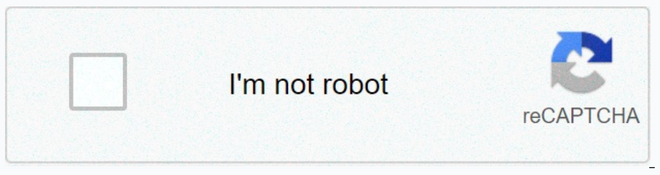

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BOTOX® Cosmetic is a prescription drug that is injected into the muscles and is used to temporarily improve the appearance of moderate to severe forehead lines, crow legs lines, and eyebrows between eyebrows in adults. IMPORTANT SAFETY INFORMATION Botox® beautician can cause serious side effects that may be life-threatening. Seek medical advice immediately if you have any of these problems at any time (hours to weeks) after injection botox® Cosmetic: Problems swallowing, speaking, or breathing, due to weakening of the bound muscles, can be severe and cause loss of life. If these problems are present before you inject, you are at greatest risk. Swallowing problems can last for several months for toxin effects. The effect of botulinum toxin can affect areas away from the injection site and cause serious symptoms including: powerlessness and muscle weakness, double vision, blurred vision and drooping eyelids, hoarseness or change of voice or loss of voice, problems with clear words, loss of bladder control, difficulty breathing and difficulty swallowing. BOTOX® cosmetic dosing devices are not the same and are not comparable to any other botulinum toxin product. Botox® Cosmetic has been used at the recommended dose for the treatment of frown lines, crow legs and/or end lines, no serious spread of the toxin effect has been confirmed. BOTOX® cosmetic may cause loss of power or general muscle weakness, visual disturbances or dizziness within hours to weeks of taking BOTOX® Cosmetic. If this happens, do not drive, use machines or carry out any other dangerous activities. Serious and/or immediate allergic reactions have been reported. These include: itching, rash, red itchy welts, wheezing, asthma symptoms, or dizziness or feeling faint. Contact your doctor immediately if you have wheezing or if you experience asthma symptoms or if you experience dizziness or fainting. Do not take BOTOX® Cosmetic if you: are allergic to any of the ingredients of BOTOX® Cosmetic (see Medicines guide ingredients); allergic reactions to another botulinum toxin medicine such as Myobloc® (rimabotulinumtoxinB), Dysport® (abobotulinumtoxinA) or Xeomini® (incobotulin toxinA); you have a skin infection planned at the injection site. Tell your doctor about any muscle or nerve conditions such as ALS or Lou Gehrig's disease, myasthenia gravis or Lambert-Eaton syndrome, as you may be at increased risk of serious side effects, including difficulty swallowing and breathing difficulties due to typical doses of BOTOX® Cosmetic. Tell your doctor about all your medical conditions, including: plans to have surgery; your face has had surgery; you have difficulty raising your eyebrows; salivation of the eyelids; other abnormal facial changes; are pregnant or are planning to have a baby (it is not known if BOTOX® Cosmetic may harm your unborn baby); breast-feeding or plan (are (on if BOTOX® is excreted in human milk). Tell your doctor about all medicines you are taking, including prescription medicines and prescription medicines, vitamins and herbal medicines. Botox® With certain other medicines can cause serious side effects. Do not start with new medicines until you have told your doctor that you have previously received BOTOX® Cosmetic. Tell your doctor if you have received another botulinum toxin in the last 4 months; you have previously received a botulinum toxin such as myobloc®, Dysport® or Xeomin injection® (tell me exactly what medicine you received); you have recently been given an antibiotic by injection. take muscle relaxlers take allergy or cold medicine; take a sleep medication; aspirin products or blood thinners. Other side effects of BOTOX® cosmetic include: dry mouth; discomfort or pain at the injection site tiredness; headache; neck pain; and eye problems: double vision, blurred vision, impaired vision, salivation of the eyelids and eyebrows, swelling of the eyelids and eyes. For more information, see the Medicines Guide or talk to your doctor. To report any adverse reactions, call Allergan at 1-800-678-1605. Please refer to BOTOX® Cosmetic for complete product information including boxed warning and medication guide. LATISSE® (bimatoprost ophthalmic solution) 0.03% Important information Approved Use latisse® is an FDA-approved treatment for eyelash growing in people with insufficient or inadequate eyelashes. Important safety information Do not use LATISSE ® if you are allergic to any of these ingredients. If you are using/use prescription products to treat eye pressure problems, use ® your doctor. Can cause brown darkening in the colored part of the eye, which is likely to be permanent. LATISSE® may cause the eyelid to darken the skin, which may be reversible. Apply only on the basis of upper eyelashes. DO