

CREAM- titanium dioxide, zinc oxide 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.

TAN DEEP- SKIN TREAT BLURRING SKIN TINT 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 skin aging caused by the sun.

Warnings

For external use only

Stop use and ask doctor if rash occurs.

Do not use on damaged or broken skin.

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 of 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 our risk of skin cancer and early skin aging. To decrease this 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.- 2p.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, Mica, 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, Iron Oxides.

Primary Package (Inner)



F: front panel width B: back panel width


CW: circumferential width

L: length (Circumferential width x 2)

L: tube length (from shoulder to bottom)
 Eyemark (3 × 6mm): mark for end sealing
 Upper edge(7mm): space for end sealing
 Bottom edge(5mm): non-printing space


Dia 25 × L110	F/B 39.25mm	CW 78.5mm	L(changeable) 110mm
shadow area:	printing area		

Attention of artwork: Use CorelDraw, Freehand or Illustrator, and do not compress or change the file's type, then outline the file and send us.

sugar  rush	
SUB-BRAND: SUGAR RUSH	FILE NAME: SKIN TREAT
COMPONENT: PRIMARY	SHADE: SPF
DATE OPENED: 5/4/18	DATE REVISED: 10/19/18
REASON FOR CHANGE: PID REVISED	
COLORS	
■ BLACK	■ 182C
■ 225C	■ 304C

Secondary Package (Outer)

tan-deep

sugar  rush
tarte™

*skin
treat*

@sugarrush
sugarrushbeauty.com

**Drug Facts
(continued)**

**Inactive
Ingredients**

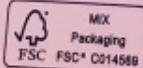
Cyclopentasiloxane,
isododecane,
polysilicone-11, mica,
polymethylsilsesqui-
oxane, hexyl laurate,
PEG-10 dimethicone,
polyglyceryl-4
isostearate, stearic
acid, cetyl
PEG/PPG-10/1
dimethicone, alumina,
triethoxycaprylylsi-
lane, dipalmitoyl
hydroxyproline,
diamond powder, iron
oxides.
OX100FZ000101.

LOT OH01
EXP: 10/21

net 30 mL / 1 Fl. Oz. Made
in USA. Dist. by tarte,
Inc. NY 10018. All rights
reserved. Biorius Sprl, Rue
de la Croye 10A, B-7170
Manage, BE.



crueltyfree
andvegan



blurring skin tint
Broad Spectrum
SPF 20 Sunscreen

Drug Facts

**Active
Ingredients Purpose**

Titanium Dioxide 4.55%.....Sunscreen
Zinc Oxide 3.50%.....Sunscreen

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FG05242-01



CREAM

titanium dioxide, zinc oxide lotion

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:61354-051
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC OXIDE - UNII:SOI2LOH54Z)	ZINC OXIDE	3.5 mg in 100 mg
TITANIUM DIOXIDE (UNII: 15FIX9V2JP) (TITANIUM DIOXIDE - UNII:15FIX9V2JP)	TITANIUM DIOXIDE	4.55 mg in 100 mg

Inactive Ingredients

Ingredient Name	Strength
POLYMETHYLSILSESQUIOXANE (11 MICRONS) (UNII: Z570VEV8XK)	
HEXYL LAURATE (UNII: 4CG9F9W01Q)	
ISODODECANE (UNII: A8289P68Y2)	
MICA (UNII: V8A1AW0880)	
POLYGLYCERYL-4 ISOSTEARATE (UNII: 820DPX33S7)	
PEG-10 DIMETHICONE (600 CST) (UNII: 8PR7V1SVM0)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
DIMETHICONE/VINYL DIMETHICONE CROSSPOLYMER (SOFT PARTICLE) (UNII: 9E4CO0W6C5)	
CYCLOMETHICONE 5 (UNII: 0THT5PCI0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:61354-051-16	1 in 1 CARTON	02/23/2021	
1		30 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	02/23/2021	

Labeler - OXYGEN DEVELOPMENT, LLC (137098492)

Establishment

Name	Address	ID/FEI	Business Operations
OXYGEN DEVELOPMENT, LLC		137098492	manufacture(61354-051)

Revised: 2/2023

OXYGEN DEVELOPMENT, LLC