

MOISTURIZING SUNBLOCK - titanium dioxide lotion

Vienna Health and Beauty Corporation

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.

Drug Facts

Active Ingredients:	Purpose
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Titanium Dioxide 8%	Sunscreen
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Octyl Methoxycinnamate 6%	Sunscreen
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Benzophenone 3%	Sunscreen
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Other Ingredients: Purified Water (Acqua), Caprylic/Capic Triglyceride, C12-15 Alkyl Benzoate, Glycerin, Cetyl Alcohol, Stearyl Alcohol, Isopropyl Myristate, Propylene Glycol, Ceteareth-6, Ceteareth-25, Dimethicone, Glyceryl Stearate, Stearic Acid, Tocopherol Acetate9Vitamin E), Alpha Bisabolol, Triethanolamine, Diazolidinyl Urea, Methylparaben, Propylparaben, Allantoin, BHT.

Vienna Health and Beauty - MOISTURIZING SUNBLOCK - UVA/UVB Protection - SPF30



MOISTURIZING SUNBLOCK

titanium dioxide lotion

Product Information				
Product Type		HUMAN OTC DRUG	Item Code (Source)	
Route of Administration		TOPICAL	NDC:52625-100	
Active Ingredient/Active Moiety				
Ingredient Name			Basis of Strength	Strength
TITANIUM DIOXIDE (UNII: 15FIX9V2JP) (TITANIUM DIOXIDE - UNII:15FIX9V2JP)			TITANIUM DIOXIDE	8 g in 100 g
OCTINOXATE (UNII: 4Y5P7MUD51) (OCTINOXATE - UNII:4Y5P7MUD51)			OCTINOXATE	6 g in 100 g
BENZOPHENONE (UNII: 701M4TTV9O) (BENZOPHENONE - UNII:701M4TTV9O)			BENZOPHENONE	3 g in 100 g
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:52625-100-02	1 in 1 CARTON		
1	NDC:52625-100-01	100 g in 1 JAR		
Marketing Information				
Marketing Category	Application Number or Monograph Citation		Marketing Start Date	Marketing End Date
OTC monograph final	part352		07/01/2010	

Labeler - Vienna Health and Beauty Corporation (831463919)

Registrant - Vienna Health and Beauty Corporation (831463919)

Establishment			
Name	Address	ID/FEI	Business Operations
Vienna Health and Beauty Corporation		831463919	manufacture