

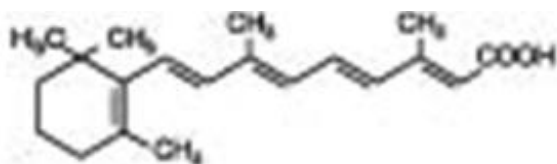
RENOVA- tretinoin cream
Rebel Distributors Corp

RENOVA®
(TRETINOIN CREAM)
0.02%

FOR TOPICAL USE ON THE FACE. NOT FOR OPHTHALMIC, ORAL OR INTRAVAGINAL USE.

DESCRIPTION

RENOVA® (tretinoin cream) 0.02% contains the active ingredient tretinoin in a cream base. Tretinoin is a yellow- to light-orange crystalline powder having a characteristic floral odor. Tretinoin is soluble in dimethylsulfoxide, slightly soluble in polyethylene glycol 400, octanol, and 100% ethanol. It is practically insoluble in water and mineral oil, and it is insoluble in glycerin. The chemical name for tretinoin is (all-E)-3,7-dimethyl-9-(2,6,6-trimethyl-1-cyclononen-1-yl)-2,4,6,8-nonatetraenoic acid. Tretinoin is also referred to as all-trans-retinoic acid and has a molecular weight of 300.44. The structural formula is represented below.



TRETINOIN

Tretinoin is available as RENOVA® at a concentration of 0.02% w/w in an oil-in-water emulsion formulation consisting of benzyl alcohol, butylated hydroxytoluene, caprylic/capric triglyceride, cetyl alcohol, edetate disodium, fragrance, methylparaben, propylparaben, purified water, stearic acid, stearyl alcohol, steareth 2, steareth 20, and xanthan gum.

CLINICAL PHARMACOLOGY

Tretinoin is an endogenous retinoid metabolite of Vitamin A that binds to intracellular receptors in the cytosol and nucleus, but cutaneous levels of tretinoin in excess of physiologic concentrations occur following application of a tretinoin-containing topical drug product. Although tretinoin activates three members of the retinoic acid (RAR) nuclear receptors (RAR α , RAR β , and RAR γ) which may act to modify gene expression, subsequent protein synthesis, and epithelial cell growth and differentiation, it has not been established whether the clinical effects of tretinoin are mediated through activation of retinoic acid receptors, other mechanisms such as irritation, or both.

The effect of tretinoin on skin with chronic photodamage has not been evaluated in animal studies. When hairless albino mice were treated topically with tretinoin shortly after a period of UVB irradiation, new collagen formation was demonstrated only in photodamaged skin. However, in human skin treated topically, adequate data have not been provided to demonstrate any increase in desmosine, hydroxyproline, or elastin mRNA. Application of 0.1% tretinoin cream to photodamaged human forearm skin was associated with an increase in antibody staining for procollagen I propeptide. No correlation was made between procollagen I propeptide staining with collagen I levels or with observed clinical effects. Thus, the relationships between the increased collagen in rodents, increased procollagen I propeptide in humans, and the clinical effects of tretinoin have not yet been clearly defined.

Tretinoin was shown to enhance UV-stimulated melanogenesis in pigmented mice. Generalized amyloid

deposition in the basal layer of tretinoin-treated skin was noted in a two-year mouse study. In a different study, hyalinization at tretinoin-treated skin sites was noted at doses beginning at 0.25 mg/kg in CD-1 mice.

The transdermal absorption of tretinoin from various topical formulations ranged from 1% to 31% of applied dose, depending on whether it was applied to healthy skin or dermatitic skin. No percutaneous absorption study was conducted with RENOVA[®] (tretinoin cream) 0.02% in human volunteers. When percutaneous absorption of the oil-in-water emulsion formulation at 0.05% concentration was assessed in healthy male subjects with radiolabeled cream after a single application (n=7), as well as after repeated daily applications (n=7) for 28 days, the absorption of tretinoin was less than 2% and the extent of bioavailability was less after repeated application. No significant difference in endogenous concentrations of tretinoin was observed between single and repeated daily applications.

INDICATIONS AND USAGE

(To understand fully the indication for this product, please read the entire INDICATIONS AND USAGE section of the labeling.)

RENOVA[®] (tretinoin cream) 0.02% is indicated as an adjunctive agent (see second bullet point below) for use in the mitigation (palliation) of fine facial wrinkles in patients who use comprehensive skin care and sunlight avoidance programs. **RENOVA[®] (tretinoin cream) 0.02% DOES NOT ELIMINATE WRINKLES, REPAIR SUN-DAMAGED SKIN, REVERSE PHOTOAGING, or RESTORE MORE YOUTHFUL or YOUNGER SKIN.** In double-blinded, vehicle-controlled clinical studies, many patients in the vehicle group achieved desired palliative effects on fine wrinkling of facial skin with the use of comprehensive skin care and sunlight avoidance programs including sunscreens, protective clothing, and non-prescription emollient creams.

