

MAGIC FOUNDATION FLAWLESS LONG-LASTING COVERAGE SUNSCREEN BROAD SPECTRUM SPF 15- titanium dioxide and zinc oxide suspension

Charlotte Tilbury Beauty Ltd.

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.

Magic Foundation Flawless Long-Lasting Coverage Sunscreen Broad Spectrum SPF 15

Drug Facts

<i>Active Ingredients</i>	<i>Purpose</i>
Titanium Dioxide 3.19%	Sunscreen
Zinc Oxide 0.94%	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. water to remove.

Keep out of reach of children. If product is swallowed, get medical help or contact a Poison Control Center right away.

Directions:

- Apply liberally and evenly 15 minutes before sun exposure.
- Children under 6 months of age: Ask a doctor.
- **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 value 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-sleeved shirts, pants, hats, and sunglasses.
- Reapply at least every 2 hours or immediately after towel drying.
- Use a water-resistant sunscreen if swimming or sweating.

Inactive Ingredients:

aqua (water), cyclopentasiloxane, cyclohexasiloxane, talc, cetyl peg/ppg-10/1 dimethicone, glycerin, nylon-12, hdi/trimethylol hexyllactone crosspolymer, acrylates/dimethicone copolymer, phenoxyethanol, benzyl alcohol, sodium chloride, magnesium sulfate, hydrogen dimethicone, sodium dehydroacetate, alumina, disteardimonium hectorite, dehydroacetic acid, potassium sorbate, benzoic acid, dimethicone, tetrasodium edta, tocopheryl acetate, aluminum hydroxide, triethoxycaprylylsilane, ascorbyl tetraisopalmitate, propylene carbonate, sodium hyaluronate, butylene glycol, silica,



Shade 2 Fair



Shade 3 Fair



Shade 3.5 Fair



Shade 4 Fair



Shade 4.5 Medium



Shade 5 Medium



Shade 6 Medium



Shade 7 Medium



Shade 8 Medium



Shade 9 Dark



Shade 9.5 Dark



Shade 10 Dark



Shade 11 Dark



Shade 12 Dark



**MAGIC FOUNDATION FLAWLESS LONG-LASTING COVERAGE SUNSCREEN
BROAD SPECTRUM SPF 15**

titanium dioxide and zinc oxide suspension

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:70186-566
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
TITANIUM DIOXIDE (UNII: 15FIX9V2JP) (TITANIUM DIOXIDE - UNII:15FIX9V2JP)	TITANIUM DIOXIDE	0.957 g in 30 mL
ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC OXIDE - UNII:SOI2LOH54Z)	ZINC OXIDE	0.282 g in 30 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
CYCLOMETHICONE 5 (UNII: 0THT5PC10R)	
CYCLOMETHICONE 6 (UNII: XHK3U310BA)	
TALC (UNII: 7SEV7J4R1U)	
CETYL PEG/PPG-10/1 DIMETHICONE (HLB 2) (UNII: V2W71V8T0X)	
GLYCERIN (UNII: PDC6A3C0OX)	
NYLON-12 (UNII: 446U8J075B)	
HEXAMETHYLENE DIISOCYANATE/TRIMETHYLOL HEXYLLACTONE CROSSPOLYMER (UNII: WB5K9Y35Y9)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
BENZYL ALCOHOL (UNII: LKG8494WBH)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
MAGNESIUM SULFATE, UNSPECIFIED (UNII: DE08037SAB)	
SODIUM DEHYDROACETATE (UNII: 8W46YN971G)	
ALUMINUM OXIDE (UNII: LM26O6933)	
DISTEARDIMONIUM HECTORITE (UNII: X687XDK09L)	
DEHYDROACETIC ACID (UNII: 2KAG279R6R)	
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)	
BENZOIC ACID (UNII: 8SKN0B0MIM)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
EDETATE SODIUM (UNII: MP1J8420LU)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
ALUMINUM HYDROXIDE (UNII: 5QB0T2IUN0)	
TRIETHOXYCAPRYLYLSILANE (UNII: LDC331P08E)	
ASCORBYL TETRAISOPALMITATE (UNII: 47143LT58A)	
PROPYLENE CARBONATE (UNII: 8D08K3S51E)	
HYALURONATE SODIUM (UNII: YSE9PPT4TH)	
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
LARICIFOMES OFFICINALIS FRUITING BODY (UNII: 7IFM8431X3)	
POLYOXYL 40 HYDROGENATED CASTOR OIL (UNII: 7YC686GQ8F)	
FERRIC OXIDE RED (UNII: 1K09F3G675)	

Product Characteristics

Color	brown (1 Fair)	Score	
Shape	capsule	Size	
Flavor		Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70 186-566-01	1 in 1 CARTON		
1		30 mL in 1 BOTTLE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part352	02/22/2016	

