MARY KAY CC CREAM SUNSCREEN BROAD SPECTRUM SPF 15 VERY DEEP- homosalate, octinoxate, oxybenzone 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.

Mary Kay CC Cream Sunscreen SPF 15 Very Deep

Drug Facts

Active ingredients

Homosalate 5%

Octinoxate 6.5%

Oxybenzone 1.2 %

Purpose

Sunscreen

Uses

- 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

Warnings

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

Keep out of reach of children.

If product is swallowed, get medical help or contact a Poison Control Center right away.

Directions

- 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 SPF of 15 or higher and other sun protection measures including:
- limit time in the sun, especially from 10 a.m. - 2 p.m.
- wear long-sleeve shirts, pants, hats and sunglasses
- children under 6 months: Ask a doctor

Other information

- Store at 25°C (77°F). Do not freeze or refrigerate.

Inactive ingredients

water, cyclopentasiloxane, butylene glycol, glycerin, PEG-9 polydimethylsiloxyethyl dimethicone, niacinamide, mica, PEG-9 dimethicone, magnesium sulfate, ascorbyl glucoside, silybum marianum fruit extract, tocopheryl acetate, salix nigra (willow) bark extract, salicylic acid, adenosine, dimethicone/PEG-10/15 crosspolymer, disodium stearoyl glutamate, xanthan gum, dipropylene glycol, disodium EDTA, cyclohexasiloxane, sodium citrate, tocopherol, sorbic acid, sodium benzoate, benzyl alcohol, aluminum hydroxide, titanium dioxide, iron oxides

Questions or comments?