- RENOVA[®] (tretinoin cream) 0.02% has NOT DEMONSTRATED A MITIGATING EFFECT on significant signs of chronic sunlight exposure such as coarse or deep wrinkling, tactile roughness, mottled hyperpigmentation, lentigines, telangiectasia, skin laxity, keratinocytic atypia, melanocytic atypia, or dermal elastosis.
- RENOVA[®] (tretinoin cream) 0.02% should be used under medical supervision as an adjunct to a comprehensive skin care and sunlight avoidance program that includes the use of effective sunscreens (minimum SPF of 15) and protective clothing.
- Patients with visible actinic keratoses and patients with a history of skin cancer were excluded from clinical trials of RENOVA[®] (tretinoin cream) 0.02%. Thus the effectiveness and safety of RENOVA[®] (tretinoin cream) 0.02% in these populations are not known at this time.
- Neither the safety nor the effectiveness of RENOVA[®] (tretinoin cream) 0.02% for the prevention or treatment of actinic keratoses or skin neoplasms has been established.
- Neither the safety nor the efficacy of RENOVA[®] (tretinoin cream) 0.02% daily for greater than 52 weeks has been established, and daily use beyond 52 weeks has not been systematically and histologically investigated in adequate and well-controlled trials. (See **WARNINGS** section.)

CLINICAL TRIALS

Four adequate and well-controlled multi-center trials and one single-center randomized, controlled trial were conducted involving a total of 324 evaluable patients treated with RENOVA[®] (tretinoin cream) 0.02% and 332 evaluable patients treated with the vehicle cream on the face for 24 weeks with a comprehensive skin care and sun avoidance program, to assess the effects on fine and coarse wrinkling, mottled hyperpigmentation, tactile skin roughness, and laxity. Patients were evaluated at baseline on a 10 unit scale and changes from the baseline rating were categorized as follows:

Worsening:	Increase of 1 unit or more.
No improvement:	No change.
Minimal improvement:	Reduction of 1 unit.

Mild improvement: Reduction of 2 units.
 Moderate improvement: Reduction of 3 units or more.

In these trials, the fine and coarse wrinkling, mottled hyperpigmentation, tactile roughness, and laxity of the facial skin were thought to be caused by multiple factors which included intrinsic aging or environmental factors, such as chronic sunlight exposure.

Two of the five trials provided adequate demonstration of efficacy for mitigation of fine facial wrinkling. No two of the five trials adequately demonstrated efficacy for mitigation of coarse wrinkling, mottled hyperpigmentation, tactile skin roughness, and laxity. Data for fine wrinkling (the indication for which RENOVA[®] (tretinoin cream) 0.02% demonstrated efficacy) from all five trials (four studies in lightly pigmented subjects with Fitzpatrick Skin Types I-III and one study in darkly pigmented subjects with Fitzpatrick Skin Types IV-VI) is provided below:

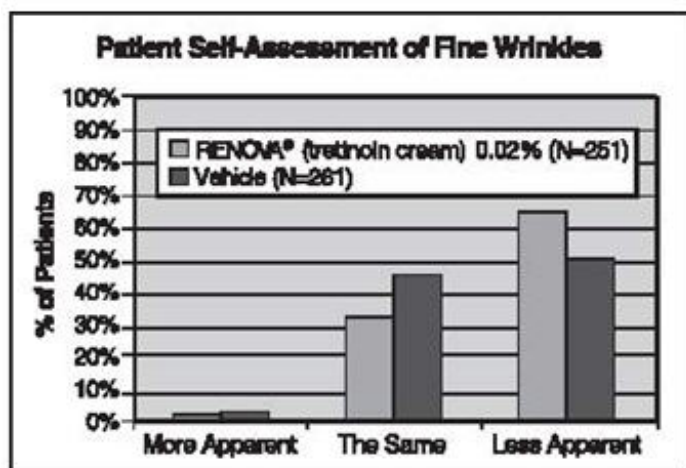
FINE WRINKLING IN LIGHTLY PIGMENTED SUBJECTS

	Subjects using RENOVA [®] (tretinoin cream) 0.02% + CSP* (N=279)	Vehicle + CSP* (N=280)
Worsened	1%	3%
No Change	40%	58%
Minimal Improvement	35%	27%
Mild Improvement	15%	9%
Moderate Improvement	10%	3%

A single-center study (N=107) in darkly pigmented, mostly African-American, subjects with Fitzpatrick Skin Types IV-VI demonstrated minimal or mild improvement in fine facial wrinkling in 43% of patients using Vehicle + CSP* compared to 29% of subjects using RENOVA[®] (tretinoin cream) 0.02% + CSP*. Although fewer darkly pigmented subjects improved with RENOVA[®] (tretinoin cream) 0.02% than with vehicle, these findings may reflect the small size of this study.

