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Latisse® (bimatoprost ophthalmic solution) 0.03%

LATISSE®* is an FDA-approved treatment to grow eyelashes for people with inadequate or not enough lashes.

SkinMedica® AN ALLERGAN COMPANY

SkinMedica®* is a physician-dispensed, cosmetic, and nonprescription skin care product line.

Juvéderm VOLBELLA® XC

JUVÉDERM VOLBELLA® XC is the first and only injectable gel for lip augmentation proven to increase lip fullness and soften the appearance of vertical lip lines for up to 1 year.†

3+ Earn 500 Points

kybella® (deoxycholic acid) injection 10 mg/mL

KYBELLA® is a prescription medicine used in adults to improve the appearance and profile of moderate to severe fat below the chin (submental fat), also called "double chin." It is not known if KYBELLA® is safe and effective for use outside of the submental area or in children less than 18 years of age.

Natrelle®

The *Natrelle*® collection of saline and silicone filled implants is the result of 25 years of experience and represents today's advanced breast augmentation technology.

REVOLVE™ ADVANCED ADIPOSE SYSTEM

REVOLVE™ System is an integrated, high-volume, fat-processing device that quickly and efficiently yields high-quality fat tissue for re-implantation in multiple body areas.

Please see Important Safety Information and accompanying F and LATISSE® (bimatoprost ophthalmic solution) 0.03% and JUVÉDERM VOLBELLA® XC, CoolSculpting®, and SkinMe

Please see Important Safety Information and accompanying Prescribing Information for KYBELLA® (deoxycholic acid) injection 10 mg/mL and **Important Safety Information** for *Natrelle*® Breast Implants and REVOLVE™ Advanced Adipose System at the back of this brochure.

All treatments and products are available by prescription only, excluding Ski
*Earn 100 points for every $150 spent on SkinMedica® and/or LATISSE® pr
†With optimal treatment.

EARNING IS AS EASY AS. . .

1 Earn 100 Points

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JUVÉDERM VOLBELLA® XC is the first and only injectable gel for lip augmentation proven to increase lip fullness and soften the appearance of vertical lip lines for up to 1 year.†

2 Earn 200 Points

BOTOX® Cosmetic onabotulinumtoxinA injection

BOTOX® Cosmetic is a prescription medicine that is injected into muscles and used to temporarily improve the look of both moderate to severe crow's feet lines and frown lines between the eyebrows in adults.

Juvéderm® XC

JUVÉDERM® XC injectable gel is for injection into the facial tissue for the correction of moderate to severe facial wrinkles and folds, such as nasolabial folds. proven to last up to 1 year.†

coolsculpting®

The CoolSculpting® fat-freezing procedure is the only FDA-cleared, non-surgical fat-reduction treatment that uses controlled cooling to eliminate stubborn fat that resists all efforts through diet and exercise.

3 Earn 300+ Points

Juvéderm VOLUMA® XC

JUVÉDERM VOLUMA® XC is the first and only injectable gel proven to last up to 2 years in the cheek area to correct age-related volume loss.†

Juvéderm VOLLURE™ XC

JUVEDERM VOLLURE™ XC is the first and only injectable gel proven to last up to 18 months in moderate to severe facial wrinkles and folds, such as nasolabial folds in adults over 21.†

Please see Important Safety Information and accompanying Prescribing Information for BOTOX® Cosmetic (onabotulinumtoxinA) and LATISSE® (bimatoprost ophthalmic solution) 0.03% and **Important Safety Information** for JUVÉDERM® XC, JUVÉDERM VOLUMA® XC, JUVÉDERM VOLLURE™ XC, JUVÉDERM VOLBELLA® XC, CoolSculpting®, and SkinMedica® at the back of this brochure.

All treatments and products are available by prescription only, excluding SkinMedica®.
*Earn 100 points for every $150 spent on SkinMedica® and/or LATISSE® products.
†With optimal treatment.