MAGIC FOUNDATION FLAWLESS LONG-LASTING COVERAGE SUNSCREEN BROAD SPECTRUM SPF 15

titanium dioxide and zinc oxide suspension

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:70 186-567
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
TITANIUM DIOXIDE (UNII: 15FIX9 V2JP) (TITANIUM DIOXIDE - UNII:15FIX9 V2JP)	TITANIUM DIOXIDE	0.957 g in 30 mL
ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC OXIDE - UNII:SOI2LOH54Z)	ZINC OXIDE	0.282 g in 30 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
CYCLOMETHICONE 5 (UNII: 0THF5PCI0R)	
CYCLOMETHICONE 6 (UNII: XHK3U310BA)	
TALC (UNII: 7SEV7J4R1U)	
CETYL PEG/PPG-10/1 DIMETHICONE (HLB 2) (UNII: V2W71V8T0X)	
GLYCERIN (UNII: PDC6A3C0OX)	
NYLON-12 (UNII: 446U8J075B)	
HEXAMETHYLENE DIISOCYANATE/TRIMETHYLOL HEXYLLACTONE CROSSPOLYMER (UNII: WB5K9Y35Y9)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
BENZYL ALCOHOL (UNII: LKG8494WBH)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
MAGNESIUM SULFATE, UNSPECIFIED (UNII: DE08037SAB)	
SODIUM DEHYDROACETATE (UNII: 8W46YN971G)	

ALUMINUM OXIDE (UNII: LM26O6933)
DISTEARDIMONIUM HECTORITE (UNII: X687XDK09L)
DEHYDROACETIC ACID (UNII: 2KAG279R6R)
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)
BENZOIC ACID (UNII: 8SKN0B0MIM)
DIMETHICONE (UNII: 92RU3N3Y1O)
EDETATE SODIUM (UNII: MP1J8420LU)
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)
ALUMINUM HYDROXIDE (UNII: 5QB0T2IUN0)
TRIETHOXYCAPRYLYLSILANE (UNII: LDC331P08E)
ASCORBYL TETRAISOPALMITATE (UNII: 47143LT58A)
PROPYLENE CARBONATE (UNII: 8D08K3S51E)
HYALURONATE SODIUM (UNII: YSE9PPT4TH)
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)
LARICIFOMES OFFICINALIS FRUITING BODY (UNII: 7IFM8431X3)
POLYOXYL 40 HYDROGENATED CASTOR OIL (UNII: 7YC686GQ8F)
FERRIC OXIDE RED (UNII: 1K09F3G675)

Product Characteristics

Color	brown (2 Fair)	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70186-567-01	1 in 1 CARTON		
1		30 mL in 1 BOTTLE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part352	02/22/2016	

MAGIC FOUNDATION FLAWLESS LONG-LASTING COVERAGE SUNSCREEN BROAD SPECTRUM SPF 15

titanium dioxide and zinc oxide suspension

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:70186-568
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
TITANIUM DIOXIDE (UNII: 15FIX9V2JP) (TITANIUM DIOXIDE - UNII:15FIX9V2JP)	TITANIUM DIOXIDE	0.957 g in 30 mL
ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC OXIDE - UNII:SOI2LOH54Z)	ZINC OXIDE	0.282 g in 30 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
CYCLOMETHICONE 5 (UNII: 0THF5PCI0R)	
CYCLOMETHICONE 6 (UNII: XHK3U310BA)	
TALC (UNII: 7SEV7J4R1U)	
CETYL PEG/PPG-10/1 DIMETHICONE (HLB 2) (UNII: V2W71V8T0X)	
GLYCERIN (UNII: PDC6A3C0OX)	
NYLON-12 (UNII: 446U8J075B)	
HEXAMETHYLENE DIISOCYANATE/TRIMETHYLOL HEXYLLACTONE CROSSPOLYMER (UNII: WB5K9Y35Y9)	
PHENOXYETHANOL (UNII: HE492ZZ3T)	
BENZYL ALCOHOL (UNII: LKG8494WBH)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
MAGNESIUM SULFATE, UNSPECIFIED (UNII: DE08037SAB)	
SODIUM DEHYDROACETATE (UNII: 8W46YN971G)	
ALUMINUM OXIDE (UNII: LM26O6933)	
DISTEARDIMONIUM HECTORITE (UNII: X687XDK09L)	
DEHYDROACETIC ACID (UNII: 2KAG279R6R)	
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)	
BENZOIC ACID (UNII: 8SKN0B0MIM)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
EDETATE SODIUM (UNII: MP1J8420LU)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
ALUMINUM HYDROXIDE (UNII: 5QB0T2IUN0)	
TRIETHOXYCAPRYLSILANE (UNII: LDC331P08E)	
ASCORBYL TETRAISOPALMITATE (UNII: 47143LT58A)	
PROPYLENE CARBONATE (UNII: 8D08K3S51E)	
HYALURONATE SODIUM (UNII: YSE9PPT4TH)	
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
LARICIFOMES OFFICINALIS FRUITING BODY (UNII: 7IFM8431X3)	
POLYOXYL 40 HYDROGENATED CASTOR OIL (UNII: 7YC686GQ8F)	
FERRIC OXIDE RED (UNII: 1K09F3G675)	

Product Characteristics

Color	brown (3 Fair)	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70 186-568-01	1 in 1 CARTON		
1		30 mL in 1 BOTTLE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part352	02/22/2016	