* CSP = Comprehensive skin protection and sunlight avoidance programs including use of sunscreens, protective clothing, and non-prescription emollient creams.

Self-assessment of fine wrinkles after 24 weeks of treatment with either RENOVA[®] (tretinoin cream) 0.02% or Vehicle from the four studies in lightly pigmented patients showed the following:



No studies have been conducted comparing the facial irritation or efficacy of RENOVA[®] (tretinoin cream) 0.02% to RENOVA[®] (tretinoin cream) 0.05% (older marketed formulation).

Patients may lose some of the mitigating effects of RENOVA[®] (tretinoin cream) 0.02% after 12 weeks of discontinuation of RENOVA[®] (tretinoin cream) 0.02% from their comprehensive skin care and

sunlight avoidance program.

CONTRAINDICATIONS

This drug is contraindicated in individuals with a history of sensitivity reactions to any of its components. It should be discontinued if hypersensitivity to any of its ingredients is noted.

WARNINGS

- RENOVA[®] (tretinoin cream) 0.02% is a dermal irritant, and the results of continued irritation of the skin for greater than 52 weeks in chronic use with RENOVA[®] (tretinoin cream) 0.02% are not known. There is evidence of atypical changes in melanocytes and keratinocytes and of increased dermal elastosis in some patients treated with RENOVA[®] (tretinoin cream) 0.05% for longer than 48 weeks. The significance of these findings and their relevance for RENOVA[®] (tretinoin cream) 0.02% are unknown.
- RENOVA[®] (tretinoin cream) 0.02% should not be administered if the patient is also taking drugs known to be photosensitizers (e.g., thiazides, tetracyclines, fluoroquinolones, phenothiazines, sulfonamides) because of the possibility of augmented phototoxicity.

Exposure to sunlight (including sunlamps) should be avoided or minimized during use of RENOVA[®] (tretinoin cream) 0.02% because of heightened sunburn susceptibility. Patients should be warned to use sunscreens (minimum SPF of 15) and protective clothing when using RENOVA[®] (tretinoin cream) 0.02%. Patients with sunburn should be advised not to use RENOVA[®] (tretinoin cream) 0.02% until fully recovered. Patients who may have considerable sun exposure, e.g., due to their occupation, and those patients with inherent sensitivity to sunlight should exercise caution when using RENOVA[®] (tretinoin cream) 0.02% and follow the precautions outlined in the Patient Package Insert.

RENOVA[®] (tretinoin cream) 0.02% should be kept out of the eyes, mouth, angles of the nose, and mucous membranes. Topical use may cause severe local erythema, pruritus, burning, stinging, and peeling at the site of application. If the degree of local irritation warrants, patients should be directed to use less medication, decrease the frequency of application, discontinue use temporarily, or discontinue use altogether and consider additional appropriate therapy.

Tretinoin has been reported to cause severe irritation on eczematous skin and should be used only with caution in patients with this condition.

Application of larger amounts of medication than recommended has not been shown to lead to more rapid or better results, and marked redness, peeling, or discomfort may occur.

PRECAUTIONS

General

RENOVA[®] (tretinoin cream) 0.02% should be used only as an adjunct to a comprehensive skin care and sunlight avoidance program. (See **INDICATIONS AND USAGE** section.)

If a drug sensitivity, chemical irritation, or a systemic adverse reaction develops, use of RENOVA[®] (tretinoin cream) 0.02% should be discontinued. Weather extremes, such as wind or cold, may be more irritating to patients using tretinoin-containing products.

Information for Patients

RENOVA[®] (tretinoin cream) 0.02% is to be used as described below unless otherwise directed by your physician:

1. It is for use on the face.
2. Avoid contact with the eyes, ears, nostrils, angles of the nose, and mouth. RENOVA[®] (tretinoin

cream) 0.02% may cause severe redness, itching, burning, stinging, and peeling if used on these areas.

