

Natrelle® GEL REWARDS

Get MORE
of what you
deserve!

Actual Natrelle INSPIRA®
primary augmentation patients.

Individual results may vary.



When you choose any Natrelle® gummy implant for breast augmentation, you and a friend can enjoy one of the complimentary treatments or products below.* First, sign up for *Brilliant Distinctions*®, as only members can take advantage of this offer. Join now at BrilliantDistinctionsProgram.com.

BOTOX®
COSMETIC
onabotulinumtoxinA
injection

|
or
|

 **coolsculpting®**

Only a doctor can determine if BOTOX® Cosmetic is right for you.

CoolSculpting® has not been studied with Natrelle® breast implants, nor is it intended to be used in combination with Natrelle® breast implants.

*Terms and conditions apply; for qualified augmentation patients only. Must be first experience with BOTOX® Cosmetic or CoolSculpting® treatment. Patient has the ability to choose any physician of her choice. BOTOX® Cosmetic is available by prescription only. CoolSculpting® treatment is available only if your physician has the CoolAdvantage™ applicator. A CoolSculpting® treatment is one CoolAdvantage™ cycle.

They have been by your side—now it's your turn to share the love!

Give your friend the gift of an Allergan Aesthetics experience by sharing the next page of this offer. **Remember, this offer must be redeemed within 6 months of your breast augmentation procedure.**

✓ **Bring this form to your consultation appointment for more details.**

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.

Please see Natrelle® Breast Implants Important Safety Information on following pages.

BOTOX® Cosmetic (onabotulinumtoxinA) Important Information

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 forehead lines, crow's feet lines, and frown lines between the eyebrows in adults.

Please see BOTOX® Cosmetic Important Safety Information, including Boxed Warning, on following pages.

Natrelle® GEL REWARDS

A GIFT from your friend!

Actual Natrelle INSPIRA®
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To redeem this offer, simply take this form to your plastic surgeon's office and mention your friend's name within 6 months of her breast augmentation procedure.

Natrelle® Breast Implants 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.

Please see additional **Natrelle® Breast Implants Important Safety Information on following page.**

BOTOX® Cosmetic 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.

Please see additional **BOTOX® Cosmetic Important Safety Information, including Boxed Warning, on following pages.**

Natrelle® Breast Implants IMPORTANT SAFETY INFORMATION (continued)

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 you 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 risks 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 see the patient brochures at www.allergan.com/products. To report a problem with Natrelle® Breast Implants, please call Allergan at 1-800-433-8871.

Natrelle® Breast Implants are available by prescription only.

BOTOX® Cosmetic (onabotulinumtoxinA) IMPORTANT SAFETY INFORMATION (continued)

- **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, and/or forehead lines.

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 receive 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; have 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 over-the-counter medicines, vitamins, and herbal supplements. 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.

Please see additional BOTOX® Cosmetic Important Safety Information on following page.

CoolSculpting® Important Information

Uses

The CoolSculpting® procedure is FDA-cleared for the treatment of visible fat bulges in the submental (under the chin) and submandibular (under the jawline) areas, thigh, abdomen and flank, along with bra fat, back fat, underneath the buttocks (also known as banana roll) and upper arm. It is also FDA-cleared to affect the appearance of lax tissue with submental area treatments. The CoolSculpting® procedure is not a treatment for weight loss.

Important Safety Information

The CoolSculpting® procedure is not for everyone. You should not have the CoolSculpting® procedure if you suffer from cryoglobulinemia, cold agglutinin disease, or paroxysmal cold hemoglobinuria.

Tell your doctor if you have any medical conditions including recent surgery, pre-existing hernia, and any known sensitivities or allergies.

During the procedure you may experience sensations of pulling, tugging, mild pinching, intense cold, tingling, stinging, aching, and cramping at the treatment site. These sensations subside as the area becomes numb. Following the procedure, typical side effects include temporary redness, swelling, blanching, bruising, firmness, tingling, stinging, tenderness, cramping, aching, itching, or skin sensitivity, and sensation of fullness in the back of the throat after submental or submandibular area treatment.

Rare side effects may also occur. CoolSculpting® may cause a visible enlargement in the treated area which may develop two to five months after treatment and requires surgical intervention for correction.

Please see full Important Safety Information on coolsculpting.com.

BOTOX® Cosmetic (onabotulinumtoxinA) IMPORTANT SAFETY INFORMATION (continued)

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, decreased eyesight, drooping eyelids and eyebrows, swelling of your eyelids and dry eyes.

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.

Please see BOTOX® Cosmetic full [Product Information](#) including [Boxed Warning](#) and [Medication Guide](#).

