

**LIP PROTECTOR SUNSCREEN SPF15 - octinoxate,zinc oxide,oxybenzone lipstick
Mary Kay INC.**

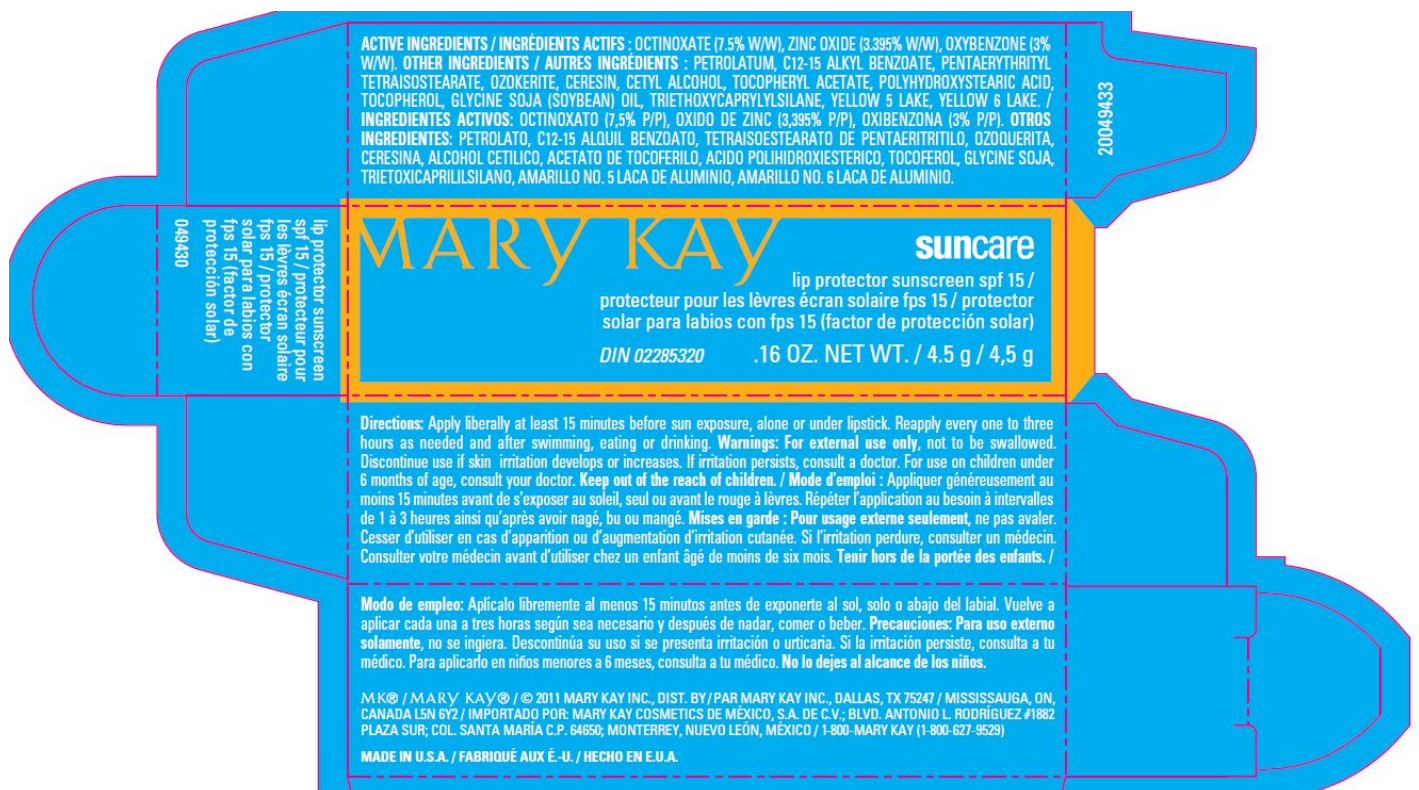
Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

ACTIVE INGREDIENTS: OCTINOXATE(7.5% W/W), ZINC OXIDE(3.395% W/W),OXYBENZONE(3%W/W)

DIRECTION: Apply liberally at least 15 minutes before sun exposure, alone or under lipstick. Reapply every one to three hours as needed and after swimming, eating or drinking .3

WARNING : For external use only. Discontinue use if irritation or rash develops. For use under 6 month of age , consult your doctor. Keep out reach of children.

PETROLATUM , C12-15 ALKYL BENZOATE , PENTAETHYTHRITYL TETRAISOSTEARATE ,
OZOKERITE , CERESIN , CETYL ALCOHOL , TOCOPHERYL ACETATE,
POLYHYDROXYSTEARIC ACID,TOCOPEROL , GLYCINE SOJA (SOYBEAN) OIL ,
TRIETHOXYCAPRYLRLSILANE , YELLOW 5 LAKE , YELLOW 6 LAKE



LIP PROTECTOR SUNSCREEN SPF15

octinoxate,zinc oxide,oxybenzone lipstick

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:51531-9480
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
Octinoxate (UNII: 4Y5P7MUD51) (Octinoxate - UNII:4Y5P7MUD51)	Octinoxate	0.338 g in 4.5 g
Oxybenzone (UNII: 95OOS7VE0Y) (Oxybenzone - UNII:95OOS7VE0Y)	Oxybenzone	0.135 g in 4.5 g
Zinc Oxide (UNII: SOI2LOH54Z) (Zinc Oxide - UNII:SOI2LOH54Z)	Zinc Oxide	0.152 g in 4.5 g

Inactive Ingredients

Ingredient Name	Strength
PETROLATUM (UNII: 4T6H12BN9U)	
ALKYL (C12-15) BENZOATE (UNII: A9EJ3J61HQ)	
CERESIN (UNII: Q1LS2UJO3A)	
CETYL ALCOHOL (UNII: 936JST6JCN)	
ACETATE ION (UNII: 569DQM74SC)	
POLYHYDROXYSTEARIC ACID (2300 MW) (UNII: YXH47AOU0F)	
ALPHA-TOCOPHEROL (UNII: H4N855PNZ1)	
SOYBEAN OIL (UNII: 241ATL177A)	
TRIETHOXYCAPRYLYLSILANE (UNII: LDC331P08E)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:51531-9480-0	4.5 g in 1 CONTAINER		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part352	03/01/2007	

Labeler - Mary Kay INC. (103978839)

Revised: 3/2012

Mary Kay INC.