MAGIC FOUNDATION FLAWLESS LONG-LASTING COVERAGE SUNSCREEN BROAD SPECTRUM SPF 15

titanium dioxide and zinc oxide suspension

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:70 186-569
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
TITANIUM DIOXIDE (UNII: 15FIX9V2JP) (TITANIUM DIOXIDE - UNII:15FIX9V2JP)	TITANIUM DIOXIDE	0.957 g in 30 mL
ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC OXIDE - UNII:SOI2LOH54Z)	ZINC OXIDE	0.282 g in 30 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
CYCLOMETHICONE 5 (UNII: 0TH5PC10R)	
CYCLOMETHICONE 6 (UNII: XHK3U310BA)	
TALC (UNII: 7SEV7J4R1U)	
CETYL PEG/PPG-10/1 DIMETHICONE (HLB 2) (UNII: V2W71V8T0X)	
GLYCERIN (UNII: PDC6A3C0OX)	
NYLON-12 (UNII: 446U8J075B)	
HEXAMETHYLENE DIISO CYANATE/TRIMETHYLOL HEXYLLACTONE CROSSPOLYMER (UNII: WB5K9Y35Y9)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
BENZYL ALCOHOL (UNII: LKG8494WBH)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
MAGNESIUM SULFATE, UNSPECIFIED (UNII: DE08037SAB)	
SODIUM DEHYDRO ACETATE (UNII: 8W46YN971G)	
ALUMINUM OXIDE (UNII: LM26O6933)	
DISTEARDIMONIUM HECTORITE (UNII: X687XDK09L)	
DEHYDROACETIC ACID (UNII: 2KAG279R6R)	
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)	
BENZOIC ACID (UNII: 8SKN0B0MIM)	

DIMETHICONE (UNII: 92RU3N3Y1O)
EDETATE SODIUM (UNII: MP1J8420LU)
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)
ALUMINUM HYDROXIDE (UNII: 5QB0T2IUN0)
TRIETHOXYCAPRYLSILANE (UNII: LDC331P08E)
ASCORBYL TETRAISOPALMITATE (UNII: 47143LT58A)
PROPYLENE CARBONATE (UNII: 8D08K3S51E)
HYALURONATE SODIUM (UNII: YSE9PPT4TH)
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)
LARICIFOMES OFFICINALIS FRUITING BODY (UNII: 7IFM8431X3)
POLYOXYL 40 HYDROGENATED CASTOR OIL (UNII: 7YC686GQ8F)
FERRIC OXIDE RED (UNII: 1K09F3G675)

Product Characteristics	
Color	brown (3.5 Fair)
Shape	
Flavor	
Contains	

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70186-569-01	1 in 1 CARTON		
1		30 mL in 1 BOTTLE; Type 0: Not a Combination Product		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part352	02/22/2016	

MAGIC FOUNDATION FLAWLESS LONG-LASTING COVERAGE SUNSCREEN BROAD SPECTRUM SPF 15
titanium dioxide and zinc oxide suspension

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:70186-570
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	Basis of Strength	Strength
TITANIUM DIOXIDE (UNII: 15FIX9V2JP) (TITANIUM DIOXIDE - UNII:15FIX9V2JP)	TITANIUM DIOXIDE	0.957 g in 30 mL
ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC OXIDE - UNII:SOI2LOH54Z)	ZINC OXIDE	0.282 g in 30 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
CYCLOMETHICONE 5 (UNII: 0THT5PCI0R)	
CYCLOMETHICONE 6 (UNII: XHK3U310BA)	
TALC (UNII: 7SEV7J4R1U)	
CETYL PEG/PPG-10/1 DIMETHICONE (HLB 2) (UNII: V2W71V8T0X)	
GLYCERIN (UNII: PDC6A3C0OX)	
NYLON-12 (UNII: 446U8J075B)	
HEXAMETHYLENE DIISO CYANATE/TRIMETHYLOL HEXYLLACTONE CROSSPOLYMER (UNII: WB5K9Y35Y9)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
BENZYL ALCOHOL (UNII: LKG8494WBH)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
MAGNESIUM SULFATE, UNSPECIFIED (UNII: DE08037SAB)	
SODIUM DEHYDRO ACETATE (UNII: 8W46YN971G)	
ALUMINUM OXIDE (UNII: LM26O6933)	
DISTEARDIMONIUM HECTORITE (UNII: X687XDK09L)	
DEHYDRO ACETIC ACID (UNII: 2KAG279R6R)	
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)	
BENZOIC ACID (UNII: 8SKN0B0MIM)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
EDETATE SODIUM (UNII: MP1J8420LU)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
ALUMINUM HYDROXIDE (UNII: 5QB0T2IUN0)	
TRIETHOXYCAPRYLYLSILANE (UNII: LDC331P08E)	
ASCORBYL TETRAISOPALMITATE (UNII: 47143LT58A)	
PROPYLENE CARBONATE (UNII: 8D08K3S51E)	
HYALURONATE SODIUM (UNII: YSE9PPT4TH)	
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
LARICIFOMES OFFICINALIS FRUITING BODY (UNII: 7IFM8431X3)	
POLYOXYL 40 HYDROGENATED CASTOR OIL (UNII: 7YC686GQ8F)	
FERRIC OXIDE RED (UNII: 1K09F3G675)	

Product Characteristics

Color	brown (4 Fair)	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70186-570-01	1 in 1 CARTON		
1		30 mL in 1 BOTTLE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part352	02/22/2016	

