TOTAL PROTECTOR 30 - zinc oxide,octinaxade,octisalate cream
MD Formulation

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.

Sun Total Protector 30

Zinc Oxide - 7.8%
Octinoxate - 7.5%
Octisalate - 4.0%

Zinc Oxide - 7.8% (Sunscreen)
Octinoxate - 7.5% (Sunscreen)
Octisalate - 4.0% (Sunscreen)

Provide the skin with maximum protection against the environmental factors that lead to premature skin aging.

md formulations®

SPF 30

SUN

total protector 30
maximum daily protection

écran protection
maximum 30
protection maximale de jour

3 ml (0.1 fl. oz.)

sample not for resale
échantillon vente interdite
muestra gratuita, no para vender

US **PROFESSIONAL TREATMENT PRODUCT**
Provide the skin with maximum protection against the environmental factors that lead to premature skin aging.

Active Ingredients: Zinc Oxide (CI 77947) -7.8%(Sunscreen), Octinoxate - 7.5%(Sunscreen), Octisalate - 4.0%(Sunscreen).

F **PRODUIT DE SOIN PROFESSIONNEL**
Assure une protection maximum de la peau contre certains facteurs liés á l'environnement, pouvant provoquer un vieillissement prématuré de la peau.

ES **PRODUCTO PARA TRATAMIENTO PROFESIONAL**
Proporciona a la piel la máxima protección contra los factores ambientales que causan un prematuro envejecimiento cutáneo.

Dist. by md formulations®
San Francisco, CA 94105 USA
Bare Escentuals UK Ltd., SE13 1ND, UK
www.mdformulations.com
Made in USA • 31039

TOTAL PROTECTOR 30

zinc oxide,octinaxade,octisalate cream

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:66078-368
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
Zinc Oxide (UNII: SOI2LOH54Z) (Zinc Oxide - UNII:SOI2LOH54Z)	Zinc Oxide	7.8 g in 100 g
Octinoxate (UNII: 4Y5P7MUD51) (Octinoxate - UNII:4Y5P7MUD51)	Octinoxate	7.5 g in 100 g
OCTISALATE (UNII: 4X49Y0596W) (OCTISALATE - UNII:4X49Y0596W)	OCTISALATE	4.0 g in 100 g

Inactive Ingredients	
Ingredient Name	**Strength**
GLYCERIN (UNII: PDC6A3C0OX)	
ALLANTOIN (UNII: 344S277G0Z)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
XANTHAN GUM (UNII: TTV12P4NEE)	
CETYL HYDROXYETHYLCELLULOSE (350000 MW) (UNII: T7SWE4S2TT)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
Diazolidinyl Urea (UNII: H5RIZ3MPW4)	
Iodopropynyl Butylcarbamate (UNII: 603P14DHEB)	
GLYCERYL MONOSTEARATE (UNII: 230OU9XXE4)	
Steareth-2 (UNII: V56DFE46J5)	
GLYCERYL MONOSTEARATE (UNII: 230OU9XXE4)	
PEG-100 STEARATE (UNII: YD01N1999R)	
Steareth-100 (UNII: 4OH5W9UM87)	
TRIACONTANYL PVP (WP-660) (UNII: N0SS3Q238D)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:66078-368-17	288 in 1 CASE		
1		3.24 g in 1 PACKET		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part352	12/13/2001	

Labeler - MD Formulation (087008363)

Registrant - Harmony Labs, Inc. (105803274)

Establishment

Name	Address	ID/FEI	Business Operations
Harmony Labs, Inc.		105803274	manufacture, label, pack, relabel, repack

Revised: 7/2011 MD Formulation