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Earn bonus points on every treatment		♦ +50 pts	♦♦ +100 pts
Receive a SkinMedica® gift		♦	♦♦
Your points never expire			♦

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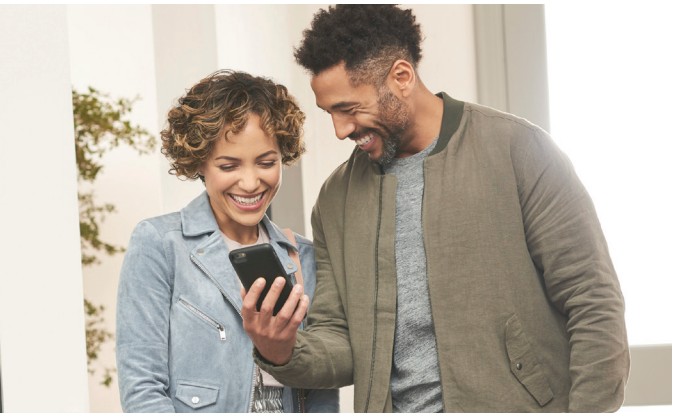

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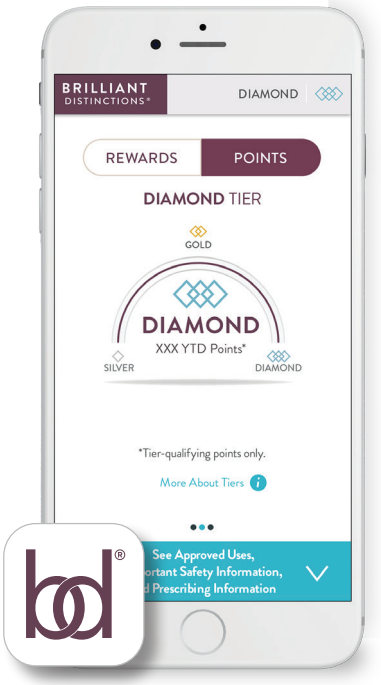

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*Points earned through referring friends and shopping at the BD Mall do not count towards tier advancement.

†Points earned vary by retailer. Retailers are subject to change.

Approved Uses, Important Safety Information, and Prescribing Information

BOTOX® Cosmetic (onabotulinumtoxinA) Important Information

IMPORTANT SAFETY INFORMATION

BOTOX® Cosmetic may cause serious side effects that can be life threatening. Get medical help right away if you have any of these problems any time (hours to weeks) after injection of BOTOX® Cosmetic:

- **Problems swallowing, speaking, or breathing**, due to weakening of associated muscles, can be severe and result in loss of life. You are at the highest risk if these problems are pre-existing before injection. Swallowing problems may last for several months
- **Spread of toxin effects.** The effect of botulinum toxin may affect areas away from the injection site and cause serious symptoms including: loss of strength and all-over muscle weakness, double vision, blurred vision and drooping eyelids, hoarseness or change or loss of voice, trouble saying words clearly, loss of bladder control, trouble breathing, trouble swallowing

BOTOX® Cosmetic dosing units are not the same as, or comparable to, any other botulinum toxin product.
There has not been a confirmed serious case of spread of toxin effect when BOTOX® Cosmetic has been used at the recommended dose to treat frown lines, crow's feet lines, or both at the same time.

BOTOX® Cosmetic may cause loss of strength or general muscle weakness, vision problems, or dizziness within hours to weeks of taking BOTOX® Cosmetic. **If this happens, do not drive a car, operate machinery, or do other dangerous activities.**

Serious and/or immediate allergic reactions have been reported. They include: itching, rash, red itchy welts, wheezing, asthma symptoms, or dizziness or feeling faint. Get medical help right away if you are wheezing or have asthma symptoms, or if you become dizzy or faint.

Do not take BOTOX® Cosmetic if you: are allergic to any of the ingredients in BOTOX® Cosmetic (see Medication Guide for ingredients); had an allergic reaction to any other botulinum toxin product such as *Myobloc®* (rimabotulinumtoxinB), *Dysport®* (abobotulinumtoxinA), or *Xeomin®* (incobotulinumtoxinA); have a skin infection at the planned injection site.

Tell your doctor about all your 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 difficulty breathing from typical doses of BOTOX® Cosmetic.

Tell your doctor about all your medical conditions, including: plans to have surgery; had surgery on your face; weakness of forehead muscles: trouble raising your eyebrows; drooping eyelids; any other abnormal facial change; are pregnant or plan to become pregnant (it is not known if BOTOX® Cosmetic can harm your unborn baby); are breast-feeding or plan to (it is not known if BOTOX® Cosmetic passes into breast milk).