3. In the evening, gently wash your face with a mild soap. Pat skin dry and wait 20-30 minutes before applying RENOVA[®] (tretinoin cream) 0.02%. Apply only a small pearl-sized (about ¼ inch or 5 millimeter diameter) amount of RENOVA[®] (tretinoin cream) 0.02% to your face at one time. This should be enough to cover the entire affected area lightly.
4. Do not wash your face for at least one hour after applying RENOVA[®] (tretinoin cream) 0.02%.
5. For best results, you are advised not to apply another skin care product or cosmetic for at least one hour after applying RENOVA[®] (tretinoin cream) 0.02%.
6. In the morning, apply a moisturizing sunscreen, SPF 15 or greater.
7. RENOVA[®] (tretinoin cream) 0.02% is a serious medication. Do not use RENOVA[®] (tretinoin cream) 0.02% if you are pregnant or attempting to become pregnant. If you become pregnant while using RENOVA[®] (tretinoin cream) 0.02%, please contact your physician immediately.
8. Avoid sunlight and other medicines that may increase your sensitivity to sunlight.
9. RENOVA[®] (tretinoin cream) 0.02% does not remove wrinkles or repair sun-damaged skin.

Please refer to the Patient Package Insert for additional patient information.

Drug Interactions

Concomitant topical medications, medicated or abrasive soaps, shampoos, cleansers, cosmetic with a strong drying effect, products with high concentrations of alcohol, astringents, spices or lime permanent wave solutions, electrolysis, hair depilatories or waxes, and products that may irritate the skin should be used with caution in patients being treated with RENOVA[®] (tretinoin cream) 0.02% because they may increase irritation with RENOVA[®] (tretinoin cream) 0.02%.

RENOVA[®] (tretinoin cream) 0.02% should not be administered if the patient is also taking drugs known to be photosensitizers (e.g., thiazides, tetracyclines, fluoroquinolones, phenothiazines, sulfonamides) because of the possibility of augmented phototoxicity.

Carcinogenesis, Mutagenesis, Impairment of Fertility

In a 91-week dermal study in which CD-1 mice were administered 0.017% and 0.035% formulations of tretinoin, cutaneous squamous cell carcinomas and papillomas in the treatment area were observed in some female mice. These concentrations are near the tretinoin concentration of this clinical formulation (0.02%). A dose-related incidence of liver tumors in male mice was observed at those same doses. The maximum systemic doses associated with the 0.017% and 0.035% formulations are 0.5 and 1.0 mg/kg/day. These doses are 10 and 20 times the maximum human systemic dose, when adjusted for total body surface area. The biological significance of these findings is not clear because they occurred at doses that exceeded the dermal maximally tolerated dose (MTD) of tretinoin and because they were within the background natural occurrence rate for these tumors in this strain of mice. There was no evidence of carcinogenic potential when 0.025 mg/kg/day of tretinoin was administered topically to mice (0.5 times the maximum human systemic dose, adjusted for total body surface area). For purposes of comparisons of the animal exposure to systemic human exposure, the maximum human systemic dose is defined as 1 gram of 0.02% RENOVA[®] (tretinoin cream) 0.02% applied daily to a 50 kg person (0.004 mg tretinoin/kg body weight).

Studies in hairless albino mice suggest that concurrent exposure to tretinoin may enhance the tumorigenic potential of carcinogenic doses of UVB and UVA light from a solar simulator. This effect has been confirmed in a later study in pigmented mice, and dark pigmentation did not overcome the enhancement of photocarcinogenesis by 0.05% tretinoin. Although the significance of these studies to humans is not clear, patients should minimize exposure to sunlight or artificial ultraviolet irradiation sources.

The mutagenic potential of tretinoin was evaluated in the Ames assay and in the *in vivo* mouse micronucleus assay, both of which were negative.

In dermal Segment I fertility studies in rats, slight (not statistically significant) decreases in sperm count and motility were seen at 0.5 mg/kg/day (20 times the maximum human systemic dose adjusted for total body surface area), and slight (not statistically significant) increases in the number and percent of nonviable embryos in females treated with 0.25 mg/kg/day (10 times the maximum human systemic dose adjusted for total body surface area) and above were observed. A dermal Segment III study with RENOVA[®] (tretinoin cream) 0.02% has not been performed in any species. In oral Segment I and Segment III studies in rats with tretinoin, decreased survival of neonates and growth retardation were observed at doses in excess of 2 mg/kg/day (83 times the human topical dose adjusted for total body surface area).

Pregnancy

Teratogenic Effects

Pregnancy Category C

ORAL tretinoin has been shown to be teratogenic in rats, mice, rabbits, hamsters, and subhuman primates. It was teratogenic and fetotoxic in Wistar rats when given orally or topically in doses greater than 1 mg/kg/day (42 times the maximum human systemic dose normalized for total body surface area). However, variations in teratogenic doses among various strains of rats have been reported. In the cynomolgus monkey, which, metabolically, is closer to humans for tretinoin than the other species examined, fetal malformations were reported at doses of 10 mg/kg/day or greater, but none were observed at 5 mg/kg/day (417 times the maximum human systemic dose adjusted for total body surface area), although increased skeletal variations were observed at all doses. A dose-related increase in embryoletality and abortion was reported. Similar results have also been reported in pigtail macaques.

