	05/06/2022	

**Labeler** - Oxygen Development LLC (137098492)

### Establishment

Name	Address	ID/FEI	Business Operations
Oxygen Development LLC		137098492	manufacture(61354-066)

Revised: 5/2022

Oxygen Development LLC