NOT WEAR ON THE LOWER COVER. Hair can grow outside the treatment area. If you have eye problems/surgeries, talk to your doctor. Common side effects are itchy and red eyes. When stopped, the eyelashes gradually return to the previous look. These are not all possible side effects of LATISSE®. Talk to your doctor for more information. Please refer to LATISSE® complete prescribing information. KYBELLA® (deoxycholic acid) injection 10 mg/ml Important information Important safety information What is KYBELLA®? KYBELLA® a prescription medicine used in adults to improve the appearance and appearance of moderate to heavy fat under the chin (submental fat), also called double chin. It is not known whether KYBELLA ® and effective for the treatment of fat outside the sub-area or in children under 18 years of age. Who shouldn't ® KYBELLA? Do not receive KYBELLA if you have ® in the treatment area. Before you can ® KYBELLA, talk to your doctor all your medical conditions, including if you: have been or are planning surgery on your face, neck or chin. has had cosmetic treatment on your face, neck, or chin; you have had or have medical conditions in or near the neck area. you have had or have had swallowing problems. you have bleeding problems are pregnant or planning to have a baby (it is not known if KYBELLA® harms your unborn child) breast-feeding or plan to breast-feed (it is not known whether KYBELLA ® excreted in breast milk). Tell your healthcare provider about all the medicines you take, including prescription and over-the-counter medicines, vitamins and herbal medicines. In particular, tell your healthcare professional if you are taking a medicine that prevents blood clotting (a medicine that inhibits platelet aggregation or an anticoagulant medicine). What are the possible side effects of KYBELLA®? KYBELLA® can cause serious side effects, including nerve damage in the jaw (which may cause uneven smile or weakness of the facial muscles) Problems with injection site shortening problems Injection site problems including: accumulation of blood under the skin (haematoma) or bruising, artery or vein damage when KYBELLA® is injected inadvertently, hair loss, open sores (ulcers), damage and dog-like death (necrosis) around the injection site. Call your health care provider if: you start to develop weakness in the muscles of your face or your smile becomes uneven; if you have difficulty swallowing or if you have already worsened any symptoms. develop open ulcers or drainage in the treatment areaKybella most common side effects® swelling, pain, numbness, redness and hardness areas in the treatment area. These are not all possible side effects ® KYBELLA. Talk to your doctor about side effects. Please refer to KYBELLA® full prescribing information. Please refer to the full prescribing provided with you or ask your healthcare professional or visit MyKybella.com. CoolSculpting® treatment important information uses CoolSculpting® procedure is an FDA-passed treatment of visible fat bulge submental (under the chin) and submandibular (under the jaw) areas of the thigh, abdomen and groin, along with bra fat, back fat under the buttocks (also known as banana roll) and upper arm. It is also FDA-passed to affect the appearance of flaccid tissue in the lower treatment. CoolSculpting® is not a cure for weight loss. Important safety information coolsculpting® procedure is not for everyone. You should not have a CoolSculpting® procedure if you have cryoglobulinaemia, cold agglutinin disease, or paroxysmal cold hemoglobinuria. Tell your doctor if you have any medical conditions, including recent surgery, an existing hernia and known sensitivities or allergies. During the procedure you may feel like pulling, tugging, light squeezing, tingling, tingling, aches and seizures at the treatment site. These sensations disappear when the area becomes numb. After the procedure, typical side effects are temporary redness, swelling, blanching, bruising, hardness, tingling, tingling, tenderness, convulsions, pain, itching, or skin sensitivity, and a feeling of fullness in the back of the throat after submental or submandibular treatment. Rare side effects may also occur. CoolSculpting® can cause visible expansion in the treated area, which can occur two to five months after treatment and requires surgical intervention to correct it. For more information, see the full, important safety information. CoolTone™ Essential Information Uses CoolTone™ device is FDA-passed to improve abdominal tone, strengthening the abdominal muscles, and developing a firmer abdomen. CoolTone™ fda-passed strengthening, toning, and firming buttocks and thighs. IMPORTANT SAFETY INFORMATION CoolTone™ procedure is not for everyone. You should not have CoolTone™ treatment areas for metal or electronic implants/devices such as pacemakers, implanted