TOPICAL tretinoin in animal teratogenicity tests has generated equivocal results. There is evidence for teratogenicity (shorted or kinked tail) of topical tretinoin in Wistar rats at doses greater than 1 mg/kg/day (42 times the maximum human systemic dose adjusted for total body surface area). Anomalies (humerus: short 13%, bent 6%, os parietal incompletely ossified 14%) have also been reported when 10 mg/kg/day was dermally applied.

There are other reports in New Zealand White rabbits administered doses of greater than 0.2 mg/kg/day (17 times the maximum human systemic dose adjusted for total body surface area) of an increased incidence of domed head and hydrocephaly, typical of retinoid-induced fetal malformations in this species.

In contrast, several well-controlled animal studies have shown that dermally applied tretinoin may be fetotoxic, but not overtly teratogenic, in rats and rabbits at doses of 1.0 and 0.5 mg/kg/day, respectively (42 times the maximum human systemic dose adjusted for total body surface area in both species).

With widespread use of any drug, a small number of birth defect reports associated temporally with the administration of the drug would be expected by chance alone. Thirty human cases of temporally-associated congenital malformations have been reported during two decades of clinical use of another formulation of topical tretinoin (Retin-A). Although no definite pattern of teratogenicity and no causal association has been established from these cases, 5 of the reports describe the rare birth defect category holoprosencephaly (defects associated with incomplete midline development of the forebrain). The significance of these spontaneous reports in terms of risk to the fetus is not known.

Non-Teratogenic Effects

Dermal tretinoin has been shown to be fetotoxic in rabbits when administered 0.5 mg/kg/day (42 times the maximum human systemic dose normalized for total body surface area). Oral tretinoin has been shown to be fetotoxic, resulting in skeletal variations and increased intrauterine death, in rats when administered 2.5 mg/kg/day (104 times the maximum human systemic dose adjusted for total body surface area).

There are, however, no adequate and well-controlled studies in pregnant women. RENOVA[®] (tretinoin

cream) 0.02% should not be used during pregnancy.

Nursing mothers

It is not known whether this drug is excreted in human milk. Since many drugs are excreted in human milk, mitigation of fine facial wrinkles with RENOVA[®] (tretinoin cream) 0.02% may be postponed in nursing mothers until after completion of the nursing period.

Pediatric Use

Safety and effectiveness in patients less than 18 years of age have not been established.

Geriatric use

In clinical studies with RENOVA[®] (tretinoin cream) 0.02%, patients aged 65 to 71 did not demonstrate a significant difference for improvement in fine wrinkling when compared to patients under the age of 65. Patients aged 65 and over may demonstrate slightly more irritation, although the differences were not statistically significant in the clinical studies for RENOVA[®] (tretinoin cream) 0.02%. Safety and effectiveness of RENOVA[®] (tretinoin cream) 0.02% in individuals older than 71 years of age have not been established.

ADVERSE REACTIONS

(See **WARNINGS** and **PRECAUTIONS** sections.)

In double-blind, vehicle-controlled studies involving 339 patients who applied RENOVA[®] (tretinoin cream) 0.02% to their faces, adverse reactions associated with the use of RENOVA[®] (tretinoin cream) 0.02% were limited primarily to the skin. Almost all patients reported one or more local reactions such as peeling, dry skin, burning, stinging, erythema, and pruritus. In 32% of all study patients, skin irritation was reported that was severe, led to temporary discontinuation of RENOVA[®] (tretinoin cream) 0.02%, or led to the use of a mild topical corticosteroid. About 7% of patients using RENOVA[®] (tretinoin cream) 0.02%, compared to less than 1% of the control patients, had sufficiently severe local irritation to warrant short-term use of mild topical corticosteroids to alleviate local irritation. About 4% of patients had to discontinue use of RENOVA[®] (tretinoin cream) 0.02% because of adverse reactions.

Approximately 2% of spontaneous post-marketing adverse event reporting for RENOVA[®] (tretinoin cream) 0.05% were for skin hypo- or hyperpigmentation. Other spontaneously reported adverse events for RENOVA[®] (tretinoin cream) 0.05% predominantly appear to be local reactions similar to those seen in clinical trials.

OVERDOSAGE

Application of larger amounts of medication than recommended has not been shown to lead to more rapid or better results, and marked redness, peeling, or discomfort may occur. Oral ingestion of the drug may lead to the same side effects as those associated with excessive oral intake of Vitamin A.

