

**SIGNATURE CARE SHEER FACE AND BODY SPF 70 SUNSCREEN- avobenzone, homosalate, octisalate, octocrylene, oxybenzone stick
SAFEWAY, 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.

SIGNATURE CARE SHEER FACE AND BODY SPF 70 SUNSCREEN STICK

Active ingredients

Avobenzone 3%,
Homosalate 15%,
Octisalate 5%,
Octocrylene 10%,
Oxybenzone 3%

Purpose

Sunscreen

• 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 swallowed, get medical help or contact a Poison Control Center right away.

Directions

- apply liberally 15 minutes before sun exposure
- 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

adipic acid/diglycol crosspolymer, beeswax, butyloctyl salicylate, C12-15 alkyl benzoate, caprylyl glycol, dimethicone, fragrance, neopentyl

glycol diethylhexanoate, neopentyl glycol diisostearate, octyldodecyl neopentanoate, ozokerite, paraffin, polyethylene, silica

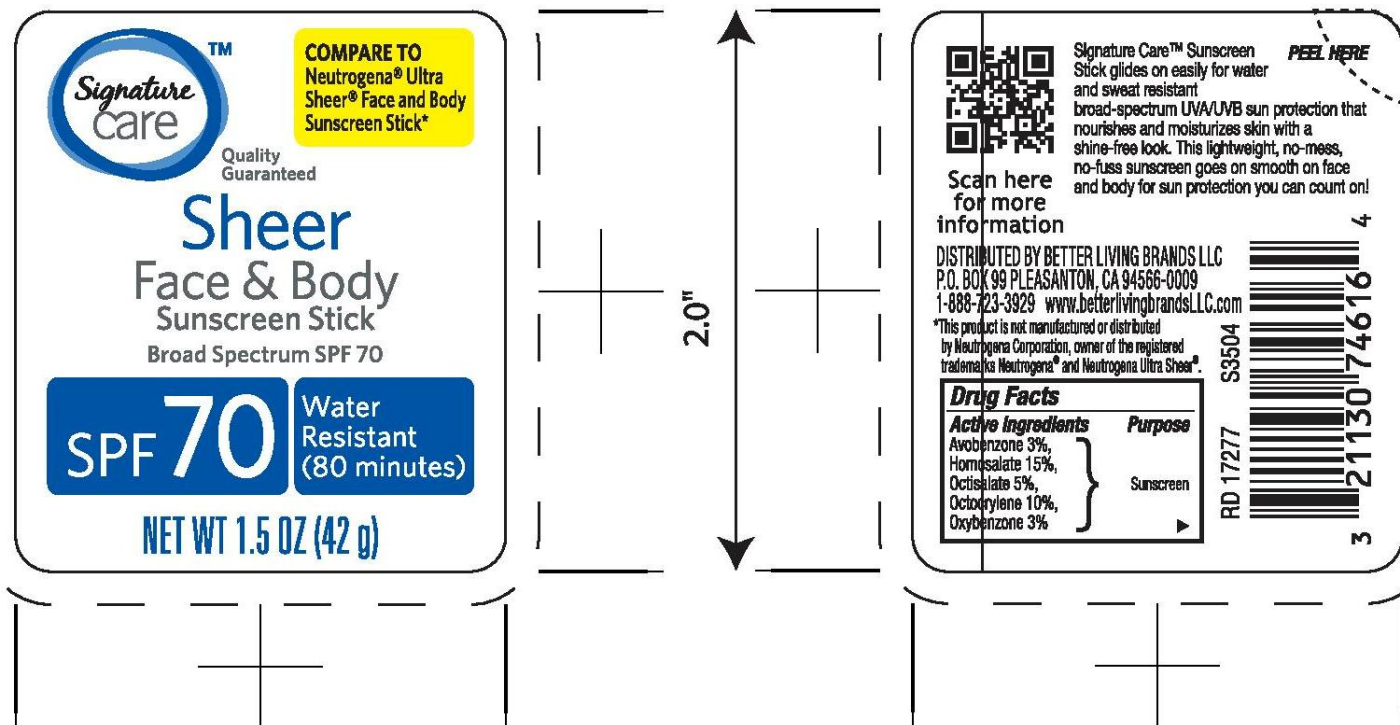
Questions or comments?

