CONSENT FOR HYALURONIC DERMAL FILLERS

Juvederm is a stabilized hyaluronic acid used to smooth moderate to severe facial wrinkles and folds around the nose and mouth or shape facial contours. Juvederm has been FDA approved for the cosmetic treatment of moderate to severe facial wrinkles and soft tissue depressions.

Hyaluronic acid is a naturally occurring substance that is found within soft tissue of all mammals. The hyaluronic acid in Juvederm is biocompatible and is a totally non-animal product; there is little risk of animal-based disease transmission or allergic reaction.

Juvederm injections are customized for every patient, depending on his or her particular needs. These can be performed in areas involving the face and eyelid region, forehead, and lips. Juvederm cannot stop the process of aging. It can however, temporarily diminish the look of wrinkles and soft tissue depressions. Juvederm injections may be performed as a singular procedure, in combination with other treatments such as BOTOX®, or as an adjunct to a surgical procedure. Juvederm injections require regional nerve blocks or local anesthetic injections to diminish discomfort. Soft tissue fillers, including Juvederm, produce temporary swelling, redness, and needle marks, which resolve after a few days time.

Continuing treatments are necessary in order to maintain the effect of Juvederm over time. Juvederm once injected will be slowly absorbed by the body. The length of effect for Juvederm injections is variable.

ALTERNATIVE TREATMENTS

Alternative forms of management include not treating the skin wrinkles or soft tissue depressions by any means. Improvement of skin wrinkles and soft tissue depressions may be accomplished by other treatments: laser treatments, chemical skin-peels, dermabrasion, or other skin procedures, alternative types of tissue fillers, or surgery such as a blepharoplasty, face or brow lift when indicated. Risks and potential complications are associated with alternative forms of medical or surgical treatment.

RISKS OF JUVEDERM® INJECTIONS

Every procedure involves a certain amount of risk and it is important that you understand these risks and the possible complications associated with them. In addition, every procedure has limitations. An individual's choice to undergo this procedure is based on the comparison of the risk to potential benefit. Although the majority of patients do not experience the following, you should discuss each of them with your physician to make sure you understand the risks, potential complications, limitations, and consequences of Juvederm injections. Additional information concerning Juvederm may be obtained from the package-insert sheets supplied by Allergan.

NORMAL OCCURRENCES DURING TISSUE FILLER INJECTIONS, INCLUDING JUVEDERM

Bleeding and Bruising - It is possible, though unusual, to have a bleeding episode from a Juvederm injection or local anesthesia used during the procedure. Bruising in soft tissues may occur. Should you develop post-injection bleeding, it may require emergency treatment or surgery. Aspirin, anti-inflammatory medications, platelet inhibitors, anticoagulants, Vitamin E, ginkgo biloba and other "herbs / homeopathic remedies" may contribute to a greater risk of a bleeding problem. Do not take any of these for seven days before or after Juvederm injections.

Swelling - Swelling (edema) is a normal occurrence following the injections. It decreases after a few days. If swelling is slow to resolve, medical treatment may be necessary.

Erythema (Skin Redness) - Erythema in the skin occurs after injections. It can be present for a few days after the procedure.

Needle Marks - Visible needle marks from the injections occur normally and resolve in a few days.
Acne-Like Skin Eruptions: Acneiform skin eruptions can occur following the injection of tissue fillers. This generally resolves within a few days.

Skin Lumpiness: Lumpiness can occur following the injection of Juvederm. This tends to smooth out over time. In some situations, it may be possible to feel the injected tissue filler material for long periods of time.

Visible Tissue Filler Material: It may be possible to see any type of tissue filler material that was injected in areas where the skin is thin.

Asymmetry: The human face is normally asymmetrical in its appearance and anatomy. It may not be possible to achieve or maintain exact symmetry with tissue filler injections. There can be a variation from one side to the other in terms of the response to Juvederm injection. This may require additional injections.

Pain: Discomfort associated with Juvederm injections is normal and usually of short duration.

Skin Sensitivity: Skin rash, itching, tenderness and swelling may occur following Juvederm injections. After treatment, you should minimize exposure of the treated area to excessive sun or UV lamp exposure and extreme cold weather until any initial swelling or redness has gone away. If you are considering laser treatment, chemical skin peeling or any other procedure based on a skin response after Juvederm treatment, or you have recently had such treatments and the skin has not healed completely, there is a possible risk of an inflammatory reaction at the filler site.

RISKS OF JUVEDERM INJECTIONS

Damage to Deeper Structures: Deeper structures such as nerves and blood vessels may be damaged during the course of injection. Injury to deeper structures may be temporary or permanent.

Infection: Although infection following injection of tissue fillers is unusual, bacterial, fungal, and viral infections can occur. Herpes simplex virus infections around the mouth can occur following a tissue filler treatment. This applies to both individuals with a past history of Herpes simplex virus infections and individuals with no known history of Herpes simplex virus infections in the mouth area. Specific medications must be prescribed and taken both prior to and following the treatment procedure in order to suppress an infection from this virus. Should any type of skin infection occur, additional treatment including antibiotics may be necessary.