DOSAGE AND ADMINISTRATION

- Do NOT use RENOVA[®] (tretinoin cream) 0.02% if the patient is pregnant or is attempting to become pregnant or is at high risk of pregnancy,
- Do NOT use RENOVA[®] (tretinoin cream) 0.02% if the patient is sunburned or if the patient has eczema or other chronic skin conditions of the face,
- Do NOT use RENOVA[®] (tretinoin cream) 0.02% if the patient is inherently sensitive to sunlight,
- Do NOT use RENOVA[®] (tretinoin cream) 0.02% if the patient is also taking drug(s) known to be photosensitizers (e.g., thiazides, tetracyclines, fluoroquinolones, phenothiazines, sulfonamides) because of the possibility of augmented phototoxicity.

Patients require detailed instruction to obtain maximal benefits and to understand all the precautions necessary to use this product with greatest safety. The physician should review the Patient Package Insert.

RENOVA[®] (tretinoin cream) 0.02% should be applied to the face once a day in the evening, using only enough to cover the entire affected area lightly. Patients should gently wash their faces with a mild soap, pat the skin dry, and wait 20 to 30 minutes before applying RENOVA[®] (tretinoin cream) 0.02%. The patient should apply a small pearl-sized (about ¼ inch or 5 millimeter diameter) amount of cream to cover the entire affected area lightly. Caution should be taken when applying the cream to avoid the eyes, ears, nostrils, and mouth.

Application of RENOVA[®] (tretinoin cream) 0.02% may cause a transitory feeling of warmth or slight stinging.

Mitigation (palliation) of fine facial wrinkling may occur gradually over the course of therapy. Up to six months of therapy may be required before the effects are seen.

With discontinuation of RENOVA[®] (tretinoin cream) 0.02% therapy, some patients may lose the mitigating effects of RENOVA[®] (tretinoin cream) 0.02% on fine facial wrinkles. **The safety and effectiveness of using RENOVA[®] (tretinoin cream) 0.02% daily for greater than 52 weeks have not been established.**

Application of larger amounts of medication than recommended may not lead to more rapid or better results, and marked redness, peeling, or discomfort may occur.

Patients treated with RENOVA[®] (tretinoin cream) 0.02% may use cosmetics but the areas to be treated should be cleansed before the medication is applied. (See **PRECAUTIONS** section.)

HOW SUPPLIED

RENOVA[®] (tretinoin cream), 0.02% is available in tubes containing 40 grams (NDC 42254-165-40).

Storage

Store at 25° (77°F), excursions permitted to 15-30°C (59°-86°F).

QUESTIONS

Physicians and Pharmacists can call 1-800-426-7762, from 8:30 a.m. to 4:30 p.m. Eastern time, Monday through Friday.

Rx only.

Marketed by: **Ortho Dermatologics™**
DIVISION OF ORTHO-McNEIL-JANSSEN
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Rebel Distributors Corp
Thousand Oaks, CA 91320

RENOVA[®]
(TRETINOIN CREAM)
0.02%

RENOVA[®] (reh-NO-vah)

Generic Name: Tretinoin Cream (0.02%)

Use only on the Face

Read this leaflet carefully before you start to use your medicine. Read the information you get every time you get more medicine. There may be new information about the drug. This leaflet does not take the place of talks with your doctor. It is important for you to talk with your doctor about how to use RENOVA[®] (tretinoin cream) 0.02% for the best results and how to reduce side effects.

What is the Most Important Information about RENOVA[®] (tretinoin cream) 0.02%?

RENOVA[®] (tretinoin cream) 0.02% is a serious medicine. **Do not use RENOVA[®] (tretinoin cream) 0.02% if you are pregnant or attempting to become pregnant.** If you become pregnant while using RENOVA[®] (tretinoin cream) 0.02%, please contact your doctor immediately.

Avoid sunlight and other medicines that may increase your sensitivity to sunlight (See "Who should not use RENOVA[®] (tretinoin cream) 0.02%?")

RENOVA[®] (tretinoin cream) 0.02% does not remove wrinkles or repair sun-damaged skin. (See "What is RENOVA[®] (tretinoin cream) 0.02%" for more details.)

What is RENOVA[®] (tretinoin cream) 0.02%?

RENOVA[®] (tretinoin cream) 0.02% is a prescription medicine that may reduce fine wrinkles. It is for patients who are using a total skin care and sunlight avoidance program. RENOVA[®] (tretinoin cream) 0.02% does not remove wrinkles or repair sun-damaged skin. RENOVA[®] (tretinoin cream) 0.02% does not work for everyone who uses it. It may work better for some patients than for others. RENOVA[®] (tretinoin cream) 0.02% should be used only under the guidance of your doctor as part of a sunlight avoidance and total skin care program. This program should include avoiding sunlight as much as possible, using clothing to protect you from sunlight, using sunscreens with a minimum SPF of 15, and using face creams that add moisture to the skin.

