TIMEWISE DAY SOLUTION SUNSCREEN SPF 35 -
homosalate,octisalate,oxybenzone,octocrylene,avobenzone cream
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.

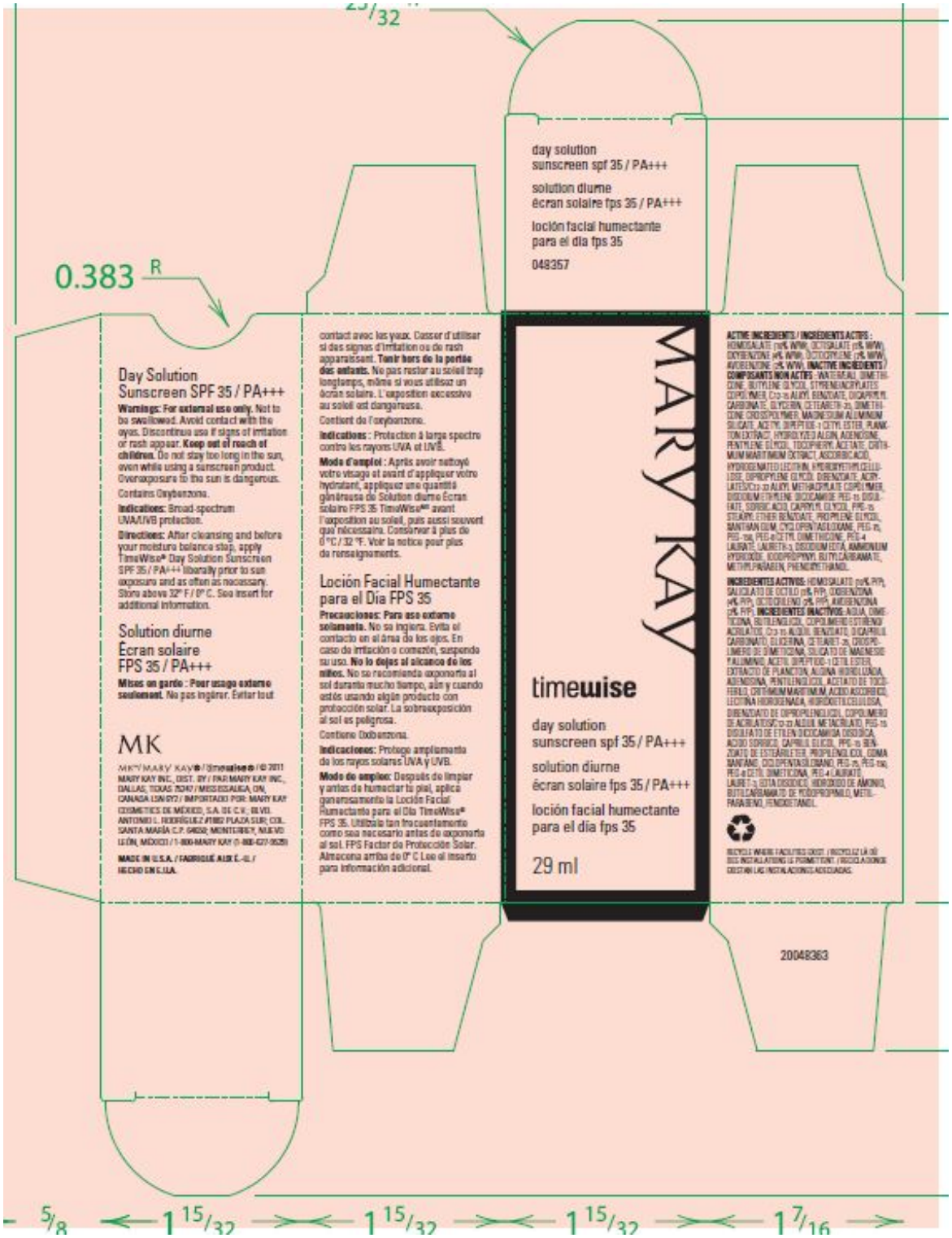
Active Ingredients ; HOMOSALATE (10% w/w) ,OCTISALATE(5% w/w),OXYBENZONE(4%w/w), OCOCRYLENE (2% w/w), AVOBENZONE(2% w/w)

Warnings; For external use only. Not to be swalloed. Avoid contact with the eyes. Discontinue use if signs of irritation or rash appear.

Keep out of reach of children. Do not stay too long in the sun, even while using a sunscreen product.

Indication; Broad-spectrum UVA/UVB protection.

Water,Dimethicone, Butylene Glycol,Styrene/Acrylates Copolymer,C12-15 Alkyl Benzoate,Dicapryl Carbonate,Glycerin,Ceteareth-25,Dimethicone Crosspolymer,Magnesium Aluminum Silicate, Acetyl Dipeptide-1,Cetyl Ester, Plankton Extract,Hydrolyzed Algin, Adenosin, Pentylene Glycol , Tocopheryl Acetate,Crithmum Marithimum Extract,Ascorbic Acid,Hydrogenated Lecithin,Hydroxyethylcellulose,Dipropylene Glycol,Dibenzoate Acrylate / C12-22 Alkyl Methacrylate Crosspolymer ,Disodium Ethylene Dicocamide PEG-15 Disulfate,Sorbic Acid ,Caprylyl Glycol, PPG-15 Stearyl Ether Benzoate,Propylene Glycol , Xanthan Gum, Cyclopentasiloxane, PEG-75,PEG-150,PEG-6 Cetyl Dimethicone,PEG-4 Laurate,Laureth-3 ,Disodium EDTA , Ammonium Hydroxide,Iodopropyl Butylcarbamate,Methylparaben,Phenoxyethanol



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Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:51531-8363
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
Oxybenzone (UNII: 95OOS7VE0Y) (Oxybenzone - UNII:95OOS7VE0Y)	Oxybenzone	1.16 mL in 29 mL
Avobenzone (UNII: G63QQF2NOX) (Avobenzone - UNII:G63QQF2NOX)	Avobenzone	0.58 mL in 29 mL
Octisalate (UNII: 4X49Y0596W) (Octisalate - UNII:4X49Y0596W)	Octisalate	1.45 mL in 29 mL
Homosalate (UNII: V06SV4M95S) (Homosalate - UNII:V06SV4M95S)	Homosalate	2.9 mL in 29 mL
OCTOCRYLENE (UNII: 5A68WGF6WM) (OCTOCRYLENE - UNII:5A68WGF6WM)	OCTOCRYLENE	0.58 mL in 29 mL

Inactive Ingredients	
Ingredient Name	**Strength**
Water (UNII: 059QF0KO0R)	
Butylene Glycol (UNII: 3XUS85K0RA)	
C12-15 Alkyl Benzoate (UNII: A9EJ3J61HQ)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:51531-8363-0	29 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	03/01/2012	

Labeler - Mary Kay Inc. (103978839)

Revised: 3/2012 Mary Kay Inc.