

PHELITYL DAY SPF 15- phelityl day spf 15 cream
MANA PRODUCTS, INC.

PHELITYL DAY SPF 15

ACTIVE INGREDIENTS

Avobenzone 2%,
Octinoxate 5%

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.

Directions

- Apply liberally 15 minutes before sun exposure
- reapply: after 40 minutes of swimming or sweating
- immediately after towel drying
- 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

Inactive Ingredients

Water, Glyceryl Stearate, Butylene Glycol, Cetyl Alcohol, Cyclopentasiloxane, C12-15 Alkyl Benzoate, Octyldodecyl Neopentanoate, Silica, Butyrospermum Parkii (Shea) Butter, Emu Oil, Glycerin, PEG-100 Stearate, Polyglyceryl-3 Laurate, Taraktogenos Kurzii Seed Oil, Aloe Barbadensis Leaf Extract, Anthemis Nobilis Flower Extract, Camellia

Sinensis Leaf Extract, Panax Ginseng Root Extract, Allantoin, Carbomer, Ceteth-20, Disodium EDTA, Ethylhexylglycerin, O-Cymen-5-ol, Tetrahexyldecyl Ascorbate, Tocopheryl Acetate, Fragrance, Steareth-2, Steareth-20, Triethanolamine, Phenoxyethanol.

Other information

- protect this product from excessive heat and direct sun

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.

Product label



PHELITYL DAY SPF 15

pHELITYL day spf 15 cream?

Product Information

| | | | |
|--------------------------------|----------------|---------------------------|---------------|
| Product Type | HUMAN OTC DRUG | Item Code (Source) | NDC:59735-475 |
| Route of Administration | TOPICAL | | |

Active Ingredient/Active Moiety

| Ingredient Name | Basis of Strength | Strength |
|---|-------------------|--------------|
| OCTINOXATE (UNII: 4Y5P7MUD51) (OCTINOXATE - UNII:4Y5P7MUD51) | OCTINOXATE | 5 g in 100 g |
| AVOBENZONE (UNII: G63QQF2NOX) (AVOBENZONE - UNII:G63QQF2NOX) | AVOBENZONE | 2 g in 100 g |

Inactive Ingredients

| Ingredient Name | Strength |
|---|----------|
| CETYL ALCOHOL (UNII: 936JST6JCN) | |
| CYCLOMETHICONE 5 (UNII: 0THT5PCI0R) | |
| SILICON DIOXIDE (UNII: ETJ7Z6XBU4) | |
| SHEA BUTTER (UNII: K49155WL9Y) | |
| EMU OIL (UNII: 344821WD61) | |
| GLYCERIN (UNII: PDC6A3C0OX) | |
| PEG-100 STEARATE (UNII: YD01N1999R) | |
| MYRISTYL ALCOHOL (UNII: V42034O9PU) | |
| COCO GLUCOSIDE (UNII: ICS790225B) | |
| ETHYLHEXYLGLYCERIN (UNII: 147D247K3P) | |
| CETETH-20 (UNII: I835H2IHHX) | |
| WATER (UNII: 059QF0KO0R) | |
| GLYCERYL MONOSTEARATE (UNII: 230OU9XXE4) | |
| BUTYLENE GLYCOL (UNII: 3XUS85K0RA) | |
| ALKYL (C12-15) BENZOATE (UNII: A9EJ3J61HQ) | |
| OCTYLDODECYL NEOPENTANOATE (UNII: X8725R883T) | |
| DIMETHICONE (UNII: 92RU3N3Y1O) | |
| HYDNOCARPUS KURZII SEED OIL (UNII: N757YEZ18Q) | |
| ALOE VERA LEAF (UNII: ZY81Z83H0X) | |
| CHAMAEMELUM NOBILE FLOWER (UNII: O2T154T6OG) | |
| GREEN TEA LEAF (UNII: W2ZU1RY8B0) | |
| ASIAN GINSENG (UNII: CUQ3A77YXI) | |
| ALLANTOIN (UNII: 344S277G0Z) | |
| EDETATE DISODIUM (UNII: 7FLD91C86K) | |
| STEARETH-20 (UNII: L0Q8IK9E08) | |
| TROLAMINE (UNII: 9O3K93S3TK) | |
| PHENOXYETHANOL (UNII: HIE492ZZ3T) | |
| O-CYMEN-5-OL (UNII: H41B6Q1I9L) | |
| TETRAHEXYLDECYL ASCORBATE (UNII: 9LBV3F07AZ) | |
| .ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0) | |
| STEARETH-2 (UNII: V56DFE46J5) | |

Product Characteristics

| | | | |
|-----------------|-------|---------------------|--|
| Color | white | Score | |
| Shape | | Size | |
| Flavor | | Imprint Code | |
| Contains | | | |

Packaging

| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
|---|------------------|--|----------------------|--------------------|
| 1 | NDC:59735-475-01 | 1 in 1 CARTON | 04/10/2013 | |
| 1 | | 90 q in 1 JAR; Type 0: Not a Combination | | |

| Product | | | |
|------------------------------|--|----------------------|--------------------|
| Marketing Information | | | |
| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
| OTC Monograph Drug | M020 | 04/10/2013 | |

Labeler - MANA PRODUCTS, INC. (078870292)

| Establishment | | | |
|----------------------|---------|-----------|------------------------|
| Name | Address | ID/FEI | Business Operations |
| MANA PRODUCTS, INC | | 032870270 | manufacture(59735-475) |

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| MANA PRODUCTS, INC. | | 078870292 | manufacture(59735-475) |

Revised: 1/2024

MANA PRODUCTS, INC.