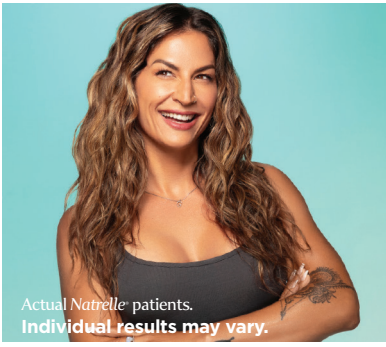
Natrelle | PERKS PROGRAM

REWARDS *for* the NEW YOU

Natrelle PERKS®

Exclusive PERKS with your *Natrelle*® Breast Augmentation.



***Natrelle*® Breast Implants IMPORTANT SAFETY INFORMATION AND APPROVED USES**

Breast implants are not considered lifetime devices. The longer people have them, the greater the chances are that they will develop complications, some of which will require more surgery.

Breast implants have been associated with the development of a cancer of the immune system called breast implant–associated anaplastic large cell lymphoma (BIA-ALCL). This cancer occurs more commonly in patients with textured breast implants than smooth implants, although rates are not well defined. Some patients have died from BIA-ALCL.

Patients receiving breast implants have reported a variety of systemic symptoms, such as joint pain, muscle aches, confusion, chronic fatigue, autoimmune diseases, and others. Individual patient risk for developing these symptoms has not been well established. Some patients report complete resolution of symptoms when the implants are removed without replacement.

Please see additional Important Safety Information on following pages.

NATRELLE PERKS®

is an exclusive offer for Allē Members who receive breast augmentation with *Natrelle*® gel breast implants.

Earn 500 Allē points on your gel breast implants with *Natrelle*®, plus get a complimentary aesthetic treatment offer with either BOTOX® Cosmetic (onabotulinumtoxinA) or JUVÉDERM® Ultra XC.

Natrelle PERKS®

BOTOX® COSMETIC onabotulinumtoxinA injection

Only BOTOX® Cosmetic is FDA approved to temporarily make moderate to severe frown lines, crow's feet, and forehead lines look better in adults.

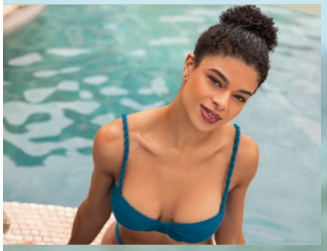
OR

JUVÉDERM® ULTRA XC

JUVÉDERM® Ultra XC is an injectable gel that temporarily adds more fullness and plumps thin lips for adults over the age of 21.

"Feeling good about my new look is my PERK."

What's yours?



Actual *Natrelle*® patients.
Individual results may vary.

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

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, and 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.**

Please see additional Important Safety Information throughout for *Natrelle*® and BOTOX® Cosmetic, including Boxed Warnings, and JUVÉDERM® Collection of Fillers.

Talk to Your Specialist

and ask for BOTOX® Cosmetic by name to see if it's right for you.

BOTOX® COSMETIC onabotulinumtoxinA injection

Only BOTOX® Cosmetic is FDA approved to temporarily make moderate to severe frown lines, crow's feet, and forehead lines look better in adults.

Since FDA approval in 2002, BOTOX® Cosmetic is the #1 selling product* of its kind on the market.

*Data collected through September 2021 (N = 970).

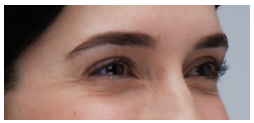
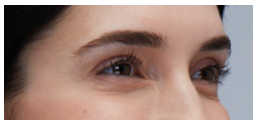
BEFORE AFTER **DAY 2** AFTER **DAY 30** AFTER **MONTH 4**

20 Units • *Moderate to severe* **frown lines**

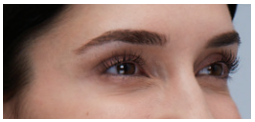

Real patient. Results may vary.
Photos taken at maximum frown before and 2 days after treatment with BOTOX® Cosmetic.


Real patient. Results may vary.
Photos taken at maximum frown 30 and 120 days after treatment with BOTOX® Cosmetic. In clinical studies, physicians assessed 80% of adults had significant improvement at day 30; in the same studies, 89% of adults who were treated saw at least moderate improvement at day 30. In the same studies, physicians assessed 25% of people to have at least moderate improvement on day 120.

