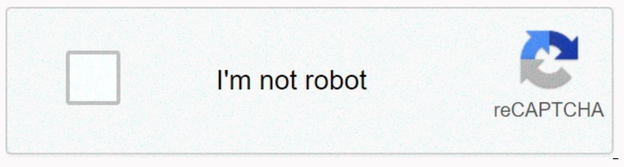

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Brilliant distinctions new app

High-quality skin care is expensive, but fortunately it can save you a lot of the treatments you need for healthy, glowing skin. Savvy skin smart patients know that colorful distinctions empower them to earn future savings on some of their favorite products. Now that version 2.0 of the brilliant distinguished smartphone app has been released, there is no better time to start scoring! Why should you subscribe to the brilliant distinction: get Botox injections regularly? Do you like the skin medica moisturizer you use every day? If allergen treatments are part of your skincare routine, the gorgeous distinction is perfect for you. With brilliant distinction, you can earn points for all qualified purchases. Most of Skin Smart's most popular products and procedures apply to this category. These include luxurious SkinMedica products such as peels, allergen injections (Botox, jubederm fillers, kibella chin treatments, etc.), Litera 2.0 pigment correction serums, glyph firming lotions, and AHA/BHA denail removal cleansers. For every 100 points earned through a brilliant distinction, you get back $10 for future purchases. Points are added quickly, and the more you spend, the more you save! Why you should download colorful distinguishing applications: colorful distinctions will not save you money; It saves you time, too. Gorgeous Distinguish free smartphone app allows you to quickly and easily earn reward points, monitor and spend. Just last February, it announced version 2.0 of the app, bringing the latest in technological innovation to the brilliant distinguishing rewards program. With the app's new update to design and functionality, it's easier than ever to view and manage colorful distinguished accounts. Users with brilliant distinction can now register for the rewards program, view available points and coupons and save coupons in their mobile wallets - all from smartphones! Who we are: The Skin Smart Dermatology and Skin Care Centre is a team of dermatologists and hairdressers who provide Flutown, Jamison, Glenside, Lansdale, Ervington, Doylestown, Roxborough, Oreland, Warrington, Chestnut Hill, New Britain, Lancedale and the surrounding area. Our practice is committed to providing patients with high quality skin care procedures and products. Want to learn more about our services or make a commitment to the process? You can reach it at 215-348-7335 or via our website. BOTOX® is a prescription drug injected into muscles and used to temporarily improve moderate to severe forehead lines, crow's foot lines and frown lines between adult eyebrows. Important safety information botox® cosmetics can cause serious side effects that can be life-threatening. If you ® this problem at any time after botox or cosmetic injections (several hours to weeks), seek medical help immediately: weakness in the muscles involved can cause serious and loss of swallowing, speaking or breathing problems. Life. These problems are at the highest risk if existing before injection. Swallowing problems can last for several months with the spread of toxin effects. The effects of botulinum toxin can affect areas far from the injection site and cause serious symptoms such as loss of strength and muscle weakness, double vision, blurred vision and drooping eyelids, loss of hoarseness or voice, problems speaking words clearly, loss of bladder control, shortness of breath and swallowing problems. BOTOX® dosing devices cannot be identical or comparable to other botulinum toxin products. BOTOX® cases of spread of toxin effects were not identified when cosmetics were used in recommended doses to treat frown lines, crow's foot lines and/or forehead lines. BOTOX® cosmetics can cause loss of strength ® general muscle weakness, vision problems or dizziness within a few weeks to a few weeks after taking Botox, cosmetics. In this case, do not drive a car, operate a machine or do any other dangerous activity. Serious and/or immediate allergic reactions have been reported. They include: itching, rash, red itching welt, whether it's asthma symptoms, or dizziness or faint feeling. Get medical help immediately if you are w whether you have whether you have asthma symptoms, or if you become dizzy or faint. Botox® take cosmetics: Botox® are allergic to ingredients in cosmetics (see drug guide for ingredients); I ® allergic reactions to other botulinum toxin products such as Myobloc (limabotulinumtoxinB), Dysport®® (abobotulinumtoxinA), or zeobotulinumtoxinA (incobotulinumtoxinA). There is a skin infection at the planned injection site. Tell your doctor about any muscle or neurological condition, such as ALS or Lou Garrick's disease, myopia gravitone or Lambert-Eaton syndrome ®. Tell your doctor about all medical conditions, including: planning surgery; I had surgery on my face. I'm having trouble raising my eyebrows. drooping eyelids; other abnormal facial changes; pregnant or planning to get pregnant (Botox® do not know if these cosmetics can harm unborn