When you use RENOVA[®] (tretinoin cream) 0.02%, you will not see improvement right away. Generally, you may notice some effects in 3 to 4 months. If RENOVA[®] (tretinoin cream) 0.02% treatment is stopped, the improvement may gradually disappear.

The use of RENOVA[®] (tretinoin cream) 0.02% in patients for more than 52 weeks has not been studied. Therefore, it is not known if RENOVA[®] (tretinoin cream) 0.02% is safe or works if used longer than 52 weeks. In a study in people with medium to dark skin color, RENOVA[®] (tretinoin cream) 0.02% has not demonstrated a benefit over a sunlight avoidance and total skin care program. RENOVA[®] (tretinoin cream) 0.02% has not been studied in people with visible actinic keratoses or in people with a history of skin cancer.

Who should not use RENOVA[®] (tretinoin cream) 0.02%?

Do not use RENOVA[®] (tretinoin cream) 0.02% if:

- you are pregnant or plan to become pregnant. If you become pregnant while using RENOVA[®] (tretinoin cream) 0.02%, please contact your doctor immediately.
- you are sunburned or your skin is irritated
- you are highly sensitive to sunlight
- you are allergic to any of the ingredients in RENOVA[®] (tretinoin cream) 0.02%. The active ingredient is tretinoin. Ask your doctor or pharmacist about the inactive ingredients.

RENOVA[®] (tretinoin cream) 0.02% can cause increased skin irritation and increased chance of sunburn. Tell your doctor if you have any skin condition, RENOVA[®] (tretinoin cream) 0.02% may not be right

for you.

Because RENOVA[®] (tretinoin cream) 0.02% may make your skin more likely to burn from sunlight, tell your doctor if you are using other medicines that increase sensitivity to sunlight. You should not use RENOVA[®] (tretinoin cream) 0.02% with such medicines. These include, but are not limited to:

- thiazides (to treat high blood pressure)
- tetracyclines, fluoroquinolones, sulfonamides (to treat infection)
- phenothiazines (to treat serious emotional problems)

If you are taking any prescription or non-prescription medicines, check with your doctor to make sure you can use RENOVA[®] (tretinoin cream) 0.02% with them.

We do not know if RENOVA[®] (tretinoin cream) 0.02% is passed to infants through breast milk. Therefore, tell your doctor if you are breast feeding.

How should I use RENOVA[®] (tretinoin cream) 0.02% ?

Use RENOVA[®] (tretinoin cream) 0.02% as a part of a total skin care and sun avoidance program. Follow your doctor's instructions on how to use RENOVA[®] (tretinoin cream) 0.02%. RENOVA[®] (tretinoin cream) 0.02% is usually applied to the face once a day in the evening, following the 3 steps listed below:

1. Gently wash your face with a mild soap.
2. Pat the skin dry and wait 20-30 minutes before applying RENOVA[®] (tretinoin cream) 0.02%.
3. TUBE: Squeeze a small amount of RENOVA[®] (tretinoin cream) 0.02% (the size of a pearl about ¼ inch or 5mm diameter) on your fingertip, and apply to your face.

PUMP: Fully depress the pump twice to dispense a small amount of RENOVA[®] (tretinoin cream) 0.02% (the size of a pearl—about ¼ inch or 5mm diameter) on your fingertip, and apply to your face.

This should be enough to cover your affected area lightly.

Be especially careful when applying RENOVA[®] (tretinoin cream) 0.02% to avoid your eyes, ears, nostrils, angles of the nose, and mouth. RENOVA[®] (tretinoin cream) 0.02% may cause severe redness, itching, burning, stinging, and peeling if used on these areas.

You may use cosmetics one hour after applying RENOVA[®] (tretinoin cream) 0.02%. If you do, be sure to clean your face before applying RENOVA[®] (tretinoin cream) 0.02% again. Skin moisturizers should be used at least every morning to protect the treated areas from dryness.

Use sunscreen and wear protective clothing to protect the treated areas from sunlight. If you sunburn easily, or if you spend a lot of time exposed to sunlight, be especially careful to protect your skin.

What should I avoid while using RENOVA[®] (tretinoin cream) 0.02% ?

