

**H.E.B SOLUTIONS SUNSCREEN- ultra spf 50 continuous spray aerosol, spray**  
**H.E.B**

*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.*

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**H.E.B Solutions Sunscreen Ultra SPF 50 Continuous Spray**

**Active ingredients**

Avobenzone 3.0%, Homosalate 10.0%, Octisalate 5.0%, Octocrylene 4.0%, Oxybenzone 5.0%

**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**

**FLAMMABLE:**

- do not use near heat, flame or while smoking
- avoid long term storage above 104°F (40°C)

**Do not use** • on damaged or broken skin.

**When using this product** • keep out of eyes. Rinse with water to remove. • do not puncture or incinerate. Contents under pressure. • do not store at temperatures above 120°F.

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

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

**Directions**

- apply liberally 15 minutes before sun exposure
- hold can 4-6 inches away from body, spray evenly to ensure complete coverage
- do not spray into face. Spray into hand and apply to face.
- use in well ventilated, but not windy areas
- reapply:
  - after 80 minutes of swimming or sweating
  - immediately after towel drying
  - at least every 2 hours
  - 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 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

***Other information***

- protect this product from excessive heat and direct sun
- may stain or damage some fabrics, materials or surfaces

***Inactive ingredients***

SD alcohol 40-B, acrylates/octylacrylamide copolymer, menthyl lactate, aloe barbadensis leaf juice, tocopherol, camellia sinensis leaf extract (green tea), chamomilla recutita (matricaria) flower extract, glycerin, saccharomyces/podophyllum peltatum ferment filtrate, stearoxytrimethylsilane, caprylic/capric triglyceride, fragrance, butylene glycol, propylene glycol

***Questions or Comments?***

Call toll free 1-800-527-7731

H.E.B Solutions Sunscreen Ultra SPF 50 Continuous Spray

5.5 OZ (156g)

NDC 37808-974-37

Dermatologist tested  
 Oil Free • PABA Free • Rub-Free Formula  
 Hypoallergenic  
 Contains Aloe, Chamomile Extract,  
 and Green Tea Extract

**Drug Facts**

Active Ingredients	Purpose
Avobenzone 3%	Sunscreen
Homosalate 10%	
Octisalate 5%	
Octocrylene 4%	
Oxybenzone 3%	

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MADE WITH PRIDE  
 AND CARE FOR H-E-B®,  
 SAN ANTONIO, TEXAS 78204

PLEASE RECYCLE



SOLUTIONS™  
 SUNSCREEN

**ULTRA 50**  
 Ultra Sunscreen  
 UVA/UVB Protection  
 Water Resistant (80 minutes)  
 Paraben Free

**BROAD SPECTRUM SPF 50**



NET WT. 5.5 OZ (156 g)

Dermatologist tested  
 Oil Free • PABA Free • Rub-Free Formula  
 Hypoallergenic  
 Contains Aloe, Chamomile Extract,  
 and Green Tea Extract

<b>Drug Facts</b>	
<b>Active Ingredients</b>	<b>Purpose</b>
Avobenzone 3% Homosalate 10% Octisalate 5% Octocrylene 4% Oxybenzone 3%	Sunscreen
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 SAN ANTONIO, TEXAS 78204



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 (80 minutes)  
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## H.E.B SOLUTIONS SUNSCREEN

ultra spf 50 continuous spray aerosol, spray

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:378 08-974
<b>Route of Administration</b>	TOPICAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
AVOBENZONE (UNII: G63QQF2NOX) (AVOBENZONE - UNII:G63QQF2NOX)	AVOBENZONE	30 mg in 1 g
HOMOSALATE (UNII: V06SV4M95S) (HOMOSALATE - UNII:V06SV4M95S)	HOMOSALATE	100 mg in 1 g

<b>OCTISALATE</b> (UNII: 4X49Y0596W) (OCTISALATE - UNII:4X49Y0596W)	OCTISALATE	50 mg in 1 g
<b>OCTOCRYLENE</b> (UNII: 5A68WGF6WM) (OCTOCRYLENE - UNII:5A68WGF6WM)	OCTOCRYLENE	40 mg in 1 g
<b>OXYBENZONE</b> (UNII: 95OOS7VE0Y) (OXYBENZONE - UNII:95OOS7VE0Y)	OXYBENZONE	50 mg in 1 g

### Inactive Ingredients

Ingredient Name	Strength
<b>ALCOHOL</b> (UNII: 3K9958V90M)	
<b>ACRYLATE/ISOBUTYL METHACRYLATE/N-TERT-OCTYLACRYLAMIDE COPOLYMER (75000 MW)</b> (UNII: JU3XHR8VWK)	
<b>MENTHYL LACTATE, (-)</b> (UNII: 2BF9E65L7I)	
<b>ALOE VERA LEAF</b> (UNII: ZY81Z83H0X)	
<b>TOCOPHEROL</b> (UNII: R0ZB2556P8)	
<b>GLYCERIN</b> (UNII: PDC6A3C0OX)	
<b>STEAROXYTRIMETHYLSILANE</b> (UNII: 9862TW94B2)	
<b>MEDIUM-CHAIN TRIGLYCERIDES</b> (UNII: C9H2L21V7U)	
<b>BUTYLENE GLYCOL</b> (UNII: 3XUS85K0RA)	
<b>GREEN TEA LEAF</b> (UNII: W2ZU1RY8B0)	
<b>CHAMOMILE</b> (UNII: FGL3685T2X)	
<b>PROPYLENE GLYCOL</b> (UNII: 6DC9Q167V3)	

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:37808-974-37	156 g in 1 CAN; Type 0: Not a Combination Product	02/17/2012	

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part352	02/17/2012	

**Labeler** - H.E.B (007924756)

**Registrant** - Fruit of the Earth, Inc. (079559467)

### Establishment

Name	Address	ID/FEI	Business Operations
Fruit of the Earth, Inc.		008193513	manufacture(37808-974)