Tell your doctor about all the medicines you take, including prescription and nonprescription medicines, vitamins, and herbal products. Using BOTOX® Cosmetic with certain other medicines may cause serious side effects. **Do not start any new medicines until you have told your doctor that you have received BOTOX® Cosmetic in the past.**

Tell your doctor if you have received any other botulinum toxin product in the last 4 months; have received injections of botulinum toxin such as *Myobloc®*, *Dysport®*, or *Xeomin®* in the past (tell your doctor exactly which product you received); have recently received an antibiotic by injection; take muscle relaxants; take an allergy or cold medicine; take a sleep medicine; take aspirin-like products or blood thinners.

Other side effects of BOTOX® Cosmetic include: discomfort or pain at the injection site; headache; and eye problems: double vision, blurred vision, drooping eyelids, and swelling of your eyelids.

For more information refer to the Medication Guide or talk with your doctor.

To report a side effect, please call Allergan at 1-800-678-1605.

Approved Uses
BOTOX® Cosmetic is a prescription medicine that is injected into muscles and used to temporarily improve the look of both moderate to severe crow's feet lines and frown lines between the eyebrows in adults.

Please see accompanying BOTOX® Cosmetic full Prescribing Information, including Boxed Warning and Medication Guide.

LATISSE® (bimatoprost ophthalmic solution) 0.03% Important Information

Approved Use
LATISSE® is an FDA-approved treatment to grow eyelashes for people with inadequate or not enough lashes.

IMPORTANT SAFETY INFORMATION

If you use/used prescription products for eye pressure problems, use **LATISSE®** under doctor care. May cause brown darkening of the colored part of the eye which is likely permanent. **LATISSE®** may cause eyelid skin darkening which may be reversible. Only apply at base of upper lashes. DO NOT APPLY to lower lid. Hair may grow outside the treatment area. If you have eye problems/surgery, consult your doctor. Common side

effects include itchy and red eyes. If discontinued, lashes gradually return to previous appearance.

To report a side effect, please call Allergan at 1-800-678-1605.

Please see accompanying LATISSE® full Prescribing Information.

KYBELLA® (deoxycholic acid) injection 10 mg/mL Important Information

Approved Use
KYBELLA is a prescription medicine used in adults to improve the appearance and profile of moderate to severe fat below the chin (submental fat), also called "double chin."

It is not known if KYBELLA is safe and effective for use outside of the submental area or in children less than 18 years of age.

IMPORTANT SAFETY INFORMATION
Who should not receive KYBELLA?

You should not receive KYBELLA if you have an infection in the treatment area.

Before receiving KYBELLA, tell your healthcare provider about all of your medical conditions, including if you: Have had or plan to have surgery on your face, neck, or chin; have had cosmetic treatments on your face, neck, or chin; have had or have medical conditions in or near the neck area; have had or have trouble swallowing; have bleeding problems; are pregnant or plan to become pregnant (it is not known if KYBELLA will harm your unborn baby); are breastfeeding or plan to breastfeed (it is not known if KYBELLA passes into your breast milk; talk to your healthcare provider about the best way to feed your baby if you receive KYBELLA).

Tell your healthcare provider about all the medicines you take, including prescription and over-the-counter medicines, vitamins, and herbal supplements. Especially tell your healthcare provider if you take a medicine that prevents the clotting of your blood (antiplatelet or anticoagulant medicine).

What are the possible side effects of KYBELLA?

KYBELLA can cause serious side effects, including nerve injury in the jaw (which can cause an uneven smile or facial muscle weakness), or trouble swallowing.

The most common side effects of KYBELLA include swelling, bruising, pain, numbness, redness, and areas of hardness in the treatment area.

These are not all of the possible side effects of KYBELLA. Call your healthcare provider for medical advice about side effects.

Please see accompanying KYBELLA® full Prescribing Information.

CoolSculpting® Treatment Important Information

The CoolSculpting® procedure is FDA-cleared for the treatment of visible fat bulges in the submental area, thigh, abdomen and flank, along with bra fat, back fat, underneath the buttocks (also known as banana roll), and upper arm. Please talk to your healthcare provider or see full **Important Safety Information** at http://www.coolsculpting.com/pdfs/IC02211-B_Patient_Safety_Info_proof.pdf for additional information.