babies); Breastfeeding or planning (Botox® not known if cosmetics are delivered to breast milk). Tell your doctor about all medications, including prescription and non-prescription medications, vitamins and herbal products. Botox® cosmetics as certain other medicines can cause serious side effects. If you tell your ® that you have received BOTOX, cosmetics in the past, do not start a new drug. If you've received any other botulinum toxin products in the last 4 months, let your doctor know. In the past® have received botulinum toxin injections such as myoblock®, idsport, or zeomin® (tell your doctor exactly which product you received); Recently received antibiotics by injection; Take relaxants; taking allergy or cold medicine; taking sleep pills; Take products like aspirin or blood thinners. Botox® side effects of cosmetics include: dry mouth; discomfort or pain at the injection site; fatigue; headaches; Neck pain; Eye problems: double vision, poor vision, swelling of eyelids and eyebrows, eyelids and dry eyes. For more information, see our medication guide or consult your doctor. To report side effects, call Allergen at 1-800-678-1605. See ® product information, including Botox, box warnings and drug guides. LATISSE® (bimatoprost eye solution) uses 0.03% critical information approval of LATISSE® an FDA-approved treatment to grow eyelashes for people with inappropriate or insufficient eyelashes. Do not use important safety ® you are allergic to one of our ingredients. If you are using/using prescription products for opsp problems, use LATISSE® care. Brown of the colored part of the eye, which is likely to be permanent, can darken. LATISSE® may cause the eyelid skin to darken, which can be reversible. Apply lashes only to base. Do not apply to the lid below. Hair can grow outside the treatment area. Consult a doctor if you have eye problems/surgery. Common side effects include itching and red eyes. When discontinued, the eyelashes gradually return to their previous appearance. These are all possible side effects of LATISSE®. Talk to your doctor for more information. Latis® see full prescribing information. KYBELLA® (deoxycholyc acid) injection 10 mg/mL Important Information Important Safety Information KYBELLA® Is It? KYBELLA® is a prescription drug used to improve the appearance and profile of moderate to severe fat (sub-fat) under the chin, also called double chin in adults. KYBELLA® know if it is safe and effective in treating fatty fats outside the sub-region or under the age of 18. Who shouldn't ® KYBELLA? If you have an infection at the ® do not take KYBELLA. KYBELLA® health care provider before receiving a medical treatment, let them know about any medical conditions, including if you have performed or planned surgery on your face, neck or jaw. I had a beauty treatment on my face, neck or chin. There is a medical condition in or near the neck area. I had or had trouble swallowing. I have a bleeding problem. PREGNANT OR PLANNING TO GET PREGNANT (KYBELLA® NOT KNOW IF IT WILL HARM THE FETUS); Breastfeeding or breastfeeding plans (KYBELLA® not known if it is delivered to breast milk). Let your healthcare provider know about all the medicines you take, including prescription and over-the-counter medicines, vitamins, and herbal supplements. Please let your healthcare provider know if you are taking medicines that prevent blood clotting in particular (anti-platelets or anticoagulant drugs). What? Side effects of KYBELLA®? KYBELLA® can cause serious side effects, Problems swallowing injection site problems including nerve injuries to the jaw (which can cause uneven smiles or facial muscle weakness): collections of blood under the skin (hematemia) or bruising, KYBELLA® damage to arteries or veins if accidentally injected ® accidental injection, hair loss, open soreness (ulcer) damage (ulcer) and damage (ulcer) around tissue cell death sites and tissue cell death sites. Call your healthcare provider if you have: start developing weaknesses in the muscles of your face, or your smile becomes uneven; difficulty swallowing or if symptoms are already worsening; THE MOST COMMON SIDE EFFECTS OFKYBELLA IN DEVELOPING OPEN WOUNDS OR DRAINAGE IN THE TREATMENT AREA® INCLUDING SWELLING, PAIN, NUMBNESS, REDNESS, AND AREAS OF HARDNESS IN THE TREATMENT AREA. These are all possible side effects of ®. Call your doctor for medical advice on side effects. KYBELLA® full prescribing information. Check the full prescribing information, contact your healthcare provider or MyKybella.com your doctor. CoolSculpting® treatment important information using coolscal® the procedure is subtly FDA-cleared for the treatment of visible fat bulges (under the chin) and submandibular (below the jawline) area, thighs, abdomen and sides, bra fat, waist fat, hips (also known as banana rolls) and forearms. It also affects the appearance of FDA-sub-regional treatments and loose tissues®. Important safety information cool ® procedure is not for everyone. If you suffer from cryoglobulinemia, cold agluthin disease, or seizure-®, you should not use CoolSculpting. Consult your