24 Units • *Moderate to severe* **crow's feet lines**

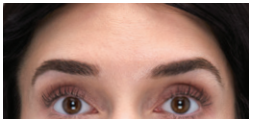
Real patient. Results may vary.
Photos taken at full smile before and 2 days after treatment with BOTOX® Cosmetic.




Real patient. Results may vary.
Photos taken at full smile 30 and 120 days after treatment with BOTOX® Cosmetic. In 2 clinical studies, 26.1% and 20.3% of adults had a ≥ 2-grade improvement at day 30. In one of these studies, 67.9% had mild or no crow's feet lines at day 30 after treatment. In the same studies, physicians assessed 25% of people to have at least moderate improvement on day 120.

20 Units • *Moderate to severe* **forehead lines**


Real patient. Results may vary.
Photos taken at maximum eyebrow elevation before and 2 days after treatment with BOTOX® Cosmetic.




Real patient. Results may vary.
Photos taken at maximum eyebrow elevation 30 and 120 days after treatment with BOTOX® Cosmetic. In 2 clinical studies of healthy adults, 61% and 46% had ≥ 2-grade improvement at day 30. In the same studies, physicians assessed 40% of people to have at least moderate improvement on day 120.

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

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).

Please see additional Important Safety Information throughout for *Natrelle*® and BOTOX® Cosmetic, including Boxed Warnings, and JUVÉDERM® Collection of Fillers.

Which Treatment

IS RIGHT FOR YOU?

A consultation with your provider will determine which treatment is right for you!

Plump the Lips with

JUVÉDERM®
ULTRA XC

JUVÉDERM® is the No. 1 chosen dermal filler collection worldwide† with 6 unique and long-lasting fillers specifically designed for different areas of the face.

JUVÉDERM® Ultra XC is a smooth hyaluronic acid injectable gel that temporarily adds more fullness and plumps thin lips and is also approved for smoothing and correcting moderate to severe facial wrinkles and folds such as the smile lines around the nose and mouth. You may see long-lasting results for up to 1 year.‡

†Based on a January 2023 healthcare provider survey in the US (n = 959) and a Q1 2022 healthcare provider survey (n = 1799).
‡With optimal treatment.

Katie – JUVÉDERM® Ultra XC Before-and-After Photos

Actual patient. Results may vary.
Unretouched photos taken before treatment and 1 month after treatment with 1 mL of JUVÉDERM® Ultra XC in the lips.

Jaskiran – JUVÉDERM® Ultra XC Before-and-After Photos

Actual patient. Results may vary.
Unretouched photos taken before treatment and 1 month after treatment with 1 mL of JUVÉDERM® Ultra XC in the lips.

JUVÉDERM® Injectable Gel Fillers Important Information
APPROVED USES

JUVÉDERM® VOLUX® XC injectable gel is for deep injection to improve moderate to severe loss of jawline definition in adults over the age of 21.

JUVÉDERM® VOLUMA® XC injectable gel is for deep injection in the cheek area to correct age-related volume loss and for augmentation of the chin region to improve the chin profile in adults over 21.

JUVÉDERM® VOLLURE® XC, JUVÉDERM® Ultra Plus XC, and JUVÉDERM® Ultra 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 injectable gel is also 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 correction of perioral lines, and for injection into the undereye hollows to improve the appearance of undereye hollows in adults over the age of 21.

IMPORTANT SAFETY INFORMATION

Are there any reasons why I should not receive any JUVÉDERM® formulation?

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

What warnings should my doctor advise me about?

- One of the risks with using dermal fillers is the unintentional injection into a blood vessel. The chances of this happening are very small, but if it does happen, 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 of the skin. Most of these events are irreversible.
- If you have changes in your vision, signs of a stroke (including sudden difficulty speaking, numbness or weakness in your face, arms or legs, difficulty walking, face drooping, severe headache, dizziness, or confusion), white appearance of the skin, or unusual pain during or shortly after treatment, you should notify your health care practitioner immediately.

Please see additional Important Safety Information throughout for *Natrelle*® and BOTOX® Cosmetic, including Boxed Warnings, and JUVÉDERM® Collection of Fillers.

Actual *Natrelle*® patients.
Individual results may vary.

