

**MICRO DAY REJUVENATING BROAD-SPECTRUM SUNSCREEN SPF 30-
octinoxate and zinc oxide lotion
Merz North America, 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.

Micro Day Rejuvenating Cream Broad-spectrum Sunscreen SPF 30

Drug Facts

Active ingredients

Octinoxate 7.5%

Zinc Oxide 7.3%

Purpose

Sunscreen

Uses

- Helps prevent sunburn and premature skin aging.
- 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.
- Higher SPF gives more sunburn protection.

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

See package insert for complete information.

For sunscreen use:

- Apply generously 15 minutes before sun exposure.
- Reapply at least every 2 hours.
- Use a water-resistant sunscreen if swimming or sweating.

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 Broad-Spectrum SPF value of 15 or higher and other sun protection measures including:

- limit time in the sun, especially from 10am - 2pm
- Wear long-sleeved shirts, pants, hats, and sunglasses.
- Children under 6 months of age, ask a doctor.

Inactive ingredients

Water, Caprylic/Capric Triglyceride, Hydrogenated C6-14 Olefin Polymers, Hexyldecanol, Glycerin, Glyceryl Stearate, PEG-100 Stearate, Potassium Cetyl Phosphate, Hydrogenated Palm Glycerides, Microcrystalline Cellulose, Camelia Sinensis Extract, Silica, Tetrapeptide-21, Capryloyl Carnosine, Palmitoyl Tripeptide-1 Acetate, Sodium Ascorbyl Phosphate, Tocopheryl Acetate, Squalane, Hydroxyethyl Acrylate/Sodium Acryloyldimethyl Taurate Copolymer, Polyhydroxystearic Acid, Steareth-21, Melanin, Cetearyl Alcohol, Sodium Hyaluronate, Polysorbate 60, Triethoxycaprylsilane, Disodium EDTA, Xanthan Gum, Cellulose Gum, Styrene/Acrylates Copolymer, Phenoxyethanol, Methylparaben, Ethylparaben, Propylparaben.

Other information

- Protect this product from excessive heat and direct sun.

Questions?

USA: 866.636.2884 • International: +800.1489.1489

Distributed in the USA by Merz North America, Inc.

PRINCIPAL DISPLAY PANEL - 50 ML Bottle Carton

micro day®

REVITALIZING
& TIGHTENING
DAY CREAM
SPF 30

NEOCUTIS®

micro day®

Drug Facts

| Active Ingredients | Purpose |
|--------------------------------------|-----------|
| Octinoxate 7.5% Zinc Oxide 7.3% } | Sunscreen |

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REVITALIZING
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Other Information

- Protect this product from excessive heat and direct sun.

Caution: For external use only. Keep out of eyes. If contact with eyes occurs, rinse eyes thoroughly with water. Discontinue use if signs of irritation or rash appear. Keep out of reach of children.

**Non-comedogenic, dermatologist-tested,
free of color additives and fragrance,
not tested on animals**

Distributed in the USA
by Merz North America, Inc.
Raleigh, NC 27615 | Made in USA

NEOCUTIS.com

USA: 866.636.2884

1.69 FL. OZ. (50 ML) | AWO0878-01

Restores elastin and
collagen to revitalize and
hydrate skin, provides
anti-oxidant care and
broad-spectrum UVA
and UVB protection

POWERED BY
PROPRIETARY PEPTIDES



Apply to face, neck and
décolleté, or as directed by
your skincare professional.

MICRO DAY REJUVENATING BROAD-SPECTRUM SUNSCREEN SPF 30

octinoxate and zinc oxide lotion

Product Information

| | | | |
|-------------------------|----------------|--------------------|---------------|
| Product Type | HUMAN OTC DRUG | Item Code (Source) | NDC:46783-173 |
| Route of Administration | TOPICAL | | |

Active Ingredient/Active Moiety

| Ingredient Name | Basis of Strength | Strength |
|--|-------------------|---------------|
| OCTINOXATE (UNII: 4Y5P7MUD51) (OCTINOXATE - UNII:4Y5P7MUD51) | OCTINOXATE | 75 mg in 1 mL |
| ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC OXIDE - UNII:SOI2LOH54Z) | ZINC OXIDE | 73 mg in 1 mL |

Inactive Ingredients

| Ingredient Name | Strength |
|---|----------|
| WATER (UNII: 059QF0KO0R) | |
| MEDIUM-CHAIN TRIGLYCERIDES (UNII: C9H2L21V7U) | |

| | |
|--|--|
| HEXYLDECANOL (UNII: 151Z7P1317) | |
| GLYCERIN (UNII: PDC6A3C0OX) | |
| GLYCERYL MONOSTEARATE (UNII: 230OU9XXE4) | |
| PEG-100 STEARATE (UNII: YD01N1999R) | |
| POTASSIUM CETYL PHOSPHATE (UNII: 03KCY6P7UT) | |
| HYDROGENATED PALM GLYCERIDES (UNII: YCZ8EM144Q) | |
| MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U) | |
| GREEN TEA LEAF (UNII: W2ZU1RY8B0) | |
| XANTHAN GUM (UNII: TTV12P4NEE) | |
| ETHYLPARABEN (UNII: 14255EXE39) | |
| PROPYLPARABEN (UNII: Z8IX2SC1OH) | |
| METHYLPARABEN (UNII: A2I8C7HI9T) | |
| TRIETHOXYCAPRYLYLSILANE (UNII: LDC331P08E) | |
| CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S) | |
| STEARETH-21 (UNII: 53J3F32P58) | |
| .ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0) | |
| HYALURONATE SODIUM (UNII: YSE9PPT4TH) | |
| SODIUM ASCORBYL PHOSPHATE (UNII: 836SJG51DR) | |
| EDETATE DISODIUM (UNII: 7FLD91C86K) | |
| POLYHYDROXYSTEARIC ACID (2300 MW) (UNII: YXH47AOU0F) | |
| HYDROGENATED POLY(C6-14 OLEFIN; 2 CST) (UNII: POTX083987) | |

Product Characteristics

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

Packaging

| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
|---|------------------|---|----------------------|--------------------|
| 1 | NDC:46783-173-05 | 1 in 1 CARTON | 03/23/2015 | 02/28/2023 |
| 1 | | 15 mL in 1 BOTTLE; Type 0: Not a Combination Product | | |
| 2 | NDC:46783-173-10 | 1 in 1 CARTON | 03/23/2015 | 12/31/2022 |
| 2 | | 30 mL in 1 BOTTLE; Type 0: Not a Combination Product | | |
| 3 | NDC:46783-173-20 | 1 in 1 CARTON | 03/23/2015 | 12/31/2022 |
| 3 | | 200 mL in 1 BOTTLE; Type 0: Not a Combination Product | | |
| 4 | NDC:46783-173-04 | 4 mL in 1 TUBE; Type 0: Not a Combination Product | 06/15/2019 | 10/31/2024 |
| 5 | NDC:46783-173-50 | 1 in 1 CARTON | 10/20/2019 | 12/31/2023 |
| 5 | | 50 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 | 03/23/2015 | 10/31/2024 |

Labeler - Merz North America, Inc. (028147846)

Revised: 6/2023

Merz North America, Inc.