JUVÉDERM® Injectable Gel Fillers Important Information

Approved Uses
JUVÉDERM VOLUMA® XC injectable gel is for deep injection in the cheek area to correct age-related volume loss in adults over 21.

JUVÉDERM® XC and JUVÉDERM VOLLURE™ XC injectable gels are for injection into the facial tissue for the correction of moderate to severe facial wrinkles and folds, such as nasolabial folds. JUVÉDERM VOLLURE™ XC injectable gel is for adults over 21.

JUVÉDERM® Ultra XC is for injection into the lips and perioral area for lip augmentation in adults over 21.

JUVÉDERM VOLBELLA® XC injectable gel is for injection into the lips for lip augmentation and for correction of perioral lines in adults over 21.

IMPORTANT SAFETY INFORMATION
Are there any reasons why I should not receive any JUVÉDERM® injectable gel formulation?

Do not use these products if you have a history of multiple severe allergies or severe allergic reactions (anaphylaxis), or if you are allergic to lidocaine or the Gram-positive bacterial proteins used in these products.

What precautions should my doctor advise me about?

- Tell your doctor if you are pregnant or breastfeeding. The safety of these products for use during pregnancy or while breastfeeding has not been studied
- The safety of JUVÉDERM VOLUMA® XC in patients under 35 years or over 65 years, the safety of JUVÉDERM® XC and JUVÉDERM® Ultra XC injectable gels in patients under 18 years, and the safety of JUVÉDERM VOLLURE™ XC and JUVÉDERM VOLBELLA® XC in patients under 22 years has not been studied
- The safety and effectiveness of JUVÉDERM VOLUMA® XC in areas other than the cheek area, JUVÉDERM® XC and JUVÉDERM VOLLURE™ XC for areas other than facial wrinkles and folds, and JUVÉDERM® Ultra XC and JUVÉDERM VOLBELLA® XC in areas other than the lips and perioral area have not been established in clinical studies

- Tell your doctor if you have a history of excessive scarring (eg, hypertrophic scarring and keloid formations) or pigmentation disorders, as use of these products may result in additional scars or changes in pigmentation
- Tell your doctor if you are planning other laser treatments or a chemical peel, as there is a possible risk of inflammation at the treatment site if these procedures are performed after treatment
- Patients who experience skin injury near the site of injection with these products may be at a higher risk for side effects
- Tell your doctor if you are on immunosuppressive therapy used to decrease the body's immune response, as use of these products may result in an increased risk of infection
- Tell your doctor if you are using medications that can prolong bleeding, such as aspirin, ibuprofen, or other blood thinners, as this may result in increased bruising or bleeding at the injection site
- Minimize strenuous exercise, exposure to extensive sun or heat, and alcoholic beverages within the first 24 hours following treatment

What are possible side effects?

The most commonly reported side effects with JUVÉDERM® injectable gels included injection-site redness, swelling, pain, tenderness, firmness, lumps/bumps, bruising, discoloration, and itching. For JUVÉDERM VOLBELLA® XC, dryness was also reported. For JUVÉDERM VOLUMA® XC, most side effects were moderate and lasted 2 to 4 weeks. For JUVÉDERM® XC, JUVÉDERM VOLLURE™ XC, and JUVÉDERM® Ultra XC injectable gels, most side effects were mild or moderate and lasted 14 days or less. For JUVÉDERM VOLBELLA® XC, most side effects were mild or moderate and lasted 30 days or less.

One of the risks with using these products is unintentional injection into a blood vessel, and, while rare, the complications can be serious and may be permanent. These complications, which have been reported for facial injections, can include vision abnormalities, blindness, stroke, temporary scabs, or permanent scarring.

As with all skin injection procedures, there is a risk of infection.

To report a side effect with any JUVÉDERM® product, please call Allergan at 1-800-433-8871.Please visit **Juvederm.com** or talk to your doctor for more information.

Available by prescription only.

***Natrelle*® Breast Implants Important Information**

Who may get breast implants?