MAGIC FOUNDATION FLAWLESS LONG-LASTING COVERAGE SUNSCREEN BROAD SPECTRUM SPF 15

titanium dioxide and zinc oxide suspension

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:70186-571
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
TITANIUM DIOXIDE (UNII: 15FIX9V2JP) (TITANIUM DIOXIDE - UNII:15FIX9V2JP)	TITANIUM DIOXIDE	0.957 g in 30 mL
ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC OXIDE - UNII:SOI2LOH54Z)	ZINC OXIDE	0.282 g in 30 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
CYCLOMETHICONE 5 (UNII: 0THT5PCI0R)	
CYCLOMETHICONE 6 (UNII: XHK3U310BA)	
TALC (UNII: 7SEV7J4R1U)	
CETYL PEG/PPG-10/1 DIMETHICONE (HLB 2) (UNII: V2W71V8T0X)	
GLYCERIN (UNII: PDC6A3C0OX)	
NYLON-12 (UNII: 446U8J075B)	
HEXAMETHYLENE DIISOCYANATE/TRIMETHYLOL HEXYLLACTONE CROSSPOLYMER (UNII: WB5K9Y35Y9)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
BENZYL ALCOHOL (UNII: LKG8494WBH)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
MAGNESIUM SULFATE, UNSPECIFIED (UNII: DE08037SAB)	
SODIUM DEHYDRO ACETATE (UNII: 8W46YN971G)	
ALUMINUM OXIDE (UNII: LM26O6933)	
DISTEARDIMONIUM HECTORITE (UNII: X687XDK09L)	
DEHYDROACETIC ACID (UNII: 2KAG279R6R)	
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)	
BENZOIC ACID (UNII: 8SKN0B0MIM)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
EDETATE SODIUM (UNII: MPIJ8420LU)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
ALUMINUM HYDROXIDE (UNII: 5QB0T2IUN0)	
TRIETHOXYCAPRYLYLSILANE (UNII: LDC331P08E)	
ASCORBYL TETRAISOPALMITATE (UNII: 47143LT58A)	

PROPYLENE CARBONATE (UNII: 8D08K3S51E)	
HYALURONATE SODIUM (UNII: YSE9PPT4TH)	
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
LARICIFOMES OFFICINALIS FRUITING BODY (UNII: 7IFM8431X3)	
POLYOXYL 40 HYDROGENATED CASTOR OIL (UNII: 7YC686GQ8F)	
FERRIC OXIDE RED (UNII: 1K09F3G675)	

Product Characteristics

Color	brown (4.5 Medium)	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70 186-571-01	1 in 1 CARTON		
1		30 mL in 1 BOTTLE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part352	02/22/2016	

MAGIC FOUNDATION FLAWLESS LONG-LASTING COVERAGE SUNSCREEN BROAD SPECTRUM SPF 15

titanium dioxide and zinc oxide suspension

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:70 186-572
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
TITANIUM DIOXIDE (UNII: 15FIX9V2JP) (TITANIUM DIOXIDE - UNII:15FIX9V2JP)	TITANIUM DIOXIDE	0.957 g in 30 mL
ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC OXIDE - UNII:SOI2LOH54Z)	ZINC OXIDE	0.282 g in 30 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	

CYCLOMETHICONE 5 (UNII: 0THI5PCI0R)
CYCLOMETHICONE 6 (UNII: XHK3U310BA)
TALC (UNII: 7SEV7J4R1U)
CETYL PEG/PPG-10/1 DIMETHICONE (HLB 2) (UNII: V2W71V8T0X)
GLYCERIN (UNII: PDC6A3C0OX)
NYLON-12 (UNII: 446U8J075B)
HEXAMETHYLENE DIISO CYANATE/TRIMETHYLOL HEXYLLACTONE CROSSPOLYMER (UNII: WB5K9Y35Y9)
PHENOXYETHANOL (UNII: HIE492ZZ3T)
BENZYL ALCOHOL (UNII: LKG8494WBH)
SODIUM CHLORIDE (UNII: 451W47IQ8X)
MAGNESIUM SULFATE, UNSPECIFIED (UNII: DE08037SAB)
SODIUM DEHYDROACETATE (UNII: 8W46YN971G)
ALUMINUM OXIDE (UNII: LM26O6933)
DISTEARDIMONIUM HECTORITE (UNII: X687XDK09L)
DEHYDROACETIC ACID (UNII: 2KAG279R6R)
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)
BENZOIC ACID (UNII: 8SKN0B0MM)
DIMETHICONE (UNII: 92RU3N3Y1O)
EDETATE SODIUM (UNII: MP1J8420LU)
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)
ALUMINUM HYDROXIDE (UNII: 5QB0T2IUN0)
TRIETHOXYCAPRYLSILANE (UNII: LDC331P08E)
ASCORBYL TETRAISOPALMITATE (UNII: 47143LT58A)
PROPYLENE CARBONATE (UNII: 8D08K3S51E)
HYALURONATE SODIUM (UNII: YSE9PPT4TH)
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)
LARICIFOMES OFFICINALIS FRUITING BODY (UNII: 7IFM8431X3)
POLYOXYL 40 HYDROGENATED CASTOR OIL (UNII: 7YC686GQ8F)
FERRIC OXIDE RED (UNII: 1K09F3G675)

Product Characteristics

Color	brown (5 Medium)	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70186-572-01	1 in 1 CARTON		
1		30 mL in 1 BOTTLE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part352	02/22/2016	