HOW IT WORKS

1. Join Allē.
   - Download the Allē app.
   - You must be an Allē Member to claim your complimentary *Natrelle* Perks® offer.
2. Confirm.
   - First, confirm your *Natrelle*® breast augmentation was logged in your Allē transaction history, which you can find in your Allē Wallet.
3. Select Your Offer.
   - Text "**PERKS**" to "**65190**" to opt in for Allē text messages.
   - Two text messages will follow.
   - Click on the second link to select your complimentary treatment offer.
   - Your selected treatment offer will be deposited into your Allē Wallet within 48 hours and cannot be changed.
4. Redeem.
   - Once the complimentary treatment offer has been added to your Allē Wallet, book an appointment with a participating Allē provider.
   - You must redeem your offer within 6 months.

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

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.

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 accompanying Summary of Information about BOTOX® Cosmetic.

JUVÉDERM® Injectable Gel Fillers IMPORTANT SAFETY INFORMATION (continued)

What warnings should my doctor advise me about? (continued)

- The use of dermal fillers where skin sores, pimples, rashes, hives, cysts, or infections are present should be postponed, as this may delay healing or make skin problems worse.
- The effectiveness of removal of any dermal filler has not been studied.

What precautions should my doctor advise me about?

- JUVÉDERM® VOLBELLA® XC should only be injected into undereye hollows by doctors who have completed the necessary training for this treatment area. To find a doctor, visit Juvederm.com/find-a-specialist. Doctors who complete the training will be listed with a symbol
- The safety of these products for use during pregnancy or while breastfeeding has not been studied
- The safety of JUVÉDERM® VOLUMA® XC has not been studied in patients under 35 years or over 65 years for cheek augmentation, or under 22 years and over 80 years for chin augmentation. The safety of JUVÉDERM® VOLUX® XC, JUVÉDERM® VOLLURE® XC and JUVÉDERM® VOLBELLA® XC has not been studied in patients under 22 years, and the safety of JUVÉDERM® Ultra Plus XC and JUVÉDERM® Ultra XC has not been studied in patients under 18 years
- The safety and effectiveness of treatment with JUVÉDERM® products in anatomical regions outside of their approved uses have not been established in clinical studies

Please see additional Important Safety Information throughout for *Natrelle*® and BOTOX® Cosmetic, including Boxed Warnings, and JUVÉDERM® Collection of Fillers.

Enjoy the Full

REWARDS OF Allē

Allē is the official loyalty program for *Natrelle*®, BOTOX® Cosmetic, the JUVÉDERM® Collection of Fillers, and more. Create an account today to earn points on 50+ products and treatments at your participating plastic surgeon, dermatologist, and med spa.

Visit Alle.com for more information.

Joining Allē is easy

To get started, visit Alle.com or download the app.

Once registered, you'll gain access to:

- Exclusive rewards
- Curated content
- Treatment information

Scan to download the Allē app.

Download on the App Store

GET IT ON Google Play

JUVÉDERM® Injectable Gel Fillers IMPORTANT SAFETY INFORMATION (continued)

What precautions should my doctor advise me about? (continued)

- If you have a history of excessive scarring (thick, hard scars) or pigmentation disorders, treatment in these patients has not been studied and may result in additional scars or changes in pigmentation
- If you are planning other procedures including laser treatments or a chemical peel, there is a possible risk of inflammation at the treatment site if these procedures are performed closely before or after JUVÉDERM® injectable gel treatment
- Tell your doctor if you are on therapy used to reduce your body's natural defense system (such as steroids, chemotherapy, and medicines to treat autoimmune diseases, HIV, and AIDs), as these may increase your risk of infection; and medications that can prolong bleeding (such as aspirin, ibuprofen, or other blood thinners), as these may result in increased bruising or bleeding at the injection site.
- Avoid applying makeup for 12 hours after treatment and minimize strenuous exercise, exposure to extensive sun or heat, and alcoholic beverages within the first 24 hours following treatment, as these may cause temporary redness, swelling, and/or itching at the injection site
- JUVÉDERM® VOLUMA® XC was not studied in patients with significant loose skin of the chin, neck, or jaw
- The effect of JUVÉDERM® VOLUMA® XC injection into the chin on facial hair growth has not been studied
- Patients who experience skin injury near the site of JUVÉDERM® VOLUMA® XC injection may be at a higher risk for adverse events
- Tell your doctor if you have already been injected with dermal fillers in the same area as the one(s) you are about to be treated for. This information helps your doctor decide when and whether you should get treatment

What are possible side effects of treatment?

