BEFORE & AFTER

See real results in patients treated with Radiesse®, Radiesse (+), and Belotero Balance® (+)

RADIESSE® | RADIESSE®+
BELOTERO BALANCE®+

BEFORE

AFTER

MIDFACE & LOWER FACE

TREATMENT PLAN

Alison, age 60. Five weeks after receiving:

- 2 syringes of Radiesse (+) to treat the jawline
- 2 syringes of Radiesse (+) in the midface to efface the lower face
- 2 syringes of Radiesse to treat nasolabial folds
- 2 syringes of Radiesse to treat pre-jowl folds
- 1 syringe of Belotero Balance (+) to treat nasolabial folds in follow-up appointment

Courtesy of Iani Silveira, MSN, FNP-BC, CPN | Bella Vida Aesthetics & Wellness
Actual patient & model. Individual results may vary.

What are RADIESSE® and RADIESSE® (+)?
RADIESSE® and RADIESSE® (+) are dermal fillers that are used for smoothing moderate to severe facial wrinkles and folds, such as nasolabial folds (the creases that extend from the corner of your nose to the corner of your mouth). RADIESSE® is also used for correcting volume loss in the back of the hands. RADIESSE® (+) is also used for improving moderate to severe loss of jawline contour in adults over 21 years old.

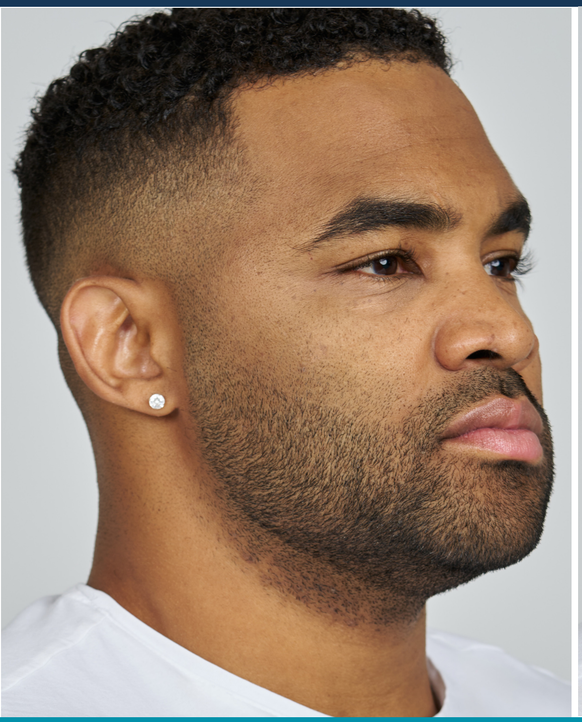
What is BELOTERO BALANCE® (+)?
BELOTERO BALANCE is a prescription injection that is approved to temporarily smooth out and fill in moderate to severe nasolabial folds (the folds or wrinkles that go from the side of the nose to the corner of the mouth) and improve the appearance of under-eye hollows in adults over the age of 21.

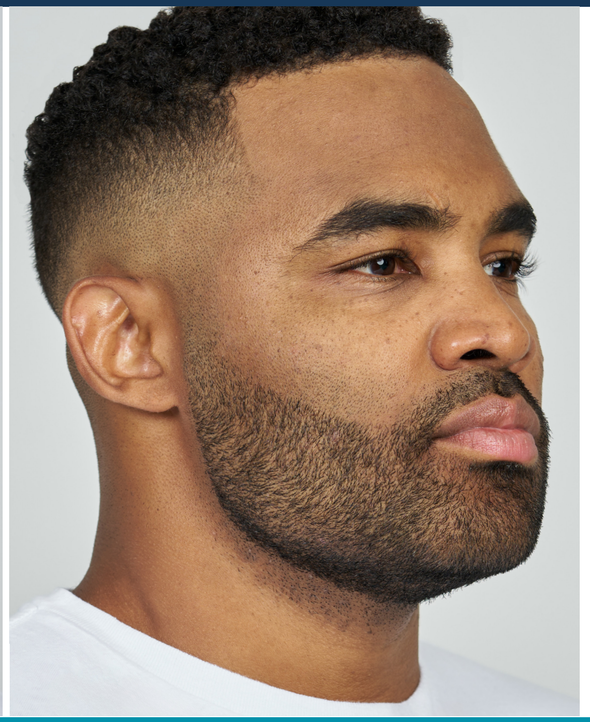
Please see Important Consumer Safety Information throughout and full Instructions for Use, including Patient Information Guides, at Belotero.com and Radiesse.com.

