What’s gummy about the gel in Natrelle INSPIRA® Cohesive?
The technical term for what’s inside Natrelle INSPIRA® Cohesive is highly cohesive gel—that means that the gel has a gummy-like consistency that sticks together, holds its shape, and doesn’t squish as much as other less cohesive silicone gels.

How is Natrelle INSPIRA® Cohesive different from other implants?
Natrelle INSPIRA® Cohesive is the most cohesive (gummiest) round gel breast implant in the US. This highly cohesive, gummy implant is designed to maintain its upper pole fullness. In other words, when you hold the implant upright in your hand, it’s designed to keep its fullness at the top.

‡Significance inside the body has not been established.
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.

- **Breast reconstruction.** Breast reconstruction includes primary 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. Breast reconstruction also includes revision surgery to correct or improve the result of a primary breast reconstruction 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 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. For more information see the patient brochures at www.allergan.com/labeling/usa.htm. To report a problem with Natrelle® Breast Implants, please call Allergan at 1-800-433-8871.

Natrelle® Breast Implants are available by prescription only.
What is AlloDerm SELECT™ Regenerative Tissue Matrix?
AlloDerm SELECT™ is a soft tissue matrix that helps to reinforce weakened or thin tissue. Your surgeon may recommend AlloDerm SELECT™ Regenerative Tissue Matrix (an acellular dermal matrix, or ADM) as part of your breast reconstruction procedure.

Is AlloDerm SELECT™ right for me?
Only your plastic surgeon can determine if AlloDerm SELECT™ is right for you. Every woman’s journey is different, and what might be a good fit for one person, may not be right for the next. Ask your care team which products are best for your unique situation.
Indication and Important Safety Information for AlloDerm SELECT™ Regenerative Tissue Matrix

AlloDerm SELECT™ Regenerative Tissue Matrix is an acellular dermal matrix that comes from donated human tissue and is processed to remove cells while maintaining structure. AlloDerm SELECT™ RTM is intended for the replacement or repair of inadequate or damaged soft tissue. AlloDerm SELECT™ RTM is not a dural substitute or for veterinary use.

Each package of AlloDerm SELECT™ RTM is for one-time use in one patient. Patients who are sensitive to any of the antibiotics listed on the product package or Polysorbate 20 should not receive AlloDerm SELECT™ RTM.

AlloDerm SELECT™ RTM is available only through a licensed physician.

Potential risks of surgical procedures associated with the use of a tissue matrix include, but are not limited to: wound or systemic infection, seroma, dehiscence, hypersensitive, allergic or other immune response, sloughing or failure of the tissue matrix, and disease transmission.

This information is not intended to replace a discussion with your surgeon. It does not describe all the potential risks associated with the use of tissue matrix in surgical procedures. Every patient’s situation is different, so please consult with your surgeon to determine if the use of AlloDerm SELECT™ RTM is right for you.

Please see AlloDerm SELECT™ RTM Instructions for Use.
How can fat transfer with REVOLVE™ System help me achieve my desired breast reconstruction results?

Women who undergo a mastectomy are left with very thin tissue. Your surgeon may suggest fat transfer with REVOLVE™ System to replace the volume you've lost and provide additional fullness.

**Shape your breasts naturally with REVOLVE™ System**

REVOLVE™ System is a system for fat transfer that allows your surgeon to collect fat from one area of your body (such as your abdomen) and process that fat for transfer to your breasts. After processing with REVOLVE™ System, surgeons can use this purified fat to help replace the volume and shape that you’ve lost after your mastectomy.

Ask your surgeon if REVOLVE™ System is right for you.

- Fat transfer can enhance your overall breast shape by adding fullness where you need it the most
- Closed system design cleans and purifies fat in one sterile environment from start to finish

**Your questions, answered**

1. How can fat transfer with REVOLVE™ System help me achieve my desired breast reconstruction results?

   Women who undergo a mastectomy are left with very thin tissue. Your surgeon may suggest fat transfer with REVOLVE™ System to replace the volume you’ve lost and provide additional fullness.

Ask your surgeon for more information about fat transfer with REVOLVE™ System
Indication and Important Safety Information for REVOLVE™ System

The 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 physician 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 undercorrection of the treatment site, tissue lumps, bleeding, and scarring.

The REVOLVE™ System is available only through a licensed physician.

This information is not intended to replace a discussion with your surgeon. It does not describe all the potential risks associated with fat grafting procedures. Every patient’s situation is different, so please consult with your surgeon to determine if the use of REVOLVE™ System is right for you.

Please see REVOLVE™ System Instructions for Use and User Manual.