

**RODAN AND FIELDS ENHANCEMENTS LIGHT- titanium dioxide, zinc oxide powder**

**RODAN AND FIELDS ENHANCEMENTS MEDIUM- titanium dioxide, zinc oxide powder**

**RODAN AND FIELDS ENHANCEMENTS BRONZE- titanium dioxide, zinc oxide powder**

**Bentley Laboratories, LLC**

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## **Drug Facts Active ingredients**

Titanium Dioxide 9.6%

Zinc Oxide 4.9%

## **Purpose**

Sunscreen

**Keep out of each of children.**

**Keep out of reach of children.**

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

## **Uses**

- Helps prevent sunburn

## **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

## **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 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

### **Inactive ingredients**

Mica, Polymethylsilsesquioxane, Squalane, Water, Silica, Bismuth Oxide, Stearic Acid, Aluminum Hydroxide, Ceramide 3, Cholesterol, Phytosphingosine, Phenoxyethanol, Butylene Glycol, Caprylyl Glycol, Barium Sulfate, Angelica Polymorpha Sinensis Root Extract, HDI/Trimethylol Hexyllactone Crosspolymer, Potassium Sorbate, Hexylene Glycol, Dimethylacrylamide/Acrylic Acid/Polystyrene Ethyl Methacrylate Copolymer, Tranexamic Acid, Triethoxycaprylylsilane, Peat Extract, Trimyristin, Hydrogenated Lecithin, Oligopeptide-10, Tetrapeptide-14, Honey Extract, Boswellia Serrata Extract, Resveratrol Lecithin, Silver. May contain: Iron Oxides, Titanium Dioxide.

### **Other information**

Protect this product from excessive heat and direct sun.

### **Rodan + Fields ENHANCEMENTS - Light, Medium, Bronze Product Labels**

description for Change™

RODAN+FIELDS®  
ENHANCEMENTS

Mineral Peptides

BROAD SPECTRUM SPF 20

RODAN+FIELDS®  
DERMATOLOGISTS

4g/0.14 Oz.

Manufactured for Rodan and Fields, LLC 111 Maiden Lane, Suite 400 San Francisco CA 94108 Made in the USA Please Recycle

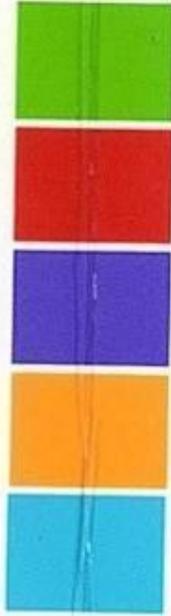
visit [rodanandfields.com](http://rodanandfields.com) QUESTIONS OR COMMENTS 1-888-995-5656

Light  
ENPL004  
08-03-0023

Medium  
ENPM004  
80-03-0024

Bronze  
ENPB004  
80-03-0025

Prescription for Change™



# RODAN+FIELDS ENHANCEMENTS

## Mineral Peptides

### BROAD SPECTRUM SPF 20

RODAN+FIELDS  
DERMATOLOGISTS

4 g/0.14 Oz.

#### Drug Facts (continued)

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

**Inactive ingredients** Mica, Polymethylsilsesquioxane, Squalane, Water, Silica, Bismuth Oxychloride, Stearic Acid, Aluminum Hydroxide, Ceramide 3, Cholesterol, Phytosphingosine, Phenoxyethanol, Butylene Glycol, Caprylyl Glycol, Barium Sulfate, Angelica Polymorpha Sinensis Root Extract, ▲

Light

ENPL004

80-03-0023

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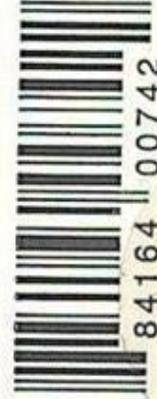
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S56978

J83772

## Drug Facts (continued)

HDI/Trimethylol Hexyllactone Crosspolymer, Potassium Sorbate, Hexylene Glycol, Dimethylacrylamide/Acrylic Acid/Polystyrene Ethyl Methacrylate Copolymer, Tranexamic Acid, Triethoxycaprylylsilane, Peat Extract, Trimyristin, Hydrogenated Lecithin, Oligopeptide-10, Tetrapeptide-14, Honey Extract, Boswellia Serrata Extract, Resveratrol, Lecithin, Silver. May Contain: Iron Oxides, Titanium Dioxide.