BEFORE & AFTER

See real results in patients treated with Radiesse® (+)

RADIESSE®+





BEFORE | IMMEDIATELY AFTER | AFTER

MIDFACE, LOWER FACE & JAWLINE

TREATMENT PLAN

Brandon, age 37. Five weeks after receiving:

- 6 syringes of Radiesse (+) to treat the jawline
- 2 syringes of Radiesse (+) in the midface to efface the lower face
- 1 syringe of Radiesse (+) to treat pre-jowl folds

Courtesy of Iani Silveira, MSN, FNP-BC, CPN | Bella Vida Aesthetics & Wellness

Actual patient & model. Individual results may vary.

RADIESSE® and RADIESSE® (+)
IMPORTANT SAFETY INFORMATION

Who should not use RADIESSE® or RADIESSE® (+)?

You should not use RADIESSE® or RADIESSE® (+) if you have an allergy to any component of the product, if you have a history of severe allergies, if you have a bleeding disorder, or if you are pregnant or breastfeeding. You should not use RADIESSE® (+) if you have an allergy to lidocaine or medicines like it.

BELOTERO BALANCE® (+)
IMPORTANT SAFETY INFORMATION

Who should not use BELOTERO BALANCE?

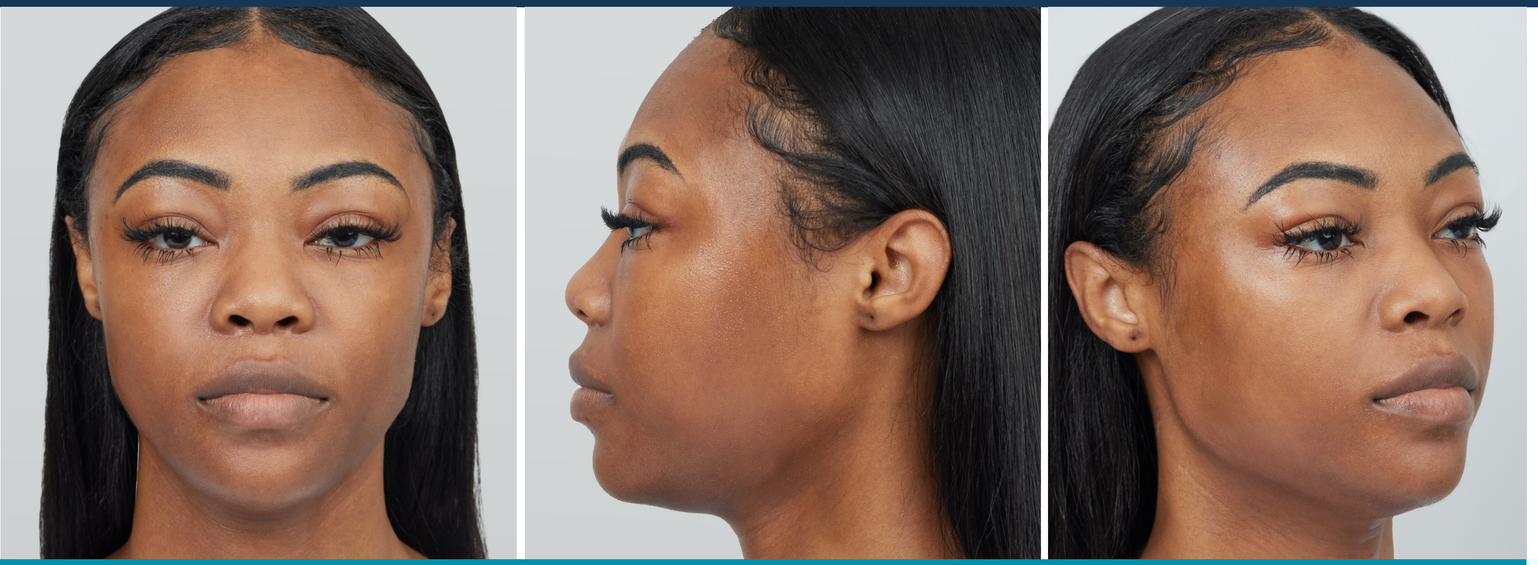
BELOTERO BALANCE should not be used in patients with a history of or presence of multiple or severe allergies, including those with a history of anaphylaxis. BELOTERO BALANCE should not be used in patients with allergies to gram-positive bacterial proteins.

Please see Important Consumer Safety Information throughout and full Instructions for Use, including Patient Information Guides, at Belotero.com and Radiesse.com.

BEFORE & AFTER

See real results in patients treated with Radiesse® (+)

RADIESSE®+



BEFORE

AFTER

JAWLINE

TREATMENT PLAN

Stonie, age 26. Four weeks after receiving:

- 4 syringes of Radiesse (+) to treat the jawline

Courtesy of Cheryl Habeeb, CCRN-K, DNP | Pura Medical Aesthetics

Actual patient & model. Individual results may vary.