hearing aids, implanted defibrillators, implanted neurostimulants, drug pumps, and hearing aids. Tell your doctor if you have conditions such as CoolTone™ should not be used in menstrual areas, areas of skin caused by normal ity, fever, malignancy, haemorrhagic conditions, epilepsy, recent surgical procedures, lung failure or pregnancy. CoolTone ™ be used with caution in Graves' disease (an autoimmune disorder that causes overactive thyroid glands), problems with active bleeding or seizure disorders. Women who are close to menstruation may find that it should be earlier, or seizures have increased or intensified with CoolTone™ therapy, therefore it is recommended not to undergo treatment at this time of month. CoolTone™ should not be used in the heart or head areas, in areas of new bone growth, over the carotid sinus nerves, or over the neck or mouth. CoolToneTM should not be applied over swollen, infected, inflamed areas or skin outbursts. Caution should be exercised in patients suspected or diagnosed with heart problems. Common side effects may be, but may not be limited, to muscle pain, temporary muscle spasm, temporary joint or tendon pain, and redness or near the treatment site. If CoolTone™ is right for you. For more information, see coolsculpting.com/cooltone. JUVÉDERM® Injectable gel fillers Important information APPROVED USES JUVÉDERM® VOLUMA™ XC injectable gel for deep injection in the cheek area to improve age-related volume loss and jaw enhancement for chin improvement in adults over 21. JUVÉDERM® VOLLURE™ XC and JUVÉDERM® XC injectable gels are injectable gels for injection into facial tissue to correct moderate to severe facial wrinkles and folds, such as nasolabial folds. JUVÉDERM® VOLLURE™ XC injectable gel is intended for adults over 21. JUVÉDERM® VOLBELLA™ XC injectable gel is intended for injection into the lips lips for oral authentication and for correction of perial lines in adults over 21 years of age. JUVÉDERM® Ultra XC injectable gel is injectable into the lips and perioral area of the lips and perioral regions of the lips for augmentation in adults over 21. IMPORTANT SAFETY INFORMATION Are there any reasons why I should not receive juvéderm® composition? Do not use these products if you have a history of a number of severe allergies or severe allergic reactions (anaphylaxis) or if you are allergic to lidocaine or gram-positive bacterial proteins used in these products. What precautions should the doctor advise me on? Reduce the use of tensioners and exposure to extensive sun or heat within the first 24 hours after treatment. Contact with any of these may cause temporary redness, swelling and/or itching at the injection site Tell your doctor if you are pregnant or breast-feeding. The safety of these medicinal products during pregnancy or breast-feeding JUVÉDERM® VOLUMA™ XC has not been studied in cheek-rearing patients below 35 years of age or over 65 years of age in cheek-rearing or in chin augmentation below 22 years of age and over 80 years of age. The safety of JUVÉDERM® VOLLURE™ XC and JUVÉDERM® VOLBELLA™ XC in patients under 22 years of age has not been studied and the safety of JUVÉDERM® XC and JUVÉDERM® Ultra XC has not been studied in juvéderm® VOLUMA™ XC patients below 18 years of age. JUVÉDERM® VOLLURE™ XC and JUVÉDERM® XC are designed for use in facial clects and folds. JUVÉDERM® VOLBELLA™ XC and JUVÉDERM® Ultra XC are intended for lip and perioral use. In clinical trials, the safety and efficacy of the product have not been contaminated, tell your doctor if you have a history of excessive scarring (thick, hard scars) or pigmentation disorders. ® has not been studied in these patients and may cause additional scars or changes in pigmentation Tell your doctor if you are receiving a treatment used to reduce the body's immune response (immunosuppressive therapy). Use may lead to an increased risk of infection Tell your doctor before treatment if you are taking substances that may prolong bleeding, such as aspirin, ibuprofen or other blood thinners. As with any injection, it may cause an increase in bruising or bleeding at the injection site Patients who develop skin lesions near the injection site may be at increased risk of adverse reactions in patients with JUVÉDERM® VOLUMA™ XC chin, neck or loose jaw ® VOLUMA™ XC injection to face growth has not been studied What are the side effects? The most commonly ® of JUVÉDERM injectable gels were redness, swelling, pain, tenderness, hardness, lumps/bumps, bruising, discoloration and itching. ® ™ WAS ALSO REPORTED FOR ™ VOLBELLA AND XC. For JUVÉDERM® VOLUMA™ XC, most of the adverse reactions were resolved by 2® to 14 days or less of THE 2 to 14 days of treatment with JUVÉDERM™ VOLUMA®™ XC®™ XC. ® 30 days or less were resolved for ™ 30 DAYS. These side effects are consistent with other facial