The most commonly reported side effects with JUVÉDERM® injectable gels were redness, swelling, pain, tenderness, firmness, lumps/bumps, bruising, discoloration, and itching. For JUVÉDERM® VOLBELLA® XC, dryness was also reported.

These side effects are consistent with other facial injection procedures and most will resolve within 30 days. Your doctor may choose to treat side effects persisting longer with antibiotics, steroids, or hyaluronidase (an enzyme that breaks down hyaluronic acid).

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

To report a side effect with any product in the JUVÉDERM® Collection, please call the Allergan® Product Support Department at 1-877-345-5372. Please also visit Juvederm.com or talk to your doctor for more information.

Products in the JUVÉDERM® Collection are available only by a licensed physician or properly licensed practitioner.

Please see additional Important Safety Information throughout for *Natrelle*® and BOTOX® Cosmetic, including Boxed Warnings, and JUVÉDERM® Collection of Fillers.

FAQs

How do I qualify for the complimentary aesthetic treatment offer of BOTOX® Cosmetic or JUVÉDERM® Ultra XC treatment?

Allē Members who undergo breast augmentation with *Natrelle*® gel implants may qualify to receive either 1 complimentary 50-unit vial of BOTOX® Cosmetic or up to two .55 mL syringes of JUVÉDERM® Ultra XC at a participating Allē provider. Injection fees may apply.

Is there an injection fee for the complimentary product?

Injection fees may apply and are up to the discretion of your provider. Allergan Aesthetics is not responsible for any associated injection costs. Please ask at a participating provider's office.

How do I know if I have the complimentary treatment offer in my Allē Wallet?

Your surgeon must submit your breast augmentation in Allē. Text "**PERKS**" to "**65190**" to claim your complimentary treatment offer. It can take up to 48 hours for your offer to be deposited into your Allē Wallet.

Do I need to be an Allē Member to receive the complimentary treatment offer?

Yes, you need to be an Allē Member. Visit Alle.com to create a free account today.

How do I redeem my Allē points?

Allē points can be used on any Allergan Aesthetics products or treatments. Points can only be redeemed at a participating Allē provider. A consultation may be required to determine if a treatment is right for you. When you've decided on a treatment with your provider, ask to apply your available points at checkout for instant savings.

Who can I contact if I have additional questions?

Please contact Allē Member Support at 1.888.912.1572, Monday–Friday, 8 am–6 pm CT.

Natrelle® **IMPORTANT SAFETY INFORMATION (continued)**

Who can get breast implants?

Natrelle® Breast Implants are approved for the following:

- **Breast augmentation for women at least 22 years old for silicone-filled implants and for women at least 18 years old for saline-filled implants.** Breast augmentation includes primary breast augmentation to increase the breast size and revision surgery to correct or improve the result of a primary breast augmentation
- **Breast reconstruction.** This includes primary breast reconstruction to replace breast tissue that has been removed due to cancer or trauma or that has failed to develop properly due to a severe breast abnormality. This also includes revision surgery to correct or improve the result of a primary breast reconstruction

Who should NOT get breast implants?

Breast implant surgery should NOT be performed in:

- 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 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 (eg, lupus and scleroderma)

Please see additional Important Safety Information throughout for *Natrelle*® and BOTOX® Cosmetic, including Boxed Warnings, and JUVÉDERM® Collection of Fillers.

To learn more about *Natrelle* PERKS®, ask your provider or scan here.

To learn more about Allē, scan here.

***Natrelle*® IMPORTANT SAFETY INFORMATION (continued)**

What should I tell my doctor? (continued)

- A weakened immune system (eg, taking medications to decrease the body's immune response)
- Planned chemotherapy or radiation therapy 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
- Those 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