What is the most important information I should know about RADIESSE® and RADIESSE® (+)?

One of the risks with using these products is 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. 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 Important Consumer Safety Information throughout and full Instructions for Use, including Patient Information Guide, at Radiesse.com.

BEFORE & AFTER

See real results in patients treated with Radiesse®

RADIESSE®

BEFORE

AFTER

HANDS

TREATMENT PLAN

Tabetha, age 57. Five weeks after receiving:

- 2 syringes of Radiesse to treat the hands (1 syringe per hand)

Courtesy of Iani Silveira, MSN, FNP-BC, CPN | Bella Vida Aesthetics & Wellness
Actual patient & model. Individual results may vary.

RADIESSE® and RADIESSE® (+) IMPORTANT SAFETY INFORMATION (cont)

If you have a history of herpes, you may experience a herpes breakout after receiving RADIESSE® or RADIESSE® (+).
Injection in the back of the hand may result in temporary difficulty performing activities. RADIESSE® may cause nodules, bumps or lumps in the back of the hand and can last up to 1 year.

Please see Important Consumer Safety Information throughout and full Instructions for Use, including Patient Information Guide, at Radiesse.com.

RADIESSE and RADIESSE (+) Important Consumer Safety Information

What are RADIESSE® and RADIESSE® (+)?

RADIESSE® and RADIESSE® (+) are dermal fillers that are used for smoothing moderate to severe facial wrinkles and folds, such as nasolabial folds (the creases that extend from the corner of your nose to the corner of your mouth). RADIESSE® is also used for correcting volume loss in the back of the hands. RADIESSE® (+) is also used for improving moderate to severe loss of jawline contour in adults over 21 years old.

RADIESSE® and RADIESSE® (+) IMPORTANT SAFETY INFORMATION

Who should not use RADIESSE® or RADIESSE® (+)?

You should not use RADIESSE® or RADIESSE® (+) if you have an allergy to any component of the product, if you have a history of severe allergies, if you have a bleeding disorder, or if you are pregnant or breastfeeding. You should not use RADIESSE® (+) if you have an allergy to lidocaine or medicines like it.

What is the most important information I should know about RADIESSE® and RADIESSE® (+)?

One of the risks with using these products is 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. 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.

As with all procedures that involve an injection through the skin, there is a risk of infection.

Do not use RADIESSE® or RADIESSE® (+) if you have a skin infection until it has healed.

It is not known if RADIESSE® or RADIESSE® (+) is safe or effective in the lips, or in the area around the eyes.

It is not known if RADIESSE® or RADIESSE® (+) is safe or effective beyond 3 years in the face and 1 year in the hand.

Injection of RADIESSE® (+) into the jawline may temporarily alter jaw function.

The microspheres in RADIESSE® and RADIESSE® (+) can be seen in X-rays and CT Scans. It is very important that you tell your health care provider that you have had RADIESSE® or RADIESSE® (+) dermal filler.

If you have a history of herpes, you may experience a herpes breakout after receiving RADIESSE® or RADIESSE® (+).

Injection in the back of the hand may result in temporary difficulty performing activities. RADIESSE® may cause nodules, bumps or lumps in the back of the hand and can last up to 1 year.

You should minimize strenuous activity and avoid extensive sun or heat exposure for about 24 hours after treatment and until any swelling or redness has resolved.

What should I tell my doctor before using RADIESSE® or RADIESSE® (+)?

Tell your health care provider if you are taking blood thinners or medicines that can interfere with the clotting of blood, such as aspirin or warfarin. These medicines might make it more likely that you will experience bruising or bleeding at the injection site. Tell your health care provider if you have any diseases, injuries or disabilities of the hand, if you have a history forming large, raised scars or if you have had any other skin treatments such as skin peels.

What are the most common adverse events with RADIESSE® or RADIESSE® (+)?

The most common adverse events seen in clinical studies of RADIESSE® used in the hands include bruising, redness, swelling, pain, itching, nodules or bumps/lumps, difficulty performing activities, loss of sensation and other local side effects. The most common adverse events seen in clinical studies of RADIESSE® or RADIESSE® (+) used in the face include bruising, redness, swelling, pain, itching, difficulty chewing, and other local side effects.

Delayed-onset inflammation near the site of injection is one of the known adverse events associated with dermal fillers. Cases of delayed-onset inflammation have been reported to occur at the treatment site following viral or bacterial illnesses or infections, vaccinations, or dental procedures. Typically, the reported inflammation was responsive to treatment or resolved on its own.

