

**MATCH PERFECTION SPF 18 NUDE 202- octinoxate titanium dioxide liquid  
Rimmel 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.*

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**Drug Facts**

Octinoxate 4%

Titanium Dioxide 4%

Sunscreen

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

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.

**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.

**Directions.** 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.

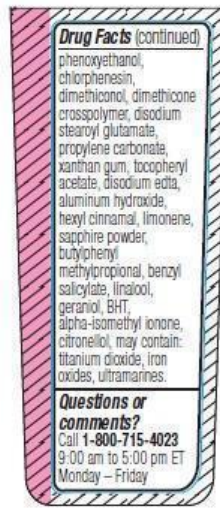
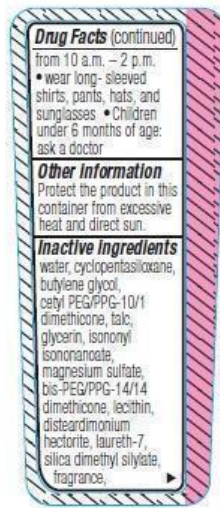
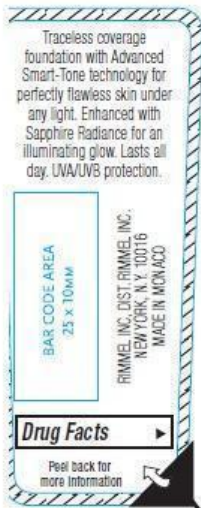
**Sun Protection Measures.** Spending time in the sun increases your risk of skin cancer and early skin aging. To decrease this risk, regularly use a sunscreen with a Broad Spectrum value of 15 SPF or higher and other sun protection measures including:

- limit time in the sun, especially from 10 a.m.-2 p.m.
- wear long-sleeved shirts, pants, hats and sunglasses
- children under 6 months of age: ask a doctor

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

water, cyclopentasiloxane, butylene glycol, cetyl PEG/PPG-10/1 dimethicone, talc, glycerin, isononyl isononanoate, magnesium sulfate, bis-PEG/PPG-14/14 dimethicone, lecithin, disteardimonium hectorite, laureth-7, silica dimethyl silylate, fragrance, phenoxyethanol, chlorphenesin, dimethiconol, dimethicone, crosspolymer, disodium stearyl glutamate, propylene carbonate, xanthan gum, tocopheryl acetate, disodium edta, aluminum hydroxide, hexyl cinnamal, limonene, sapphire powder, butylphenyl methylpropional, benzyl salicylate, linalool, geraniol, BHT, alpha-isomethyl ionone, citronellol, may contain: titanium dioxide, iron oxides, ultramarines.

**Front Label**



## MATCH PERFECTION SPF 18 NUDE 202

octinoxate titanium dioxide liquid

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:76485-1031
<b>Route of Administration</b>	TOPICAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>OCTINOXATE</b> (UNII: 4Y5P7MUD51) (OCTINOXATE - UNII:4Y5P7MUD51)	OCTINOXATE	1.2 mL in 30 mL
<b>TITANIUM DIOXIDE</b> (UNII: 15FIX9V2JP) (TITANIUM DIOXIDE - UNII:15FIX9V2JP)	TITANIUM DIOXIDE	1.5 mL in 30 mL

## Inactive Ingredients

Ingredient Name	Strength
Water (UNII: 059QF0KO0R)	
CYCLOMETHICONE 5 (UNII: 0THT5PCI0R)	
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	
CETYL PEG/PPG-10/1 DIMETHICONE (HLB 1.5) (UNII: V2W71V8T0X)	
TALC (UNII: 7SEV7J4R1U)	
GLYCERIN (UNII: PDC6A3C0OX)	
ISONONYL ISONONANOATE (UNII: S4V5BS6GCX)	
MAGNESIUM SULFATE (UNII: DE08037SAB)	
LAURETH-7 (UNII: Z95S6G8201)	
SILICA DIMETHYL SILYLATE (UNII: EU2PSP0G0W)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
CHLORPHENESIN (UNII: I670DAL4SZ)	
DISODIUM STEAROYL GLUTAMATE (UNII: 45ASM2L11M)	
PROPYLENE CARBONATE (UNII: 8D08K3S51E)	
XANTHAN GUM (UNII: TTV12P4NEE)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
ALUMINUM HYDROXIDE (UNII: 5QB0T2IUN0)	
.ALPHA.-HEXYLCINNAMALDEHYDE (UNII: 7X6O37OK2I)	
BUTYLPHENYL METHYLPROPIONAL (UNII: T7540GJV69)	
BENZYL SALICYLATE (UNII: WAO5MKN9TU)	
LINALOOL, (+/-)- (UNII: D81QY6I88E)	
GERANIOL (UNII: L837108USY)	
BUTYLATED HYDROXYTOLUENE (UNII: 1P9D0Z171K)	
ISOMETHYL-.ALPHA.-IONONE (UNII: 9XP4LC555B)	
.BETA.-CITRONELLOL, (R)- (UNII: P010UT964K)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
FERRIC OXIDE RED (UNII: 1K09F3G675)	

## Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:76485-1031-1	30 mL in 1 BOTTLE, DISPENSING		

## Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part352	09/15/2012	

**Labeler** - Rimmel Inc. (949791073)

## Establishment

Name	Address	ID/FEI	Business Operations
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Revised: 10/2012

Rimmel Inc.