MAGIC FOUNDATION FLAWLESS LONG-LASTING COVERAGE SUNSCREEN BROAD SPECTRUM SPF 15

titanium dioxide and zinc oxide suspension

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:70186-573
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
TITANIUM DIOXIDE (UNII: 15FIX9V2JP) (TITANIUM DIOXIDE - UNII:15FIX9V2JP)	TITANIUM DIOXIDE	0.957 g in 30 mL
ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC OXIDE - UNII:SOI2LOH54Z)	ZINC OXIDE	0.282 g in 30 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
CYCLOMETHICONE 5 (UNII: 0THT5PCI0R)	
CYCLOMETHICONE 6 (UNII: XHK3U310BA)	
TALC (UNII: 7SEV7J4R1U)	
CETYL PEG/PPG-10/1 DIMETHICONE (HLB 2) (UNII: V2W71V8T0X)	
GLYCERIN (UNII: PDC6A3C0OX)	
NYLON-12 (UNII: 446U8J075B)	
HEXAMETHYLENE DIISOCYANATE/TRIMETHYLOL HEXYLLACTONE CROSSPOLYMER (UNII: WB5K9Y35Y9)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
BENZYL ALCOHOL (UNII: LKG8494WBH)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
MAGNESIUM SULFATE, UNSPECIFIED (UNII: DE08037SAB)	
SODIUM DEHYDRO ACETATE (UNII: 8W46YN971G)	
ALUMINUM OXIDE (UNII: LM26O6933)	
DISTEARDIMONIUM HECTORITE (UNII: X687XDK09L)	
DEHYDROACETIC ACID (UNII: 2KAG279R6R)	
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)	
BENZOIC ACID (UNII: 8SKN0B0MIM)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
EDETATE SODIUM (UNII: MP1J8420LU)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
ALUMINUM HYDROXIDE (UNII: 5QB0T2IUN0)	
TRIETHOXYCAPRYLYLSILANE (UNII: LDC331P08E)	
ASCORBYL TETRAISOPALMITATE (UNII: 47143LT58A)	
PROPYLENE CARBONATE (UNII: 8D08K3S51E)	
HYALURONATE SODIUM (UNII: YSE9PPT4TH)	
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	

LARICIFOMES OFFICINALIS FRUITING BODY (UNII: 7IFM8431X3)

POLYOXYL 40 HYDROGENATED CASTOR OIL (UNII: 7YC686GQ8F)

FERRIC OXIDE RED (UNII: 1K09F3G675)

Product Characteristics

Color	brown (6 Medium)	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70 186-573-01	1 in 1 CARTON		
1		30 mL in 1 BOTTLE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part352	02/22/2016	

MAGIC FOUNDATION FLAWLESS LONG-LASTING COVERAGE SUNSCREEN BROAD SPECTRUM SPF 15

titanium dioxide and zinc oxide suspension

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:70186-574
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
TITANIUM DIOXIDE (UNII: 15FIX9V2JP) (TITANIUM DIOXIDE - UNII:15FIX9V2JP)	TITANIUM DIOXIDE	0.957 g in 30 mL
ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC OXIDE - UNII:SOI2LOH54Z)	ZINC OXIDE	0.282 g in 30 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
CYCLOMETHICONE 5 (UNII: 0TH5PC10R)	
CYCLOMETHICONE 6 (UNII: XHK3U310BA)	
TALC (UNII: 7SEV7J4R1U)	
CETYL PEG/PPG-10/1 DIMETHICONE (HLB 2) (UNII: V2W71V8T0X)	
GLYCERIN (UNII: PDC6A3C0OX)	
NYLON-12 (UNII: 446U8J075B)	

HEXAMETHYLENE DIISOCYANATE/TRIMETHYLOL HEXYLLACTONE CROSSPOLYMER (UNII: WB5K9Y35Y9)
PHENOXYETHANOL (UNII: HIE492ZZ3T)
BENZYL ALCOHOL (UNII: LKG8494WBH)
SODIUM CHLORIDE (UNII: 451W47IQ8X)
MAGNESIUM SULFATE, UNSPECIFIED (UNII: DE08037SAB)
SODIUM DEHYDROACETATE (UNII: 8W46YN971G)
ALUMINUM OXIDE (UNII: LM26O6933)
DISTEARDIMONIUM HECTORITE (UNII: X687XDK09L)
DEHYDROACETIC ACID (UNII: 2KAG279R6R)
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)
BENZOIC ACID (UNII: 8SKN0B0MIM)
DIMETHICONE (UNII: 92RU3N3Y1O)
EDETATE SODIUM (UNII: MP1J8420LU)
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)
ALUMINUM HYDROXIDE (UNII: 5QB0T2IUN0)
TRIETHOXYCAPRYLSILANE (UNII: LDC331P08E)
ASCORBYL TETRAISOPALMITATE (UNII: 47143LT58A)
PROPYLENE CARBONATE (UNII: 8D08K3S51E)
HYALURONATE SODIUM (UNII: YSE9PPT4TH)
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)
LARICIFOMES OFFICINALIS FRUITING BODY (UNII: 7IFM8431X3)
POLYOXYL 40 HYDROGENATED CASTOR OIL (UNII: 7YC686GQ8F)
FERRIC OXIDE RED (UNII: 1K09F3G675)

Product Characteristics

Color	brown (7 Medium)	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70186-574-01	1 in 1 CARTON		
1		30 mL in 1 BOTTLE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part352	02/22/2016	