Natrelle® Breast Implants are approved for women for the following:

- **Breast augmentation for women at least 22 years old for silicone-filled implants.**
- **Breast augmentation for women at least 18 years old for saline-filled implants.**

Breast augmentation includes primary breast augmentation to increase breast size, as well as revision surgery to correct or improve the result of a primary breast augmentation surgery.

IMPORTANT SAFETY INFORMATION
Who should NOT get breast implants?

- Women with active infection anywhere in their body.
- Women with existing cancer or precancer of their breast who have not received adequate treatment for those conditions.
- Women who are currently pregnant or nursing.

What should I know before getting breast implants?

- Breast implants are not lifetime devices, and not necessarily a one-time surgery.
- Many of the changes to your breasts following implantation cannot be undone. If you later choose to have your implant(s) removed and not replaced, you may experience unacceptable dimpling, puckering, wrinkling, or other cosmetic changes of the breast, which may be permanent.
- Breast implants may affect your ability to breast-feed, either by reducing or eliminating milk production.
- Rupture of a silicone-filled breast implant is most often silent and may not be detected by you or your doctor. You should have an MRI 3 years after your surgery and then every 2 years after that for as long as you have your breast implants to determine if rupture is present. If implant rupture is noted on an MRI, you should have the implant removed, with or without replacement.
- With breast implants, a routine screening mammography and self-examinations for breast cancer will be more difficult. Ask your doctor to help you distinguish the implant from your breast tissue. Symptoms of a ruptured implant may be hard knots or lumps surrounding the implant or in the armpit, change or loss of size or shape of the breast or implant, pain, tingling, swelling, numbness, burning, or hardening. Tell your doctor of these symptoms and remove ruptured implants.
- Inform any other doctor who treats you of the presence of your implants to minimize the risk of damage to the implants.

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 (for example, lupus and scleroderma).
- A weakened immune system (for example, currently taking drugs that weaken the body's natural resistance to disease).
- Planned chemotherapy following breast implant placement.
- Planned radiation therapy to the breast following breast implant placement.
- Conditions or medications that interfere with wound healing and blood clotting.
- Reduced blood supply to breast tissue.
- Clinical diagnosis of depression or other mental health disorders, including body dysmorphic disorder and eating disorders. Please discuss any history of mental health disorders with your surgeon prior to surgery. Patients with a diagnosis of depression or other mental health disorders should wait for resolution or stabilization of these conditions prior to undergoing breast implantation surgery.

What are some complications with breast implants?

Key complications are reoperation, implant removal with or without replacement, implant rupture with silicone-filled implants, implant deflation with saline-filled implants, and severe capsular contracture (severe scar tissue around the implant). Other complications include 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 to report a problem with *Natrelle*® Breast Implants, please call Allergan at 1-800-433-8871. Please also see the patient brochures at www.allergan.com/labeling/usa.htm.

Natrelle® Breast Implants are available by prescription only.

REVOLVE™ System

REVOLVE™ System is a single use, sterile, disposable canister that is intended for processing, filtering, and transferring of autologous adipose tissue. The device is intended to be used by a plastic surgeon to perform autologous fat grafting in aesthetic body contouring procedures.

Autologous fat grafting should not be performed in the presence of any disease process that adversely affects wound healing, and in patients who are in poor overall health. Some common adverse effects associated with autologous fat transfer are asymmetry, over- and/or under correction of the treatment site, tissue lumps, bleeding, and scarring.

Every patient's situation is different, so be sure to ask your surgeon if the use of REVOLVE™ System is right for you. This Safety Information is not intended to replace a patient's discussion with a surgeon. It does not describe all the potential risks and potential benefits associated with fat grafting procedures.

REVOLVE™ System is available only through licensed plastic surgeons. Complete product and safety information is available at lifecell.com.

SkinMedica®

Most of the SkinMedica® products described here are intended to meet the FDA's definition of a cosmetic product, an article applied to the human body to cleanse, beautify, promote attractiveness, and alter appearances. These SkinMedica® products are not intended to be drug products that diagnose, treat, cure, or prevent any disease or condition. These products have not been approved by the FDA, and the statements on these pages have not been evaluated by the FDA.

SkinMedica® is a physician-dispensed, cosmetic and nonprescription skin care product line.

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