

INFORMED CONSENT - RADIESSE INJECTABLE

INTRUCTIONS

This consent form is designed to give you the information you need to make an informed decision about whether or not to undergo treatment with the dermal filler Radiesse. If you have any questions, please ask the clinician at Alamo Hills Advanced Aesthetics & Laser Center.

INTRODUCTION

Radiesse treatments involve injections that are planted intradermally through a fine gauge needle into the treated area. Radiesse is comprised of calcium hydroxylapatite (CaHA) microspheres. Multiple treatments may be necessary to achieve desired results. Treatments generally last 9-12 months. Touch up treatments may be necessary to maintain desired results.

ALTERNATIVE TREATMENT

Alternatives to Radiesse treatments include, but are not limited to, other dermal fillers (e.g. collagen, fat, synthetic polymers), laser resurfacing, surgical facelift, lasers for skin laxity, or no treatment at all.

PATIENTS THAT MAY NOT BE ELIGIBLE FOR RADIESSE TREATMENTS

Patients with the following conditions may not receive Radiesse treatments: previous allergic reactions to injectable products, history of a serious allergic reaction (anaphylactic), multiple severe allergies, abnormal raised scarring or keloid formation, active inflammation or infection in the treatment area (e.g. pimples, rash, hives), pregnancy, or nursing.

Certain conditions require caution with injectable fillers and may preclude a patient from receiving the treatment: poor healing (due to diabetes or other conditions), long-term use of Prednisone or other steroid therapy. Recurrent viral infections such as herpes simplex (cold sores) may be activated by Radiesse treatments. The physician, registered nurse, physician assistant, or nurse practitioner must be notified of these conditions prior to treatments.

RISKS

The possible risks, side effects, and complications with Radiesse injectable include, but are not limited to:

- 1. Pain and tenderness during and after treatments at/around the treated site which typically resolves within a few days to a week.
- 2. Redness and swelling at/around the injection site is common. Itchiness may also occur. These reactions are generally present immediately after treatment and lessen or disappear within a few days to 1 week. Some patients may experience prolonged swelling and/or tenderness/pain at the injection site lasting up to 2 weeks. Some patients may experience a delayed onset of these symptoms up to several weeks after treatment. On rare occasions, pustules (acne-like lesions) may form. The



- physician must be notified if symptoms persist for more than 1 week, pustules are present, or symptoms appear in a delayed fashion after treatment.
- 3. Bruising which usually resolves within 1-2 weeks after the injection. Patients taking medications that interfere with coagulation (e.g. aspirin, ibuprofen) have an increased risk of bruising and bleeding.
- 4. Infection at the treated site.
- 5. Although rare, local tissue damage can occur with skin breakdown, scab formation, and/or scarring in the treated area.
- 6. Visible raised areas and lumpiness at/around the treated site grayish discoloration of the skin. These symptoms may persist from a few weeks to several months and may be permanent (rarely).
- 7. Failure to reduce a contour defect or wrinkle (under correction) or overcorrection. Placement of filler adjacent to or outside the desired treatment area; undesired changes in facial contour. Asymmetry, where the correction on one side may be different from the correction on the other side of the face. Swelling at time of injection may create the appearance of asymmetry or under correction which usually resolves as described above. However, you may need to return for additional treatment if under correction of asymmetry persists.
- 8. Radiesse injectable may have an unpredictable duration of action and may not last as long as anticipated or may persist in some areas longer than anticipated.
- 9. All the risks of Radiesse injectable use may not be known. Alamo Hills Advanced Aesthetics & Laser Center is not responsible for any Radiesse injectable risk or unforeseen complication not yet discovered or not commonly known.

DISCLAIMER

Informed-consent documents are used to communicate information about the proposed treatment of a disease or condition along with disclosure of risks and alternative forms of treatment. The informed-consent process attempts to define principles of risk disclosure that should generally meet the needs of most patients in most circumstances. However, informed-consent documents should not be considered all inclusive in defining other methods of care and risks encountered. Your physician may provide you with additional or different information which is based on all the facts in your particular case and the state of medical knowledge.

Informed-consent documents are not intended to define or serve as the standard of medical care. Standards of medical care are determined on the basis of all the facts involved in an individual case and are subject to change as scientific knowledge and technology advance and as practice patterns evolve.

It is important that you read the above information carefully and have all of your questions answered before signing the consent on the next page.



INFORMED CONSENT – RADIESSE INJECTABLE

I consent to administration of any related treatments that may be deemed necessary or advisable for my procedure. This includes, but is not limited to, local anesthetic such as anesthetic injections with lidocaine 1%-2% with or without epinephrine; and/or topical anesthetics such as benzocaine/lidocaine/tetracaine cream or ointment; and/or topical oral benzocaine preparations. The risks, side effects, complications of these anesthetics include, but are not limited to, skin irritation (itching or redness), lightheadedness, rapid heart rate, visual disturbances, and tongue numbness. I will inform the physician, registered nurse, physician assistant, or nurse practitioner immediately if I experience any of these symptoms. I do not have an allergy to lidocaine or anesthetics.
 No guarantees can be made or have been made that I will benefit from treatment or achieve a desired level of correction. There is no guarantee that wrinkles or folds will be reduced. I understand that I may require additional treatments to achieve correction.
 I understand that the fees for Radiesse injectable treatments are not covered by insurance. Should I require a touch-up treatment, I am responsible for the cost of that additional treatment.
 I understand that Radiesse will not correct the underlying cause of facial fat loss but will improve the appearance in the treated area.
 Microspheres in Radiesse can be seen on X-Rays & CT Scans. I understand I must inform my doctor and other health professionals that I have received Radiesse injections.
 If I take any blood thinners such as ibuprofen, aspirin, or herbal preparations prior to my procedure, I will advise my physician, registered nurse, physician assistant, or nurse practitioner. I understand that use of these medications may increase my risk of bruising.
 Possible side effects can include but are not limited to: Allergic reaction or infection. Bleeding, tenderness or pain, redness, bruising, scarring, Keloid formation/hypertrophic scarring or swelling at injection site.
 I understand if I have a history of Keloid formation or hypertrophic scarring, I must advise my physician and I am aware that will not be eligible for this treatment.
 I have fully read and agree to adhere to pre-treatment and post- treatment instructions. I understand that failure to carefully follow these instructions may affect my treatment outcome and increase the likelihood or severity of complications.



Witness Signature		Print Name	
Patient Signature		Print Name	Date
	O THE TREATMENT OR PR AM SATISFIED WITH THE	OCEDURE AND THE ABOVE EXPLANATION.	Listed Items
	It has been explained A. The above treath B. There may be alt	d to me in a way that I und nent or procedure to be ur ernative procedures or me the procedure or treatmen	erstand: ndertaken thods of treatment
	assistant, or nurse practice conter to perform Ratio all Hyaluronic acid		anced Aesthetics & Laser hts. This consent shall apply ent, I will call Alamo Hills
	I have fully read this of provided to me regal adequately informed benefits, limitations, a consider the informal answered to my satis	east 18 years of age. My sig consent form and understanding the proposed proced diabout the procedure involuternative treatments. I ha tion, and I have had all questaction. I understand and complications associated	nd the information dure. I have been olving the potential we had enough time to estions and concerns accept the risks, side
	responsibility to inforr	all of my medical history. I m and update the physicial r nurse practitioner of any o	
		apny; tnese pnotos wiil be i medical record with Alamo	•