MAGIC FOUNDATION FLAWLESS LONG-LASTING COVERAGE SUNSCREEN BROAD SPECTRUM SPF 15

titanium dioxide and zinc oxide suspension

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:70186-575
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
TITANIUM DIOXIDE (UNII: 15FIX9V2JP) (TITANIUM DIOXIDE - UNII:15FIX9V2JP)	TITANIUM DIOXIDE	0.957 g in 30 mL
ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC OXIDE - UNII:SOI2LOH54Z)	ZINC OXIDE	0.282 g in 30 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
CYCLOMETHICONE 5 (UNII: 0TH5PCI0R)	
CYCLOMETHICONE 6 (UNII: XHK3U310BA)	
TALC (UNII: 7SEV7J4R1U)	
CETYL PEG/PPG-10/1 DIMETHICONE (HLB 2) (UNII: V2W71V8T0X)	
GLYCERIN (UNII: PDC6A3C0OX)	
NYLON-12 (UNII: 446U8J075B)	
HEXAMETHYLENE DIISOCYANATE/TRIMETHYLOL HEXYLACTONE CROSSPOLYMER (UNII: WB5K9Y35Y9)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
BENZYL ALCOHOL (UNII: LKG8494WBH)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
MAGNESIUM SULFATE, UNSPECIFIED (UNII: DE08037SAB)	
SODIUM DEHYDRO ACETATE (UNII: 8W46YN971G)	
ALUMINUM OXIDE (UNII: LM26O6933)	
DISTEARDIMONIUM HECTORITE (UNII: X687XDK09L)	
DEHYDROACETIC ACID (UNII: 2KAG279R6R)	
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)	
BENZOIC ACID (UNII: 8SKN0B0MIM)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
EDETATE SODIUM (UNII: MP1J8420LU)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
ALUMINUM HYDROXIDE (UNII: 5QB0T2IUN0)	
TRIETHOXYCAPRYLSILANE (UNII: LDC331P08E)	
ASCORBYL TETRAISOPALMITATE (UNII: 47143LT58A)	
PROPYLENE CARBONATE (UNII: 8D08K3S51E)	
HYALURONATE SODIUM (UNII: YSE9PPT4TH)	
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
LARICIFOMES OFFICINALIS FRUITING BODY (UNII: 7IFM8431X3)	
POLYOXYL 40 HYDROGENATED CASTOR OIL (UNII: 7YC686GQ8F)	
FERRIC OXIDE RED (UNII: 1K09F3G675)	

Product Characteristics				
Color	brown (8 Medium)		Score	
Shape			Size	
Flavor			Imprint Code	
Contains				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70 186-575-01	1 in 1 CARTON		
1		30 mL in 1 BOTTLE; Type 0: Not a Combination Product		
Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph final	part352	02/22/2016		

MAGIC FOUNDATION FLAWLESS LONG-LASTING COVERAGE SUNSCREEN BROAD SPECTRUM SPF 15				
titanium dioxide and zinc oxide suspension				
Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:70 186-576	
Route of Administration	TOPICAL			
Active Ingredient/Active Moiety				
Ingredient Name		Basis of Strength	Strength	
TITANIUM DIOXIDE (UNII: 15FIX9 V2JP) (TITANIUM DIOXIDE - UNII:15FIX9 V2JP)		TITANIUM DIOXIDE	0.957 g in 30 mL	
ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC OXIDE - UNII:SOI2LOH54Z)		ZINC OXIDE	0.282 g in 30 mL	
Inactive Ingredients				
Ingredient Name				Strength
WATER (UNII: 059QF0KO0R)				
CYCLOMETHICONE 5 (UNII: 0THF5PCI0R)				
CYCLOMETHICONE 6 (UNII: XHK3U310BA)				
TALC (UNII: 7SEV7J4R1U)				
CETYL PEG/PPG-10/1 DIMETHICONE (HLB 2) (UNII: V2W71V8T0X)				
GLYCERIN (UNII: PDC6A3C0OX)				
NYLON-12 (UNII: 446U8J075B)				
HEXAMETHYLENE DIISOCYANATE/TRIMETHYLOL HEXYLLACTONE CROSSPOLYMER (UNII: WB5K9Y35Y9)				
PHENOXYETHANOL (UNII: HIE492ZZ3T)				
BENZYL ALCOHOL (UNII: LKG8494WBH)				
SODIUM CHLORIDE (UNII: 451W47IQ8X)				
MAGNESIUM SULFATE, UNSPECIFIED (UNII: DE08037SAB)				

SODIUM DEHYDRO ACETATE (UNII: 8W46YN971G)
ALUMINUM OXIDE (UNII: LM26O6933)
DISTEARDIMONIUM HECTORITE (UNII: X687XDK09L)
DEHYDRO ACETIC ACID (UNII: 2KAG279R6R)
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)
BENZOIC ACID (UNII: 8SKN0B0MIM)
DIMETHICONE (UNII: 92RU3N3Y1O)
EDETATE SODIUM (UNII: MP1J8420LU)
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)
ALUMINUM HYDROXIDE (UNII: 5QB0T2IUN0)
TRIETHOXYCAPRYLSILANE (UNII: LDC331P08E)
ASCORBYL TETRAISOPALMITATE (UNII: 47143LT58A)
PROPYLENE CARBONATE (UNII: 8D08K3S51E)
HYALURONATE SODIUM (UNII: YSE9PPT4TH)
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)
LARICIFOMES OFFICINALIS FRUITING BODY (UNII: 7IFM8431X3)
POLYOXYL 40 HYDROGENATED CASTOR OIL (UNII: 7YC686GQ8F)
FERRIC OXIDE RED (UNII: 1K09F3G675)

Product Characteristics			
Color	brown (9 Dark)	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70186-576-01	1 in 1 CARTON		
1		30 mL in 1 BOTTLE; Type 0: Not a Combination Product		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part352	02/22/2016	

MAGIC FOUNDATION FLAWLESS LONG-LASTING COVERAGE SUNSCREEN BROAD SPECTRUM SPF 15
titanium dioxide and zinc oxide suspension