**Other information** Protect this product from excessive heat and direct sun.

## Drug Facts

### Active ingredients

Titanium Dioxide 9.6%  
Zinc Oxide 4.9% }

### 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.

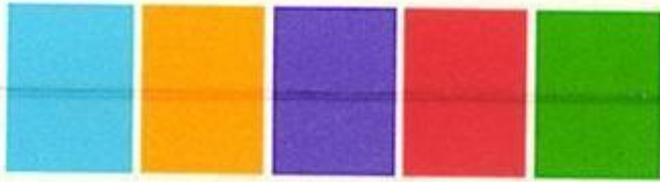
**Stop use and ask a doctor** if rash occurs

**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

Prescription for Change™



# RODAN+FIELDS® ENHANCEMENTS

## Mineral Peptides

### BROAD SPECTRUM SPF 20

RODAN+FIELDS®  
DERMATOLOGISTS

4 g/0.14 Oz.

Medium

ENPM004

80-03-0024

A73A

EXP. 07/2015

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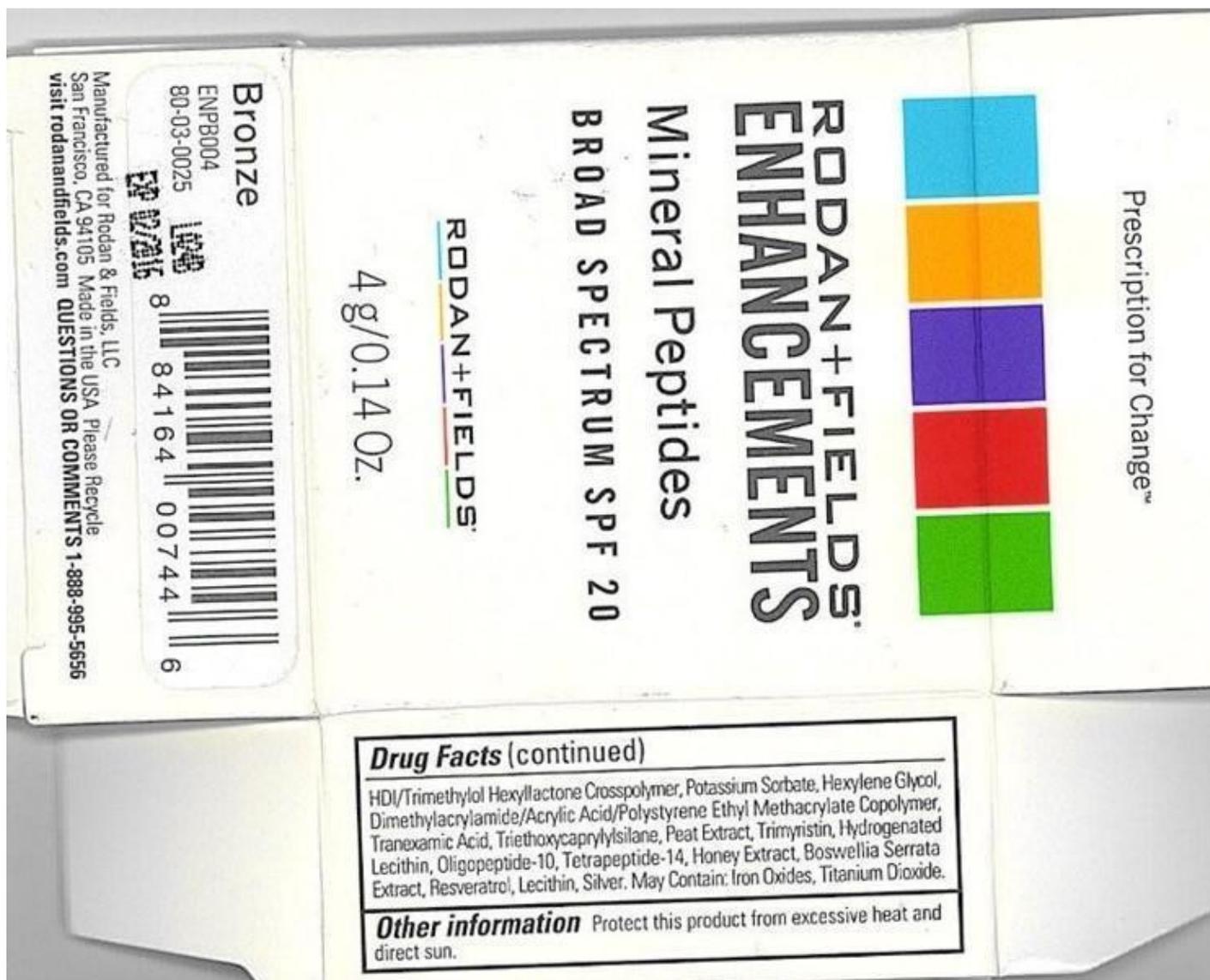
#### Drug Facts (continued)