Call toll free 1-800-627-9529

Principal Display Panel - 29 mL carton

Mary Kay

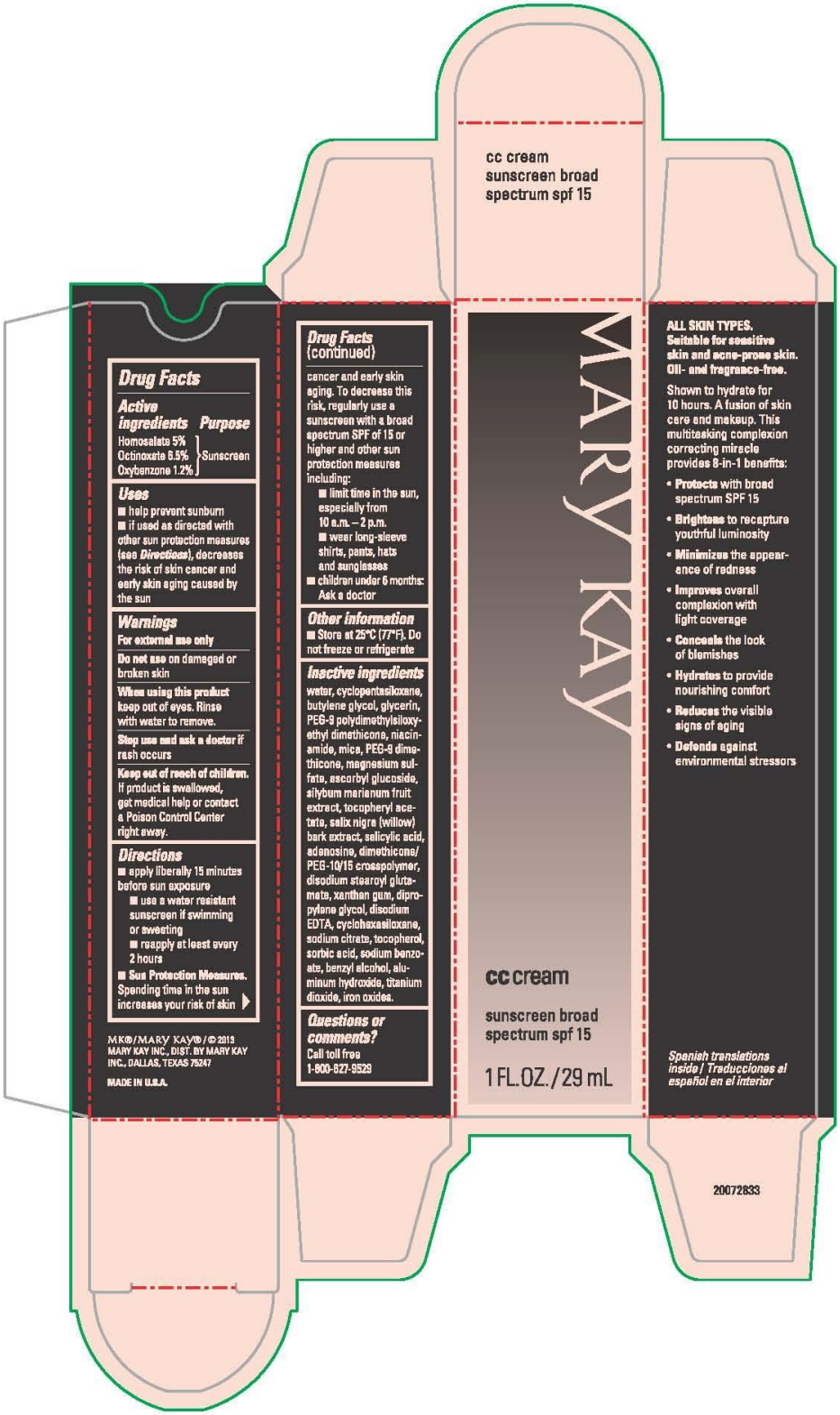
cc cream

sunscreen broad

spectrum spf 15

cream

1 FL. OZ. / 29 mL



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DEEP

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Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:51531-6620
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
HOMOSALATE (UNII: V06SV4M95S) (HOMOSALATE - UNII:V06SV4M95S)	HOMOSALATE	5 g in 100 mL
OCTINOXATE (UNII: 4Y5P7MUD51) (OCTINOXATE - UNII:4Y5P7MUD51)	OCTINOXATE	6.5 g in 100 mL
OXYBENZONE (UNII: 95OOS7VE0Y) (OXYBENZONE - UNII:95OOS7VE0Y)	OXYBENZONE	1.2 g in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
CYCLOMETHICONE 5 (UNII: 0THT5PCI0R)	
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	
GLYCERIN (UNII: PDC6A3C0OX)	
PEG-9 POLYDIMETHYLSILOXYETHYL DIMETHICONE (UNII: TYP81E471F)	
NIACINAMIDE (UNII: 25X51I8RD4)	
MICA (UNII: V8A1AW0880)	
PEG-9 DIMETHICONE (400 CST) (UNII: 9OZ27X065D)	
MAGNESIUM SULFATE, UNSPECIFIED FORM (UNII: DE08037SAB)	
ASCORBYL GLUCOSIDE (UNII: 2V52R0NHXW)	
MILK THISTLE (UNII: U946SH95EE)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
SALIX NIGRA BARK (UNII: QU52J3A5B3)	
SALICYLIC ACID (UNII: O414PZ4LPZ)	
ADENOSINE (UNII: K72T3FS567)	
DISODIUM STEAROYL GLUTAMATE (UNII: 45ASM2L11M)	
XANTHAN GUM (UNII: TTV12P4NEE)	
DIPROPYLENE GLYCOL (UNII: E107L85C40)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
CYCLOMETHICONE 6 (UNII: XHK3U310BA)	
SODIUM CITRATE, UNSPECIFIED FORM (UNII: 1Q73Q2JULR)	
TOCOPHEROL (UNII: R0ZB2556P8)	
SORBIC ACID (UNII: X045WJ989B)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
BENZYL ALCOHOL (UNII: LKG8494WBH)	
ALUMINUM HYDROXIDE (UNII: 5QB0T2IUN0)	
FERRIC OXIDE RED (UNII: 1K09F3G675)	
FERRIC OXIDE YELLOW (UNII: EX438O2MRT)	
FERROSOFERRIC OXIDE (UNII: XM0M87F357)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
DIMETHICONE/PEG-10/15 CROSSPOLYMER (UNII: 21AS8B1BSS)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:51531-6620-1	1 in 1 CARTON	02/16/2016	
1		29 mL in 1 TUBE; Type 0: Not a Combination Product		
2	NDC:51531-6620-3	1 mL in 1 PACKET; Type 0: Not a Combination Product	02/16/2016	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	M020	02/16/2016	

Labeler - Mary Kay Inc. (049994452)

Establishment

Name	Address	ID/FEI	Business Operations
Englewood Lab Inc.		172198223	manufacture(51531-6620)

Establishment

Name	Address	ID/FEI	Business Operations
Mary Kay Inc.		103978839	manufacture(51531-6620)

Revised: 7/2023 Mary Kay Inc.