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:70186-577
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
TITANIUM DIOXIDE (UNII: 15FIX9V2JP) (TITANIUM DIOXIDE - UNII:15FIX9V2JP)	TITANIUM DIOXIDE	0.957 g in 30 mL
ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC OXIDE - UNII:SOI2LOH54Z)	ZINC OXIDE	0.282 g in 30 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
CYCLOMETHICONE 5 (UNII: 0THT5PCI0R)	
CYCLOMETHICONE 6 (UNII: XHK3U310BA)	
TALC (UNII: 7SEV7J4R1U)	
CETYL PEG/PPG-10/1 DIMETHICONE (HLB 2) (UNII: V2W71V8T0X)	
GLYCERIN (UNII: PDC6A3C0OX)	
NYLON-12 (UNII: 446U8J075B)	
HEXAMETHYLENE DIISO CYANATE/TRIMETHYLOL HEXYLLACTONE CROSSPOLYMER (UNII: WB5K9Y35Y9)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
BENZYL ALCOHOL (UNII: LKG8494WBH)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
MAGNESIUM SULFATE, UNSPECIFIED (UNII: DE08037SAB)	
SODIUM DEHYDRO ACETATE (UNII: 8W46YN971G)	
ALUMINUM OXIDE (UNII: LM26O6933)	
DISTEARDIMONIUM HECTORITE (UNII: X687XDK09L)	
DEHYDRO ACETIC ACID (UNII: 2KAG279R6R)	
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)	
BENZOIC ACID (UNII: 8SKN0B0MIM)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
EDETATE SODIUM (UNII: MP1J8420LU)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
ALUMINUM HYDROXIDE (UNII: 5QB0T2IUN0)	
TRIETHOXYCAPRYLSILANE (UNII: LDC331P08E)	
ASCORBYL TETRAISOPALMITATE (UNII: 47143LT58A)	
PROPYLENE CARBONATE (UNII: 8D08K3S51E)	
HYALURONATE SODIUM (UNII: YSE9PPT4TH)	
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
LARICIFOMES OFFICINALIS FRUITING BODY (UNII: 7IFM8431X3)	
POLYOXYL 40 HYDROGENATED CASTOR OIL (UNII: 7YC686GQ8F)	
FERRIC OXIDE RED (UNII: 1K09F3G675)	

Product Characteristics

Color	brown (9.5 Dark)	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70 186-577-01	1 in 1 CARTON		
1		30 mL in 1 BOTTLE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part352	02/22/2016	

MAGIC FOUNDATION FLAWLESS LONG-LASTING COVERAGE SUNSCREEN BROAD SPECTRUM SPF 15

titanium dioxide and zinc oxide suspension

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:70 186-579
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
TITANIUM DIOXIDE (UNII: 15FIX9V2JP) (TITANIUM DIOXIDE - UNII:15FIX9V2JP)	TITANIUM DIOXIDE	0.957 g in 30 mL
ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC OXIDE - UNII:SOI2LOH54Z)	ZINC OXIDE	0.282 g in 30 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
CYCLOMETHICONE 5 (UNII: 0THF5PC10R)	
CYCLOMETHICONE 6 (UNII: XHK3U310BA)	
TALC (UNII: 7SEV7J4R1U)	
CETYL PEG/PPG-10/1 DIMETHICONE (HLB 2) (UNII: V2W71V8T0X)	
GLYCERIN (UNII: PDC6A3C0OX)	
NYLON-12 (UNII: 446U8J075B)	
HEXAMETHYLENE DIISOCYANATE/TRIMETHYLOL HEXYLLACTONE CROSSPOLYMER (UNII: WB5K9Y35Y9)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
BENZYL ALCOHOL (UNII: LKG8494WBH)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
MAGNESIUM SULFATE, UNSPECIFIED (UNII: DE08037SAB)	
SODIUM DEHYDROACETATE (UNII: 8W46YN971G)	
ALUMINUM OXIDE (UNII: LM26O6933)	
DISTEARDIMONIUM HECTORITE (UNII: X687XDK09L)	
DEHYDROACETIC ACID (UNII: 2KAG279R6R)	
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)	

BENZOIC ACID (UNII: 8SKN0B0MIM)
DIMETHICONE (UNII: 92RU3N3Y1O)
EDETATE SODIUM (UNII: MP1J8420LU)
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)
ALUMINUM HYDROXIDE (UNII: 5QB0T2IUN0)
TRIETHOXYCAPRYLYLSILANE (UNII: LDC331P08E)
ASCORBYL TETRAISOPALMITATE (UNII: 47143LT58A)
PROPYLENE CARBONATE (UNII: 8D08K3S51E)
HYALURONATE SODIUM (UNII: YSE9PPT4TH)
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)
LARICIFOMES OFFICINALIS FRUITING BODY (UNII: 7IFM8431X3)
POLYOXYL 40 HYDROGENATED CASTOR OIL (UNII: 7YC686GQ8F)
FERRIC OXIDE RED (UNII: 1K09F3G675)

Product Characteristics

Color	brown (10 Dark)	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70186-579-01	1 in 1 CARTON		
1		30 mL in 1 BOTTLE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part352	02/22/2016	

MAGIC FOUNDATION FLAWLESS LONG-LASTING COVERAGE SUNSCREEN BROAD SPECTRUM SPF 15

titanium dioxide and zinc oxide suspension

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:70186-578
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
TITANIUM DIOXIDE (UNII: 15FIX9V2JP) (TITANIUM DIOXIDE - UNII:15FIX9V2JP)	TITANIUM DIOXIDE	0.957 g in 30 mL

ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC OXIDE - UNII:SOI2LOH54Z)