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

#### Inactive ingredients

Mica, Polymethylsilsesquioxane, Squalane, Water, Silica, Bismuth Oxichloride, Stearic Acid, Aluminum Hydroxide, Ceramide 3, Cholesterol, Phytosphingosine, Phenoxethanol, Butylene Glycol, Caprylyl Glycol, Barium Sulfate, Angelica Polymorpha Sinensis Root Extract. ▼



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## RODAN AND FIELDS ENHANCEMENTS LIGHT

titanium dioxide, zinc oxide powder

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:54111-119
<b>Route of Administration</b>	TOPICAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>TITANIUM DIOXIDE</b> (UNII: 15FIX9V2JP) (TITANIUM DIOXIDE - UNII:15FIX9V2JP)	TITANIUM DIOXIDE	0.38400 g in 4 g
<b>ZINC OXIDE</b> (UNII: SOI2LOH54Z) (ZINC OXIDE - UNII:SOI2LOH54Z)	ZINC OXIDE	0.19600 g in 4 g

### Inactive Ingredients

Ingredient Name	Strength
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<b>MICA</b> (UNII: V8A1AW0880)
<b>POLYMETHYLSILSESQUIOXANE (4.5 MICRONS)</b> (UNII: 59Z907ZB69)
<b>SQUALANE</b> (UNII: GW89575KF9)
<b>WATER</b> (UNII: 059QF0K00R)
<b>SILICON DIOXIDE</b> (UNII: ETJ7Z6XBU4)
<b>BISMUTH OXYCHLORIDE</b> (UNII: 4ZR792I587)
<b>STEARIC ACID</b> (UNII: 4ELV7Z65AP)
<b>ALUMINUM HYDROXIDE</b> (UNII: 5QB0T2IUN0)
<b>CERAMIDE 3</b> (UNII: 4370DF050B)
<b>CHOLESTEROL</b> (UNII: 97C5T2UQ7J)
<b>PHYTOSPHINGOSINE</b> (UNII: GIN46U9Q2Q)
<b>PHENOXYETHANOL</b> (UNII: HIE492ZZ3T)
<b>BUTYLENE GLYCOL</b> (UNII: 3XUS85K0RA)
<b>CAPRYLYL GLYCOL</b> (UNII: 00YIU5438U)
<b>BARIUM SULFATE</b> (UNII: 25BB7EKE2E)
<b>ANGELICA SINENSIS ROOT</b> (UNII: B66F4574UG)
<b>HEXAMETHYLENE DIISOCYANATE/TRIMETHYLOL HEXYLLACTONE CROSSPOLYMER</b> (UNII: WB5K9Y35Y9)
<b>POTASSIUM SORBATE</b> (UNII: 1VPU26JZZ4)
<b>HEXYLENE GLYCOL</b> (UNII: KEH0A3F75J)
<b>TRANEXAMIC ACID</b> (UNII: 6T84R30KC1)
<b>TRIETHOXYCAPRYLYLSILANE</b> (UNII: LDC331P08E)
<b>TRIMYRISTIN</b> (UNII: 18L31PSR28)
<b>HYDROGENATED SOYBEAN LECITHIN</b> (UNII: H1109Z9J4N)
<b>HONEY</b> (UNII: Y9H1V576FH)
<b>INDIAN FRANKINCENSE</b> (UNII: 4PW41QCO2M)
<b>RESVERATROL</b> (UNII: Q369O8926L)
<b>LECITHIN, SOYBEAN</b> (UNII: 1DI56QDM62)
<b>SILVER</b> (UNII: 3M4G523W1G)
<b>FERRIC OXIDE RED</b> (UNII: 1K09F3G675)
<b>FERRIC OXIDE YELLOW</b> (UNII: EX438O2MRT)