RENOVA[®] (tretinoin cream) 0.02% can make your treated skin more sensitive to sunlight. Therefore, keep out of the sunlight as much as possible and do not use sunlamps. Avoid as much as possible products that can increase skin irritation, such as:

- other skin medicines
- medicated or abrasive (rough) soaps
- permanent wave solutions
- chemical hair removers or waxes
- electrolysis
- products with alcohol, spices, astringents, or lime
- cleansers, shampoos, or cosmetics with a strong drying effect
- other products that may irritate your skin

What are the possible side effects of RENOVA[®] (tretinoin cream) 0.02% ?

You may feel brief warmth or stinging on your skin after you use RENOVA[®] (tretinoin cream) 0.02%. Most patients report peeling, dry skin, burning, stinging, itching, and redness. These are usually mild to moderate and occur early in treatment. Contact your doctor if the side effects are a problem.

General advice about prescription medicines

Medicines are sometimes prescribed for conditions that are not mentioned in patient information leaflets. Only use RENOVA[®] (tretinoin cream) 0.02% to treat the condition that your doctor has prescribed it for. Do not give RENOVA[®] (tretinoin cream) 0.02% to other people. It may harm them.

This leaflet summarizes the most important information about RENOVA[®] (tretinoin cream) 0.02%. If you would like more information, talk with your doctor. You can ask your pharmacist or doctor for information about RENOVA[®] (tretinoin cream) 0.02% that is written for health professionals.

Storage: Store at 25° (77°F), excursions permitted to 15-30°C (59°-86°F).

Marketed by: **Ortho Dermatologics™**
DIVISION OF ORTHO-McNEIL-JANSSEN
PHARMACEUTICALS, INC. Los Angeles, CA 90045

Manufactured by:
DRAXIS Specialty Pharmaceuticals Inc., Qc, Canada H9H 4J4

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Repackaged by:

Rebel Distributors Corp
Thousand Oaks, CA 91320


PRINCIPAL DISPLAY PANEL

Peel

Renova 0.02%
40gm Tretinoin Cream
Lot #:00000 Exp. Date: 00/00/00
NDC 42254-165-40 AWP:\$411.34

Peel

Renova 0.02%
40gm Tretinoin Cream
Lot #:00000 Exp. Date: 00/00/00
NDC 42254-165-40 AWP:\$411.34



PHYSICIAN PARTNER
A DIVISION OF REBEL DISTRIBUTORS CORP.

NDC 42254-165-40

Renova 0.02%

40gm Tretinoin Cream

Each tube contains: Tretinoin 0.02%

See package insert

Product ID: SR016540

Peel

Renova 0.02%
40gm Tretinoin Cream
Lot #:00000 Exp. Date: 00/00/00
NDC 42254-165-40 AWP:\$411.34

Peel

Renova 0.02%
40gm Tretinoin Cream
Lot #:00000 Exp. Date: 00/00/00
NDC 42254-165-40 AWP:\$411.34

RX#Master RX Only Mfr. By: Ortho Pharmaceutical a Division of Janssen Ortho LLC Manati, Puerto Rico 00674-0463

Distributed by: Physician Partner, Thousand Oaks, CA 91320 www.physicianpartner.com
Store at controlled room temperature 15°-30° C (59°-86° F) Keep medication out of the reach of children.

RENOVA			
tretinoin cream			
Product Information			
Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:42254-165(NDC:0062-0187)
Route of Administration	TOPICAL		
Active Ingredient/Active Moiety			
	Ingredient Name	Basis of Strength	Strength
	Tretinoin (UNII: 5688UTC01R) (Tretinoin - UNII:5688UTC01R)	Tretinoin	0.2 mg in 1 g
Inactive Ingredients			
	Ingredient Name	Strength	
	Benzyl Alcohol (UNII: LKG8494WBH)		
	Butylated Hydroxytoluene (UNII: 1P9D0Z171K)		
	Medium-Chain Triglycerides (UNII: C9H2L21V7U)		
	Cetyl Alcohol (UNII: 936JST6JCN)		
	Edetate Disodium (UNII: 7FLD91C86K)		
	Methylparaben (UNII: A218C7HI9T)		

Propylparaben (UNII: Z8IX2SC1OH)	
Water (UNII: 059QF0KO0R)	
Stearic Acid (UNII: 4ELV7Z65AP)	
Stearyl Alcohol (UNII: 2KR89I4HIY)	
Steareth-2 (UNII: V56DFE46J5)	
Steareth-20 (UNII: L0Q8IK9E08)	
Xanthan Gum (UNII: TTV12P4NEE)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:42254-165-40	1 in 1 CARTON		
1		40 g in 1 TUBE		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
NDA	NDA021108	10/18/2011	

Labeler - Rebel Distributors Corp (118802834)

Establishment

Name	Address	ID/FEI	Business Operations
Rebel Distributors Corp		118802834	RELABEL, REPACK

Revised: 2/2012

Rebel Distributors Corp