Call toll free 1-888-723-3929

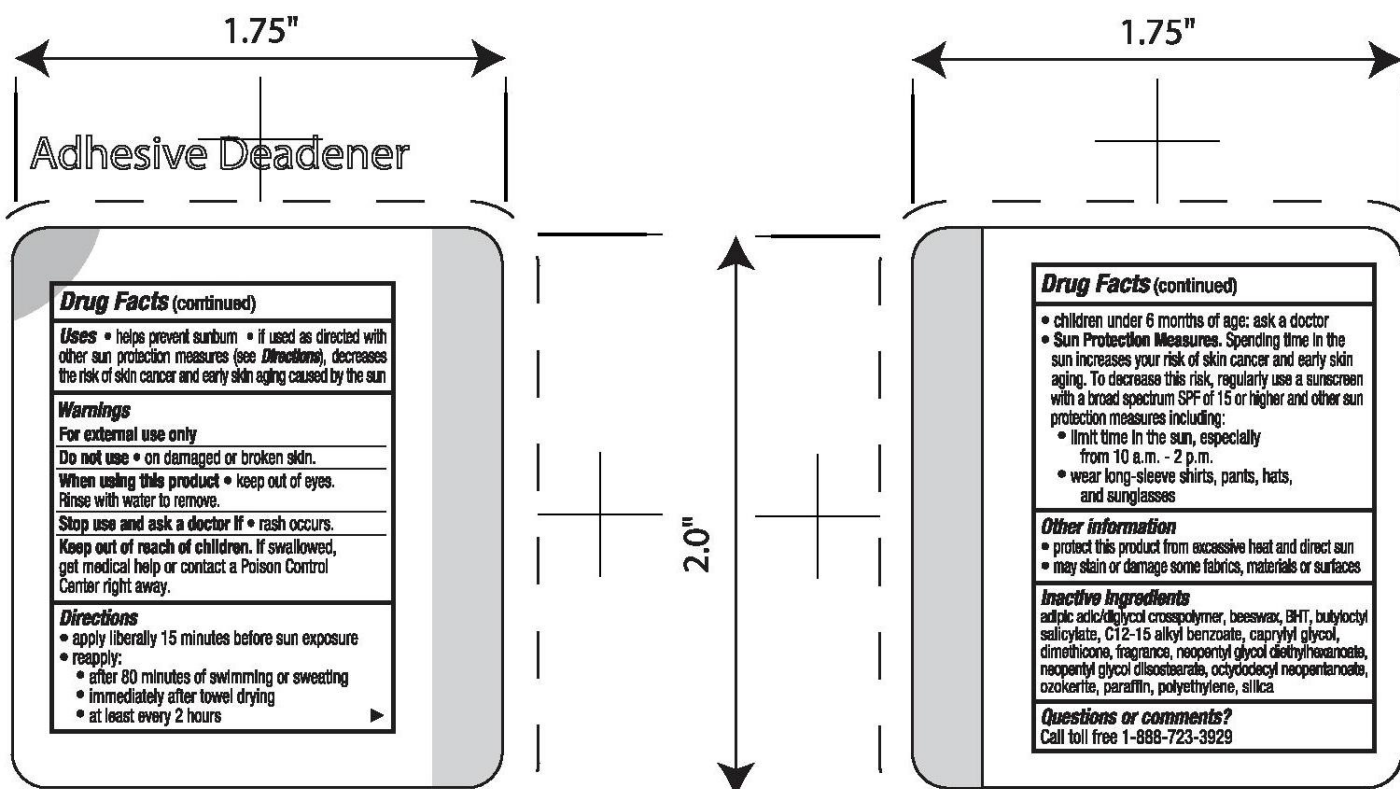
SIGNATURE CARE SHEER FACE AND BODY SPF 70 SUNSCREEN STICK

1.5 OZ. (42g)

NDC 21130-721-36



Ply 1 Front
3/32" from die edges



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avobenzone, homosalate, octisalate, octocrylene, oxybenzone stick

| Product Information | | | | |
|--|------------------|---|----------------------|--------------------|
| Product Type | | HUMAN OTC DRUG | Item Code (Source) | NDC:21130-721 |
| Route of Administration | | 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 | 150 mg in 1 g |
| OCTISALATE (UNII: 4X49 Y0596 W) (OCTISALATE - UNII:4X49 Y0596 W) | | | OCTISALATE | 50 mg in 1 g |
| OCTOCRYLENE (UNII: 5A68 WGF6 WM) (OCTOCRYLENE - UNII:5A68 WGF6 WM) | | | OCTOCRYLENE | 100 mg in 1 g |
| OXYBENZONE (UNII: 95OOS7VE0 Y) (OXYBENZONE - UNII:95OOS7VE0 Y) | | | OXYBENZONE | 30 mg in 1 g |
| | | | | |
| Inactive Ingredients | | | | |
| Ingredient Name | | | | Strength |
| HIGH DENSITY POLYETHYLENE (UNII: UG00KM4WR7) | | | | |
| CERESIN (UNII: Q1LS2UJO3A) | | | | |
| ADIPIC ACID/DIGLYCOL CROSSPOLYMER (20000 MPA.S) (UNII: R9TPS68K19) | | | | |
| BUTYLOCTYL SALICYLATE (UNII: 2EH13UN8D3) | | | | |
| ALKYL (C12-15) BENZOATE (UNII: A9EJ3J61HQ) | | | | |
| CAPRYLYL GLYCOL (UNII: 00YIU5438U) | | | | |
| DIMETHICONE (UNII: 92RU3N3Y1O) | | | | |
| NEOPENTYL GLYCOL DIETHYLHEXANOATE (UNII: U68ZV6W62C) | | | | |
| NEOPENTYL GLYCOL DIISOSTEARATE (UNII: 4M6OQ34JWW) | | | | |
| OCTYLDODECYL NEOPENTANOATE (UNII: X8725R883T) | | | | |
| PARAFFIN (UNII: I9O0E3H2ZE) | | | | |
| SILICON DIOXIDE (UNII: ETJ7Z6XBU4) | | | | |
| YELLOW WAX (UNII: 2ZA36H0S2V) | | | | |
| | | | | |
| Packaging | | | | |
| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
| 1 | NDC:21130-721-36 | 42 g in 1 TUBE; Type 0: Not a Combination Product | 11/13/2017 | |
| | | | | |
| Marketing Information | | | | |
| Marketing Category | | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
| OTC monograph not final | | part352 | 11/13/2017 | |

Labeler - SAFEWAY, INC. (009137209)

Registrant - FRUIT OF THE EARTH, INC. (079559467)

Establishment

| Name | Address | ID/FEI | Business Operations |
|--|---------|-----------|------------------------|
| Fruit Of The Earth Research Laboratories, Inc. | | 008193513 | manufacture(21130-721) |

Revised: 5/2020

SAFEWAY, INC.