doctor if you have a medical condition that includes recent surgery, an existing hernia, known sensitivity, or allergies. During the procedure, you may experience pulling, pulling, light pinching, intense cold, tingling, tingling, pain and spasms in the treatment area. This sensation subsides when the area is paralyzed. Depending on the procedure, typical side effects include temporary redness, swelling, faintness, bruising, stiffness, tingling, tingling, tenderness, convulsions, pain or itching, or skin sensitivity, and a feeling of fullness behind the neck after treatment of the appytal or submandib area. Rare side effects may also occur. CoolSculpting® develop 2-5 months after treatment and can cause visible enlargement of the treatment site, which requires surgical intervention for correction. For more information, see Full Critical Safety Information. CoolTone™ important information on treatment uses cooltones™ and the device is cleared by the FDA for improvement of abdominal tone, strengthening abdominal muscles, and development for a tighter abdomen. Coolton™ FDA Clear Strengthening, toning and firming of the hips and thighs. Important safety information Coolton™ procedure is not for everyone. In areas with metal or electronic implants/devices such as heart raters, implanted hearing aids, implanted defibrillation, implanted neurosyclars, drug pumps and hearing aids, there should be no treatment ™ CoolTone. Talk to your doctor if you have any medical conditions such as CoolTone™ should not be used through the menstrual uterus, over areas of skin where normal feeling is lacking, fever, malignant tumors, bleeding conditions, epilepsy, recent surgical procedures, lung inability, or pregnancy. CoolTone™ be used with caution in patients with Graves disease (an autoimmune disorder that causes irritable thyroid), active bleeding disorders or seizure disorders. Women close to menstruation may find that it comes sooner, or convulsions increase or CoolTone™ treatment and strengthening, therefore, it is not recommended to undergo treatment during this period of the month. CoolTone™ not be used on the heart or head area, new bone growth areas, carotid sinus nerves or over the neck or mouth. CoolToneTM should not be applied to swollen, infected, inflamed areas or skin eruptions. Caution should be used for patients with doubts or diagnosed heart problems. Common side effects may include, but may not be limited to, muscle pain, temporary muscle spasms, temporary joint or tendon pain, and seizures at or near treatment sites. CoolTone™ provider to see if it is appropriate. For more information coolsculpting.com/cooltone information about the safety of your device, see Full Critical Safety Information. JUVÉDERM® injectable gel filler important information approval use JUVÉDERM® VOLUMA™ XC injectable gel is injected deeply into the cheek area to correct age-related volume loss and improve the jaw profile of adults 21 years of age or more for enlargement of the jaw area. JUVÉDERM® VOLLURE™ XC and JUVÉDERM® XC injectable gels are injected into the face tissue for moderate to moderate to severe face wrinkles and wrinkles for nasal wrinkle-like folds. JUVÉDERM® VOLLURE™ XC injectable gel is for adults 21 years of age and older. JUVÉDERM®™ XC injectable gel is injected into the lips for lip enlargement and corrects the line of benefits in adults over the age of 21. JUVÉDERM® Ultra XC Injectable Gel is for infusion into lips and biplane areas for lip enlargement in adults over 21 years of age. Important safety information JUVÉDERM® why shouldn't you get a formula? Do not use these products if you have a history of various serious allergies or severe allergic reactions (anaphylaxis) or are allergic to lidocaine or the gram-positive bacterial proteins used in this product. What precautions should a doctor advise? Minimize strenuous exercise and exposure to a wide range of sun or heat within the first 24 hours of treatment. Exposure is The safety of these products for use during pregnancy or breastfeeding is JUVÉDERM® VOLUMA™ XC has not been studied for more than 80 years for enlarged cheeks or jaw enlargement under the age of 35 or over 65. The safety of JUVÉDERM® VOLLURE™ XC and JUVÉDERM®™ Volvéderm™ XC was not studied in patients under the age of 22, and the safety of JUVÉDERM® XV and JUVÉDERM® Ultra XC was not studied in patients under the age of 18® VOLUMA™ XC is for use in jaw and cheeks. JUVÉDERM® bolouer™ XC and JUVÉDERM® XC are intended for use in facial wrinkles and wrinkles. JUVÉDERM® volbella™ XC and JUVÉDERM® ultra XC are intended for use on the lips and surrounding areas. Safety and effectiveness for treatment in other areas tell your doctor if you have a history of excessive scarring if it is not established in clinical studies that have not been established in clinical studies (thick, hard scars) or pigmentation disorders. THE SAFETY OF JUVÉDERM® PRODUCT HAS NOT BEEN STUDIED IN THESE PATIENTS AND IF IT IS IN THE TREATMENT USED TO REDUCE THE BODY'S IMMUNE RESPONSE, ADDITIONAL SCARRING OR CHANGES IN PIGMENTATION MAY OCCUR TO THE