injection procedures. Most side effects resolve over time. Your doctor may decide to treat side effects that persist for more than 30 days with antibiotics, steroids or hyaluronidase (an enzyme that breaks down hyaluronic acid). One of the risks associated with these drugs is involuntary injection into a blood vessel. The chances of this happening are very small, but when it happens, complications can be serious and can be permanent. These complications, which are reported as facial injections, may include visual disturbances, blindness, stroke, temporary scabs, or permanent scarring of the skin. As with all skin injection procedures, there is a risk of infection. Visit Juvederm.com or tell your doctor for more information. To report ® of the Juvéderm product, call Allergan at 1-800-433-8871. THE JUVÉDERM® collection products are only available to a licensed doctor or a properly licensed practitioner. Natrelle® Breast Implants Important Information Who Can Get Breast Implants? Natrelle® implants are approved for women as follows: Breast augmentation for women at least 22 years old with silicone-filled implants. Breast augmentation for women aged 18 years and over for saline-filled implants. Breast augmentation includes an initial breast augmentation to increase breast size, as well as a review of the surgery to correct or improve the outcome of primary breast augmentation surgery. Breast reconstruction. Breast reconstruction involves a primary reconstruction to replace breast tissue that has been removed due to cancer or trauma or which is not properly developed due to the abnormal breast. Breast reconstruction also includes a revision surgery to improve or improve the outcome of primary breast reconstruction surgery. IMPORTANT SAFETY INFORMATION Who should not receive breast implants? Women with an active infection everywhere in their bodies. Women with pre-existing cancer or prectal breast cancer who have not received adequate treatment for these conditions. Women who are currently pregnant or breastfeeding. What should I know before breast implants? Breast implants are not not necessarily a single operation. Many changes in breasts after implantation cannot be withdrawn. If you later choose to have your implant(s) removed and not replaced, you may experience unacceptable dimpling, puckering, wrinkling, or other cosmetic changes in the breast that may be permanent. Breast implants can affect your ability to breastfeed, either by reducing or eliminating milk production. A rupture of a silicone-filled breast implant is most often quiet and you or your doctor may not detect it. You should have an MRI 3 years after surgery and then every 2 years after that as long as you have breast implants to determine if a rupture exists. If an implant rupture is indicated by an MRI, then the implant should be removed, with or without replacement. With breast implants, routine screening mammography and self-control of breast cancer are more difficult. Ask your doctor to help you distinguish the implant from breast tissue. Symptoms of a ruptured implant may include hard nodules or lumps surrounding the implant or armpit, a change or loss of the size or shape of the breast or implant, pain, tingling, swelling, numbness, burning or hardening. Tell your doctor about these symptoms and remove the ruptured implants. Tell other doctors who are treating you to reduce the risk of implant damage. What should I tell my doctor? Tell your doctor if you have any of the following conditions, as the risk of breast implant surgery may be higher: Autoimmune diseases (e.g. lupus and scleroderma). Weakened immune systems (for example, currently taking medications that weaken the body's natural resistance to the disease). Scheduled chemotherapy after breast implant placement. Planned radiation therapy for breastfeeding after breast implant placement. Conditions or medications that interfere with wound healing and blood clotting. Decreased blood flow to breast tissue. Clinical diagnosis of depression or other mental health disorders, including body dysmorphic disorder and eating disorders. Before surgery, talk to your surgeon about your history of mental health problems. Patients diagnosed with depression or other mental health disorders should wait for these conditions to resolve or stabilise before starting breast implant surgery. What are some complications of breast implants? The main complications are reworking, removal of the implant with or without replacement, implant rupture with silicone-filled implants, deflation of the implant with saline-filled implants and heavy capsule contracture (heavy scar tissue around the implant). Other complications are asymmetry, nipple/breast/skin sensation changes, scarring, or wrinkling/rippling. Talk to your doctor about other complications. Talk to your doctor. For