***Natrelle* PERKS® Terms and Conditions**

- Allē Members who undergo breast augmentation with *Natrelle*® gel implants ("Member") may qualify to receive either (a) one (1) complimentary treatment of BOTOX® Cosmetic (onabotulinumtoxinA) up to 50 units OR (b) one (1) complimentary treatment of JUVÉDERM® Ultra XC up to two (2) .55 mL syringes at participating Allē provider offices only. Allergan Aesthetics is not responsible for any associated injection costs.
- Members enrolled in Medicare, Medicaid, or other federal or state healthcare programs are not eligible for this offer.
- Member must claim offer in the Allē app via text message link within 6 months of their *Natrelle*® breast augmentation and select their complimentary treatment.
- Once claimed, the selected complimentary treatment cannot be changed and the offer must be redeemed within 6 months of the offer being deposited into Member's Allē Wallet. Offer expires 6 months after issue date into Allē Wallet.
- Limit one 1 per Member.
- A healthcare provider will determine if Member is an appropriate candidate for a BOTOX® Cosmetic or JUVÉDERM® Ultra XC treatment.
  – If Member is an appropriate candidate, offer can be redeemed at a participating provider's office.
- Standard Allē Loyalty Program Terms and Conditions apply.
- Members will earn Allē points on all qualifying Earnings Eligible Product purchases, subject to applicable earnings caps.
- The value of this offer cannot be redeemed or exchanged for cash.
- Offer cannot be applied to past transactions.
- Offer cannot be combined with other Allē offers on BOTOX® Cosmetic or the JUVÉDERM® Collection of Fillers but can be combined with Allē and Allē brand-specific gift cards, Allē points offers, including Double Points offers, and other Allē brand-specific offers.
- The complete value of this offer must be used in a single transaction.
- If you have questions, please contact Allē Customer Support at 1-888-912-1572, Monday - Friday, 8 am– 6 pm CT.
- Allergan Aesthetics, an AbbVie company, reserves the right to alter or cancel this offer at any time.

What else should I consider?

- There is a Boxed Warning for breast implants. Please see bold text at beginning
- Many changes to your breasts following implantation are irreversible. If you later choose to have your implants removed and not replaced, you may experience dimpling, puckering, wrinkling, or other cosmetic changes, which may be permanent
- Breast implantation is likely not a one-time surgery. The longer implants are in place, the greater the potential risk for complications. You will likely need additional surgeries on your breasts due to complications or unacceptable cosmetic results. Thus, you should also consider the complication rates for later (revision) surgery since you may experience these risks in the future
- Cancer treatments and surgery will affect the outcome and timing of breast reconstruction
- Breast implants may affect your ability to breastfeed, either by reducing or eliminating milk production

Please see additional Important Safety Information throughout for *Natrelle*® and BOTOX® Cosmetic, including Boxed Warnings, and JUVÉDERM® Collection of Fillers.

***Natrelle*® IMPORTANT SAFETY INFORMATION (continued)**

What else should I consider? (continued)

- Rupture of a silicone-filled breast implant is most often silent. Even if you have no symptoms, you should have your first ultrasound or MRI at 5 to 6 years after your initial implant surgery and then every 2 to 3 years thereafter regardless of whether your implants are for augmentation or reconstruction. If you have symptoms of or uncertain ultrasound results for breast implant rupture, an MRI is recommended. Additional imaging may be required depending on your medical history and status. The health consequences of a ruptured silicone gel-filled breast implant have not been fully established
- Routine screening mammography for breast cancer will be more difficult, and implants may rupture during the procedure. Perform self-examination every month for cancer screening and ask your surgeon to help you distinguish the implant from your breast tissue. Lumps, persistent pain, swelling, hardening, or changes in implant shape should be reported to your surgeon and possibly evaluated with imaging

What are key complications with breast implants?

Key complications include reoperation, implant removal with or without replacement, implant rupture with silicone-filled implants, implant deflation with saline-filled implants, and capsular contracture (severe scar tissue around the implant). Other complications include breast pain, swelling, asymmetry, wrinkling/rippling, implant malposition nipple complications, hypertrophic scarring, and implant palpability/visibility.

Talk to your doctor about other complications.

For more information, see the patient brochures at www.allergan.com/products.

To report a problem with *Natrelle*® Breast Implants, please call Allergan® at 1-800-624-4261.

The sale and distribution of *Natrelle*® Breast Implants is restricted to licensed physicians who provide information to patients about the risks and benefits of breast implant surgery.

Learn more at natrelle.com/rewards

Natrelle® Allē® BOTOX® COSMETIC onabotulinumtoxinA injection JUVÉDERM® ULTRA XC

Allergan Aesthetics
an AbbVie company