These are not all of the possible side effects with RADIESSE® or RADIESSE® (+). Merz collects information about adverse events seen with RADIESSE® and RADIESSE® (+) outside of clinical studies. These events are included in the RADIESSE® and RADIESSE® (+) Patient Information Guide based on an assessment of seriousness and potential causal relationship to RADIESSE® or RADIESSE® (+). Please see the Patient Information Guide available at www.radiesse.com for list of these events. Tell your health care provider about any side effects that bother you or do not go away.

Important: For full safety information, please visit www.Radiesse.com or call MyMerz Solutions at 1-844-469-6379.

RADIESSE® and RADIESSE® (+) are available by prescription only.

Please see full Instructions for Use and Patient Information Guide at Radiesse.com.

BELOTERO BALANCE (+) Important Consumer Safety Information

What is BELOTERO BALANCE® (+)?

BELOTERO BALANCE is a prescription injection that is approved to temporarily smooth out and fill in moderate to severe nasolabial folds (the folds or wrinkles that go from the side of the nose to the corner of the mouth) and improve the appearance of under-eye hollows in adults over the age of 21.

Who should not use BELOTERO BALANCE?

BELOTERO BALANCE should not be used in patients with a history of or presence of multiple or severe allergies, including those with a history of anaphylaxis. BELOTERO BALANCE should not be used in patients with allergies to gram-positive bacterial proteins.

What is the most important information I should know about BELOTERO BALANCE?

Introduction of BELOTERO BALANCE into the vasculature may lead to embolization, occlusion of the vessels, ischemia, or infarction. Take extra care when injecting soft tissue fillers, for example inject the product slowly and apply the least amount of pressure necessary. Rare but serious adverse events associated with the intravascular injection of soft tissue fillers in the face have been reported and include temporary or permanent vision impairment, blindness, cerebral ischemia or cerebral hemorrhage, leading to stroke, skin necrosis, and damage to underlying facial structures. Immediately stop the injection if patient exhibits any of the following symptoms, including changes in vision, signs of a stroke, blanching of the skin, or unusual pain during or shortly after the procedure. Patients should receive prompt medical attention and possibly evaluation by an appropriate health care practitioner specialist should an intravascular injection occur.

As with all events that involve an injection through the skin, there is a risk of infection. Laser treatments or chemical peels or any other treatments that affect the skin can increase the risk of infection. Do not use BELOTERO BALANCE if you have a skin inflammation or a skin infection. Do not use until the infection is healed.

Patients getting BELOTERO BALANCE may have an injection site reaction. These reactions can include inflammation and usually last less than seven days.

For approximately 24 hours after treatment, avoid:

- strenuous activity
- extensive sun or heat exposure
- aspirin or non-steroidal anti-inflammatory drugs
- alcoholic beverages

Exposure to any of the above can cause temporary redness, swelling, and/or itching at the injection site.

It is not known how BELOTERO BALANCE will work in areas of the face other than the smile lines.

It is not known how BELOTERO BALANCE will work in women who are pregnant or breastfeeding or people who are less than 21 years of age.

What should I tell my doctor before injections with BELOTERO BALANCE?

Tell your doctor if you are taking medicines that affect blood clotting, like aspirin, an NSAID or warfarin. These medicines may put you at an increased risk of bruising or bleeding at the treatment site. Tell your doctor if you have a skin reaction like cold sores, cysts, pimples, rashes, hives, or an infection. Treatment with BELOTERO BALANCE should be delayed until the reaction goes away. Tell your doctor if you are taking medicines that affect your immune system.

What are the most common adverse events seen with BELOTERO BALANCE?

The most common adverse events seen in clinical studies with BELOTERO BALANCE were swelling, bruising, redness, hardening of the skin, pain, altered color, or itching. Other side effects that have occurred in clinical studies of BELOTERO BALANCE include headache, swelling of the side of the nose, moderate cold sore, lip numbness, and lip dryness. Side effects were often mild to moderate and often resolved within 7 days.

Delayed-onset inflammation near the site of injection is one of the known adverse events associated with dermal fillers. Cases of delayed-onset inflammation have been reported to occur at the treatment site following viral or bacterial illnesses or infections, vaccinations, or dental procedures. Typically, the reported inflammation was responsive to treatment or resolved on its own.

Information on adverse events from post-market surveillance of BELOTERO BALANCE are included in the Package Insert (PI) and Patient Information Guide (PIG) based on an assessment of seriousness and potential causal relationship to BELOTERO BALANCE. Please see the PI and PIG available on www.belotero.com for a complete list of these events.

Important: For full safety information, please visit www.belotero.com or call MyMerz Solutions at 1-844-469-6379 by email AxUS-adverse.events@merz.com.

Rx only

Please see full Instructions for Use and Patient Information Guide at Belotero.com.

MERZ AESTHETICS® | confidence to be