more information or With Natrelle® breast implants, please call Allergan at 1-800-433-8871. Please also check out www.allergan.com/products. Natrelle® breast implants are available on prescription only. REVOLVE™ Advanced Adipose System Important Information Approved Uses What Is ™ System? REVOLVE™ Advanced Adipose System (REVOLVE™ System) is used for aspiration, harvesting, filtering and transferring fat to aesthetic body molding. THE ™ system is designed for use in the following operations for fat-pulling: plastic and reconstructive surgery, gastrointestinal and associated organ surgery, urological surgery, general surgery, bone or muscle surgery, gynecologic surgery, thoracic surgery, and minimally invasive surgery. IMPORTANT SAFETY INFORMATION Who should not use the ™ system? REVOLVE™ System should not be used by your doctor if you currently have a disease that damages wound healing, and a poor overall health condition. What warnings should I be aware of? THE ™ does not in itself lead to significant weight loss. This device should be used with extreme caution by your doctor if you have a chronic disease such as diabetes, heart, lung or circulatory disease or obesity. What precautions should I be aware of? THE ™ System is designed to remove localized deposits of excess fat through a small incision and then transfer the tissue back to you. The use of this device is limited to doctors with appropriate medical education and surgical experience in appropriate surgical procedures. The results of the procedure vary depending on your age, surgical site and your doctor's experience. The results of the procedure may or may not be permanent. What are the possible side effects? Some of the most common side effects associated with fat transfer are unevenness, overcorrection and/or undercorrection, tissue fragments, bleeding and scarring. Possible ™ to the immune system include death of fat cells, cyst formation, infection, chronic immune system, allergic reaction and inflammation. THE ™ is only available on prescription. This information is not intended to replace the discussion with your surgeon. It does not describe all the potential risks associated with fat transplantation. Each patient's situation is different, so please consult your surgeon to determine whether the ™ of the revolve system is right for you. For more information, see the User Manual (IFU) and the ™ of the SYSTEM. To report adverse reactions, call Allergan at 1.800.367.5737. DiamondGlow™ Treatment Important Information Uses DiamondGlow™ device is a microdermabrasion device that gently removes the top layer of the skin and provides topical cosmetic serums apart IMPORTANT SAFETY INFORMATION DiamondGlow™ treatment is not for everyone. If you have a skin quality disorder, you should not have ™ treatment. Tell your service provider if you are pregnant or breast-sucking, or if you have a medical condition, including an allergy, and if you are taking a topical medicine in the area you are being treated for. Typical side effects include scratching, tingling during treatment and temporary tightness, redness or mild swelling after treatment. Serious side effects, including severe skin irritation and allergic reactions, may also occur rarely. Pro-Infusion Seraes Disclaimer Pro-Infusion Serums are designed to meet the FDA definition of cosmetic product, a product used to cleanse, decorate, promote attractiveness, and change appearance. These products are not intended to be medications that diagnose, treat, treat, or prevent diseases or conditions. These products have not been approved by the FDA and these claims have not been evaluated by the FDA. For more information, contact your service provider. SkinMedica® Most SkinMedica® products described on this website are designed to meet the FDA definition of cosmetic product, a product used to cleanse, decorate, promote attractiveness and change appearance. These SkinMedica® products are not designed to be medications that diagnose, treat, treat, or prevent diseases or conditions. These products have not been approved by the FDA and these reports have not been evaluated by the FDA on these pages. SkinMedica® Total Defense + Repair Broad Spectrum Sunscreen (SPF 34, SPF 34 Tinted and SPF 50+) and Essential Defense Broad Spectrum Sunscreen (Everyday Clear SPF 47, Mineral Shield Tinted SPF 32 and Mineral Shield SPF 35) are over-the-counter drug products that are formulated and marketed in accordance with FDA regulatory regulations set out in Section 21 C.F.R. Part 352. SkinMedica® Acne System, Acne Treatment Cream, Purifying Foaming Wash, and Purifying Toner, are over-the-counter drug products that are formulated and marketed in accordance with FDA regulatory rules set out in Section 333,301 et seq. of section 333,301 et seq.

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