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:54111-119-50	4 g in 1 JAR; Type 0: Not a Combination Product	08/15/2012	

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M020	08/15/2012	01/01/2025

## RODAN AND FIELDS ENHANCEMENTS MEDIUM

titanium dioxide, zinc oxide powder

**Product Information**

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:54111-120
<b>Route of Administration</b>	TOPICAL		

**Active Ingredient/Active Moiety**

<b>Ingredient Name</b>	<b>Basis of Strength</b>	<b>Strength</b>
<b>TITANIUM DIOXIDE</b> (UNII: 15FIX9V2JP) (TITANIUM DIOXIDE - UNII:15FIX9V2JP)	TITANIUM DIOXIDE	0.38400 g in 4 g
<b>ZINC OXIDE</b> (UNII: SOI2LOH54Z) (ZINC OXIDE - UNII:SOI2LOH54Z)	ZINC OXIDE	0.19600 g in 4 g

**Inactive Ingredients**

<b>Ingredient Name</b>	<b>Strength</b>
<b>MICA</b> (UNII: V8A1AW0880)	
<b>POLYMETHYLSILSESQUIOXANE (4.5 MICRONS)</b> (UNII: 59Z907ZB69)	
<b>SQUALANE</b> (UNII: GW89575KF9)	
<b>WATER</b> (UNII: 059QF0K00R)	
<b>SILICON DIOXIDE</b> (UNII: ETJ7Z6XBU4)	
<b>BISMUTH OXYCHLORIDE</b> (UNII: 4ZR792I587)	
<b>STEARIC ACID</b> (UNII: 4ELV7Z65AP)	
<b>ALUMINUM HYDROXIDE</b> (UNII: 5QB0T2IUN0)	
<b>CERAMIDE 3</b> (UNII: 4370DF050B)	
<b>CHOLESTEROL</b> (UNII: 97C5T2UQ7J)	
<b>PHYTOSPHINGOSINE</b> (UNII: GIN46U9Q2Q)	
<b>PHENOXYETHANOL</b> (UNII: HIE492ZZ3T)	
<b>BUTYLENE GLYCOL</b> (UNII: 3XUS85K0RA)	
<b>CAPRYLYL GLYCOL</b> (UNII: 00YIU5438U)	
<b>BARIUM SULFATE</b> (UNII: 25BB7EKE2E)	
<b>ANGELICA SINENSIS ROOT</b> (UNII: B66F4574UG)	
<b>HEXAMETHYLENE DIISOCYANATE/TRIMETHYLOL HEXYLLACTONE CROSSPOLYMER</b> (UNII: WB5K9Y35Y9)	
<b>POTASSIUM SORBATE</b> (UNII: 1VPU26JZZ4)	
<b>HEXYLENE GLYCOL</b> (UNII: KEH0A3F75J)	
<b>TRANEXAMIC ACID</b> (UNII: 6T84R30KC1)	
<b>TRIETHOXYCAPRYLYLSILANE</b> (UNII: LDC331P08E)	
<b>TRIMYRISTIN</b> (UNII: 18L31PSR28)	
<b>HYDROGENATED SOYBEAN LECITHIN</b> (UNII: H1109Z9J4N)	
<b>HONEY</b> (UNII: Y9H1V576FH)	
<b>INDIAN FRANKINCENSE</b> (UNII: 4PW41QCO2M)	
<b>RESVERATROL</b> (UNII: Q369O8926L)	
<b>LECITHIN, SOYBEAN</b> (UNII: 1DI56QDM62)	
<b>SILVER</b> (UNII: 3M4G523W1G)	
<b>FERRIC OXIDE RED</b> (UNII: 1K09F3G675)	
<b>FERRIC OXIDE YELLOW</b> (UNII: EX438O2MRT)	