DOCTOR (IMMUNOSUPPLYTIC THERAPY). Using substances that can prolong bleeding, such as aspirin, ibuprofen or other blood thinners, may increase the risk of infection before treatment. As with any injection, this may result in increased bruises or bleeding at the injection site experiencing skin damage near the injection site, patients have not studied side effectsJUVÉDERM® VOLUMA™ XC has not been studied in patients with significant loose skin of the jaw, neck, or in patients with finite loose skin of the jaw juVÉDERM® VOLUMA™ the effect of XC injection has not been studied any side effects? THE MOST COMMONLY REPORTED SIDE EFFECTS WITH JUVÉDERM® WEEKLY GELS INCLUDE REDNESS, SWELLING, PAIN, TENDERNESS, ELASTICITY, LUMPS/BUMPS, BRUISING, DISCOLORATION, AND ITCHING. Dryness was ® for juvéderm™ and XC. FOR JUVÉDERM® VOLUMA™ XC, MOST SIDE EFFECTS ARE RESOLVED IN 2 RECEIVING HOURS 4 WEEKS®®™®. JUVÉDERM® Volbella™ XC were mostly resolved within 30 days. These side effects are consistent with other face injection procedures. Most side effects are resolved over time. Your doctor may choose to treat side effects that last more than 30 days, along with antibiotics, steroids, or hyaluronidase (an enzyme that breaks down hyaluronic acid). One of the risks with this product is the intended chipping infusion into blood vessels. The chances of this happening are very small, but if it happens, the complications can be serious and may be permanent. This reported complication Face injections can include vision problems, blindness, stroke, temporary scabs or permanent scarring of the skin. As with all skin injection procedures, there is a risk of infection. Juvederm.com or consult your doctor for more information. JUVÉDERM® to report side effects to your product, call allergens at 1-800-433-8871. PRODUCTS FROM ® JUVÉDERM COLLECTION CAN ONLY BE USED BY LICENSED PHYSICIANS OR PROPERLY LICENSED CONTRACTORS. Natrelle® Breast Implants Important Information Who Can Get Breast Implants? Natrelle® breast implants are approved for women for breast augmenting for women at least 22 years old for silicone-filled implants. Breast enlargement for at least 18-year-old women for saline-filled implants. Breast augmentation includes primary breast enlargement to increase breast size, as well as revision surgery to correct or improve the results of primary breast enlargement surgery. Breast reconstruction. Breast reconstruction includes primary reconstruction to replace breast tissue that has been removed due to cancer or trauma or has failed to develop properly due to severe breast injury. Breast reconstruction also includes revision surgery to correct or improve the outcome of primary breast reconstruction surgery. Important Safety Information Who Shouldn't Get Breast Implants? Women with active infections anywhere in their bodies. Women with pre-existing cancer or precancery of their breasts who have not received proper treatment for that condition. Women currently pregnant or nursing. What do I need to know before I get breast implants? Breast implants are not a lifelong device and are not necessarily a one-time operation. Many changes to the chest after transplantation are irrevers undoable. If you choose not to remove and replace the implant later, you may experience unacceptable dimples, puffering, wrinkles, or other cosmetic changes in the breast that may be permanent. Breast implants can affect your ability to breastfeed by reducing or eliminating milk production. Ruptures of silicone-filled breast implants are most often silent and may not be detected by you or your doctor. You should have an MRI 3 years after surgery and then every two years as long as you have breast implants to see if a rupture exists. With breast implants, routine screening mammography and self-examination for breast cancer will be more difficult. Ask your doctor to help you distinguish the implant from breast tissue. Symptoms of ruptured implants can be tight knots or lumps surrounding the implant, or changes or loss of size or shape of the armpits, breast or implant, pain, tingling, swelling, numbness, burning or hardening. Talk to your doctor about these symptoms and get rid of ruptured implants. Tell someone who's treating you to another doctor Implants are present to minimize the risk of damage to the implant. What should I tell my doctor? The risk of breast implant surgery may be higher, so let your doctor know if you have the following conditions: autoimmune diseases (e.g., lupus and sericosis): A weakened immune system (for example, taking drugs that weaken the body's natural resistance to current diseases). Breast implant placement and then planned chemotherapy. Radiotherapy was planned on the breast following the placement of breast implants. Conditions or drugs that interfere with wound healing and blood clotting. Reduces blood supply to breast tissue. Discuss all the history of mental health disorders with your surgeon before surgery. Patients with a diagnosis of depression or other mental health disorders should wait for the stabilization of these conditions before