# CK ONE WATERFRESH FACE MAKEUP SPF 15 200 FAIR- titanium dioxide cream Coty US 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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### **Drug Facts**

Octinoxate 2.5%

Titanium dioxide 5.0%

Sunscreen

Keep out of reach children. If swallowed, get medical help or contact a Poison Control Center right away.

Helps prevent sunburn

**Skin cancer/Skin Aging Alert**: Spending time int he sun increases your risk of skin cancer and early skin aging. This product has been shown only to prevent sunburn, **not** skin cancer or early skin aging.

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.

#### For sunscreen 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

aqua/water/eau, hydogentated polyisobutene, isohexadecane, dextrin palmitate, butylene glycol, hydrogenated lecithin, alcohol, triethylhexanoin, quaternium-90, bentonite, pentylene glycol, glycerin, peg-15/lauryl dimethicone crosspolymer, stearoyl inulin, phenoxyethanol, isopropl titanium triisostearate, octyldodectl pca, lysine, isododecane, dipropylene glycol, magnesium chloride, sodium chloride, sodium citrate, may contain: iron oxides, titanium dioxide, mica.



ck one airlight pressed powder SPF 15









### CK ONE WATERFRESH FACE MAKEUP SPF 15 200 FAIR

titanium dioxide cream

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:66184-445
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
Titanium Dioxide (UNII: 15FIX9 V2JP) (Titanium Dioxide - UNII:15FIX9 V2JP)	Titanium Dioxide	4.95 mL in 9.9 g	
Octinoxate (UNII: 4Y5P7MUD51) (Octinoxate - UNII:4Y5P7MUD51)	Octinoxate	2.4 mL in 9.9 g	

Inactive Ingredients		
Ingredient Name	Strength	
Water (UNII: 059QF0KO0R)		
Isohexadecane (UNII: 918X1OUF1E)		

Butylene Glycol (UNII: 3XUS85K0RA)			
Hydrogenated Soybean Lecithin (UNII: H1109Z9J4N)			
Alcohol (UNII: 3K9958V90M)			
Triethylhexanoin (UNII: 7K3W1BIU6K)			
Pentylene Glycol (UNII: 50C1307PZG)			
Glycerin (UNII: PDC6A3C0OX)			
Lysine (UNII: K3Z4F929H6)			
Isododecane (UNII: A8289P68Y2)			
Dipropylene Glycol (UNII: E107L85C40)			
Magnesium Chloride (UNII: 02F3473H9O)			
Sodium Chloride (UNII: 451W47IQ8X)			
Potassium Chloride (UNII: 660 YQ 98 I10)			
Sodium Citrate (UNII: 1Q73Q2JULR)			
Mica (UNII: V8A1AW0880)			

Packaging				
#	tem Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:66184-445-01	9.9 g in 1 CARTRIDGE		
2	NDC:66184-445-02	9.9 g in 1 BOX		

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph not final	part352	0 1/23/20 12		

# Labeler - Coty US LLC (789573201)

## Registrant - Coty Inc. (958662223)

Establishment				
Name	Address	ID/FEI	Business Operations	
Intercos Europe S.p.A.		438961310	manufacture	

Revised: 1/2012 Coty US LLC