**Packaging**

<b>#</b>	<b>Item Code</b>	<b>Package Description</b>	<b>Marketing Start Date</b>	<b>Marketing End Date</b>
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<b>1</b>	NDC:54111-120-50	4 g in 1 JAR; Type 0: Not a Combination Product	09/05/2012
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## Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M020	09/05/2012	

## RODAN AND FIELDS ENHANCEMENTS BRONZE

titanium dioxide, zinc oxide powder

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:54111-121
<b>Route of Administration</b>	TOPICAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>TITANIUM DIOXIDE</b> (UNII: 15FIX9V2JP) (TITANIUM DIOXIDE - UNII:15FIX9V2JP)	TITANIUM DIOXIDE	0.38400 g in 4 g
<b>ZINC OXIDE</b> (UNII: SOI2LOH54Z) (ZINC OXIDE - UNII:SOI2LOH54Z)	ZINC OXIDE	0.19600 g in 4 g

### Inactive Ingredients

Ingredient Name	Strength
<b>MICA</b> (UNII: V8A1AW0880)	
<b>POLYMETHYLSILSESQUIOXANE (4.5 MICRONS)</b> (UNII: 59Z907ZB69)	
<b>SQUALANE</b> (UNII: GW89575KF9)	
<b>WATER</b> (UNII: 059QF0KO0R)	
<b>SILICON DIOXIDE</b> (UNII: ETJ7Z6XBU4)	
<b>BISMUTH OXYCHLORIDE</b> (UNII: 4ZR792I587)	
<b>STEARIC ACID</b> (UNII: 4ELV7Z65AP)	
<b>ALUMINUM HYDROXIDE</b> (UNII: 5QB0T2IUN0)	
<b>CERAMIDE 3</b> (UNII: 4370DF050B)	
<b>CHOLESTEROL</b> (UNII: 97C5T2UQ7J)	
<b>PHYTOSPHINGOSINE</b> (UNII: GIN46U9Q2Q)	
<b>PHENOXYETHANOL</b> (UNII: HIE492ZZ3T)	
<b>BUTYLENE GLYCOL</b> (UNII: 3XUS85K0RA)	
<b>CAPRYLYL GLYCOL</b> (UNII: 00YIU5438U)	
<b>BARIUM SULFATE</b> (UNII: 25BB7EKE2E)	
<b>ANGELICA SINENSIS ROOT</b> (UNII: B66F4574UG)	
<b>HEXAMETHYLENE DIISOCYANATE/TRIMETHYLOL HEXYLLACTONE CROSSPOLYMER</b> (UNII: WB5K9Y35Y9)	
<b>POTASSIUM SORBATE</b> (UNII: 1VPU26JZZ4)	
<b>HEXYLENE GLYCOL</b> (UNII: KEH0A3F75J)	
<b>TRANEXAMIC ACID</b> (UNII: 6T84R30KC1)	
<b>TRIETHOXYCAPRYLYLSILANE</b> (UNII: LDC331P08E)	

<b>TRIMYRISTIN</b> (UNII: 18L31PSR28)	
<b>HYDROGENATED SOYBEAN LECITHIN</b> (UNII: H1109Z9J4N)	
<b>HONEY</b> (UNII: Y9H1V576FH)	
<b>INDIAN FRANKINCENSE</b> (UNII: 4PW41QCO2M)	
<b>RESVERATROL</b> (UNII: Q369O8926L)	
<b>LECITHIN, SOYBEAN</b> (UNII: 1DI56QDM62)	
<b>SILVER</b> (UNII: 3M4G523W1G)	
<b>FERRIC OXIDE RED</b> (UNII: 1K09F3G675)	
<b>FERRIC OXIDE YELLOW</b> (UNII: EX438O2MRT)	

<b>Packaging</b>				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:54111-121-50	4 g in 1 JAR; Type 0: Not a Combination Product	04/19/2013	01/01/2025

<b>Marketing Information</b>			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M020	04/19/2013	01/01/2025

**Labeler** - Bentley Laboratories, LLC (068351753)

**Registrant** - Bentley Laboratories, LLC (068351753)