undergoing a resolution or breast transplant. What are the complications with breast implants? The main complications are re-action, removal of implants without replacement, rupture of implants with silicone-filled implants, implant deflation of implants filled with saline, and severe capticular shrinkage (severe scar tissue around implants). Other complications include asymmetry, nipple/breast/skin sensory changes, scarring, or wrinkles/ripples. Talk to your doctor about other complications. Talk to your doctor. For more information or to ® or breast implants, call Allergen at 1-800-433-8871. See www.allergan.com/products patient brochure for more information. Natrelle® breast implants are only available with a prescription. REVOLVE™ use revolve-based systems for advanced ™ sensitive information approvals? REVOLVE™ advanced fat system (REVOLVE™ system) is used for fat aspiration, harvesting, filtering and transmission for aesthetic body formation. REVOLVE™ system is intended for use in the following surgeries when drawing fat: plastic and reconstructive surgery, gastrointestinal and affiliated organ surgery, urology surgery, general surgery, bone or muscle surgery, gynecology surgery, thoracic surgery, minimally invasive surgery. Important safety information Who should not ™ rotational, or system? REVOLVE™ currently has a disease that negatively affects wound healing and is in poor overall health, then the system should not be used by a doctor. Which alerts should I be aware of? REVOLVE™ system does not in itself produce significant weight loss. The device should be used with the care of a doctor if you have a chronic disease such as diabetes, heart, lung or circulatory system disease or obesity. What precautions should I be aware of? REVOLVE™ system is designed to eliminate localized deposits of excess fat through small incisions Pass the organization back. The use of this device is limited to doctors with an appropriate level of medical education and surgical experience in appropriate surgical procedures. The outcome of the procedure depends on age, the site of the operation and the experience of the doctor. The results of the procedure may or may not be permanent. What are the possible side effects? Some common side effects associated with fat transfer are uneven, excessive and/or corrective, tissue lumps, bleeding, and scarring. REVOLVE™ associated with the immune system include fat cell death, cyst formation, infection, chronic immune system response, allergic reactions and inflammation. REVOLVE™ system is only available with a prescription. This information is not intended to replace discussions with surgeons. It does not account for all the potential risks associated with fat transplant procedures. Every patient's situation is different, so talk to your surgeon to ™ use of the REVOLVE-based surgical system is appropriate. For more information, see the Ifu and revolve™ user's guide for your system. To report an adverse reaction, call allergens at 1.800.367.5737. DiamondGlow™ treatment uses diamond glow™ device is a micro skin abrasion device that gently removes the upper layer of the skin and delivers a topical cosmetic serum to the skin. Important Safety Information Diamond ™ treatment is not for everyone. If the quality of the skin is impaired™ not receive a diamond glow or treatment. If you are pregnant or nursing, have a medical condition including allergies, or use topical medications at the site to be treated, please contact your provider. Common side effects include scratched, stinging sensations and temporary tightness during treatment, redness or slight swelling after treatment. Rare serious side effects can also occur and include severe skin irritation and allergic reactions. Pro Injection Serum Disclaimer Pro Injection Serum is intended to meet the FDA's definition of cosmetics, purification, beautifying, articles applied to the human body to promote attractiveness, and change appearance. This product is not a drug that diagnoses, treats, treats, or prevents any disease or condition. These products have not been approved by the FDA and the statement has not been evaluated by the FDA. Contact your provider for more information. SkinMedica® the skin medica described on this website® most of the products are intended to meet the FDA's definition of cosmetics, purifying, beautifying, promoting attractiveness, and articles applied to the human body to change shape. These SkinMedica® are not drug products that diagnose, treat, treat or prevent a disease or condition. These products are not approved by the FDA, and the statements on this page by the FDA. SkinMedica® Total Defense + Repair Wide Spectrum Sunscreen (SPF 34, SPF 34 Tint, SPF 50+) and Essential Defense Wide Spectrum Sunscreen (Everyday Clear SPF 47, Mineral Shield Tint SPF 32, Mineral Shield SPF 35) are over-the-counter drug products manufactured and sold under fda regulations. SkinMedica® Acne System, Acne Treatment Lotion, Forming Wash Cleanse and Purifying Toner is 21 C.F.R. § 333.301 et seq.

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