ZINC OXIDE

0.282 g in 30 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
CYCLOMETHICONE 5 (UNII: 0THT5PCI0R)	
CYCLOMETHICONE 6 (UNII: XHK3U310BA)	
TALC (UNII: 7SEV7J4R1U)	
CETYL PEG/PPG-10/1 DIMETHICONE (HLB 2) (UNII: V2W71V8T0X)	
GLYCERIN (UNII: PDC6A3C0OX)	
NYLON-12 (UNII: 446U8J075B)	
HEXAMETHYLENE DIISOCYANATE/TRIMETHYLOL HEXYLLACTONE CROSSPOLYMER (UNII: WB5K9Y35Y9)	
PHENOXYETHANOL (UNII: HE492ZZ3T)	
BENZYL ALCOHOL (UNII: LKG8494WBH)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
MAGNESIUM SULFATE, UNSPECIFIED (UNII: DE08037SAB)	
SODIUM DEHYDRO ACETATE (UNII: 8W46YN971G)	
ALUMINUM OXIDE (UNII: LM26O6933)	
DISTEARDIMONIUM HECTORITE (UNII: X687XDK09L)	
DEHYDRO ACETIC ACID (UNII: 2KAG279R6R)	
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)	
BENZOIC ACID (UNII: 8SKN0B0MIM)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
EDETATE SODIUM (UNII: MP1J8420LU)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
ALUMINUM HYDROXIDE (UNII: 5QB0T2IUN0)	
TRIETHOXYCAPRYLSILANE (UNII: LDC331P08E)	
ASCORBYL TETRAISOPALMITATE (UNII: 47143LT58A)	
PROPYLENE CARBONATE (UNII: 8D08K3S51E)	
HYALURONATE SODIUM (UNII: YSE9PPT4TH)	
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
LARICIFOMES OFFICINALIS FRUITING BODY (UNII: 7IFM8431X3)	
POLYOXYL 40 HYDROGENATED CASTOR OIL (UNII: 7YC686GQ8F)	
FERRIC OXIDE RED (UNII: 1K09F3G675)	

Product Characteristics

Color	brown (11 Dark)	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70186-578-01	1 in 1 CARTON		

1	30 mL in 1 BOTTLE; Type 0: Not a Combination Product		
Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part352	02/22/2016	

MAGIC FOUNDATION FLAWLESS LONG-LASTING COVERAGE SUNSCREEN BROAD SPECTRUM SPF 15
titanium dioxide and zinc oxide suspension

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:70186-564
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety			
	Ingredient Name	Basis of Strength	Strength
	TITANIUM DIOXIDE (UNII: 15FIX9V2JP) (TITANIUM DIOXIDE - UNII:15FIX9V2JP)	TITANIUM DIOXIDE	0.957 g in 30 mL
	ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC OXIDE - UNII:SOI2LOH54Z)	ZINC OXIDE	0.282 g in 30 mL

Inactive Ingredients		
	Ingredient Name	Strength
	WATER (UNII: 059QF0KO0R)	
	CYCLOMETHICONE 5 (UNII: 0THF5PCI0R)	
	CYCLOMETHICONE 6 (UNII: XHK3U310BA)	
	TALC (UNII: 7SEV7J4R1U)	
	CETYL PEG/PPG-10/1 DIMETHICONE (HLB 2) (UNII: V2W71V8T0X)	
	GLYCERIN (UNII: PDC6A3C0OX)	
	NYLON-12 (UNII: 446U8J075B)	
	HEXAMETHYLENE DIISO CYANATE/TRIMETHYLOL HEXYLLACTONE CROSSPOLYMER (UNII: WB5K9Y35Y9)	
	PHENOXYETHANOL (UNII: HIE492ZZ3T)	
	BENZYL ALCOHOL (UNII: LKG8494WBH)	
	SODIUM CHLORIDE (UNII: 451W47IQ8X)	
	MAGNESIUM SULFATE, UNSPECIFIED (UNII: DE08037SAB)	
	SODIUM DEHYDRO ACETATE (UNII: 8W46YN971G)	
	ALUMINUM OXIDE (UNII: LM26O6933)	
	DISTEARDIMONIUM HECTORITE (UNII: X687XDK09L)	
	DEHYDRO ACETIC ACID (UNII: 2KAG279R6R)	
	POTASSIUM SORBATE (UNII: 1VPU26JZZ4)	
	BENZOIC ACID (UNII: 8SKN0B0MIM)	
	DIMETHICONE (UNII: 92RU3N3Y1O)	
	EDETATE SODIUM (UNII: MP1J8420LU)	
	.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
	ALUMINUM HYDROXIDE (UNII: 5QB0T2IUN0)	

TRIETHOXYCAPRYLYLSILANE (UNII: LDC331P08E)	
ASCORBYL TETRAISOPALMITATE (UNII: 47143LT58A)	
PROPYLENE CARBONATE (UNII: 8D08K3S51E)	
HYALURONATE SODIUM (UNII: YSE9PPT4TH)	
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
LARICIFOMES OFFICINALIS FRUITING BODY (UNII: 7IFM8431X3)	
POLYOXYL 40 HYDROGENATED CASTOR OIL (UNII: 7YC686GQ8F)	
FERRIC OXIDE RED (UNII: 1K09F3G675)	

Product Characteristics

Color	brown (12 Dark)	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70186-564-01	1 in 1 CARTON		
1		30 mL in 1 BOTTLE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part352	02/22/2016	

Labeler - Charlotte Tilbury Beauty Ltd. (218353520)