Skin Necrosis: It is very unusual to experience death of skin and deeper soft tissues after Juvederm injections. Skin necrosis can produce unacceptable scarring. Should this complication occur, additional treatments, or surgery may be necessary.

Allergic Reactions and Hypersensitivity: As with all biologic products, allergic and systemic anaphylactic reactions may occur. Juvederm should not be used in patients with a history of multiple severe allergies, severe allergies manifested by a history of anaphylaxis, or allergies to gram-positive bacterial proteins. Allergic reactions may require additional treatment.

Scarring: Juvederm should not be used in patients with know susceptibility to keloid formation or hypertrophic scarring. The safety of patients has not been studied.

Granulomas: Painful masses in the skin and deeper tissues after a Juvederm injection are extremely rare. Should these occur, additional treatments including surgery may be necessary.

Skin Disorders: Juvederm should not be used in areas with active inflammation or infections (e.g., cysts, pimples, rashes or hives). In rare instances, granuloma or abscess formation, localized necrosis and urticaria have been reported.
Accidental Intra-Arterial Injection- It is extremely rare that during the course of injection, Juvederm could be accidentally injected into arterial structures and produce a blockage of blood flow. This may produce skin necrosis in facial structures or damage blood flow to the eye, resulting in loss of vision. The risk and consequences of accidental intravascular injection of Juvederm is unknown and not predictable.

Under / Over Correction- The injection of soft tissue fillers including Juvederm to correct wrinkles and soft tissue contour deficiencies may not achieve the desired outcome. The amount of correction may be inadequate or excessive. It may not be possible to control the process of injection of tissue fillers due to factors attributable to each patient's situation. If under correction occurs, you may be advised to consider additional injections of tissue filler materials.

Migration of Juvederm- Juvederm may migrate from its original injection site and produce visible fullness in adjacent tissue or other unintended effects.

Drug and Local Anesthetic Reactions- There is the possibility that a systemic reaction could occur from either the local anesthetic or epinephrine used for sensory nerve block anesthesia when tissue filler injections are performed. This would include the possibility of light-headedness, rapid heart beat (tachycardia), and fainting. Medical treatment of these conditions may be necessary.

ADDITIONAL ADVISORIES

Unsatisfactory Result- Juvederm injections alone may not produce an outcome that meets your expectations for improvement in wrinkles or soft tissue depressions. There is the possibility of a poor or inadequate response from Juvederm injection(s). Additional Juvederm injections may be necessary. Surgical procedures or other treatments may be recommended in additional to Juvederm treatments.

Unknown Risks- The long term effect of Juvederm beyond one year is unknown.

Combination of Procedures- In some situations, Botox® injections or other types of tissue filler materials may be used in addition to Juvederm in order to specifically treat areas of the face or to enhance the outcome from tissue filler therapy. The effect of other forms of external skin treatments (laser and other light therapies, microdermabrasion, dermabrasion, or chemical peels) on skin that has been treated with Juvederm is unknown.

Pregnancy and Nursing Mothers- Animal reproduction studies have not been performed to determine if Juvederm could produce fetal harm. It is not known if Juvederm or its breakdown products can be excreted in human milk. It is not recommended that pregnant women or nursing mothers receive Juvederm treatments.

Drug Interactions- It is not known if Juvederm reacts with other drugs within the body.

Long-Term Effects- Juvederm injections should not be considered as a permanent treatment for the correction of wrinkles and soft tissue depressions. Over time, the Juvederm material is slowly absorbed by the body and wrinkles or soft tissue depressions will reappear. Continuing Juvederm treatment (injections) is necessary in order to maintain the effect of Juvederm. Subsequent alterations in face and eyelid appearance may occur as the result of aging, weight loss or gain, sun exposure, or other circumstances not related to Juvederm injections. Future surgery or other treatments may be necessary. Juvederm injection does not arrest the aging process or produce permanent tightening of the skin or improvement in wrinkles.

ADDITIONAL TREATMENT NECESSARY

There are many variable conditions in addition to risk and potential complications that may influence the long-term result of Juvederm injections. Even though risks and complications occur infrequently, the risks cited are the ones that are particularly associated with Juvederm injections. Other complications and risks can occur but are even more uncommon. Should complications occur, additional surgery or other treatments may be necessary. The practice of medicine and surgery
is not an exact science. Although good results are expected, there is no guarantee or warranty expressed or implied, on the results that may be obtained.

PHOTOGRAPHIC RELEASE CONSENT:

I give permission to take photographs of my treatment areas for diagnostic purposes and to document for the medical record. I agree that these photographs are the property of the doctor's office, and I give my permission to use these photographs for teaching purposes, for use in scientific publications, books, journals, lectures, seminars and electronic media. It is understood that in any such publication I shall not be identified by name and that appropriate measures shall be made to protect my identity. I understand that I will not receive any compensation for use of my photos for scientific and teaching/educational purposes.

| I CONSENT TO THE TREATMENT OF HYALURONIC DERMAL FILLERS AND I HAVE READ THE ABOVE LISTED ITEMS. I AM SATISFIED WITH THE INFORMED CONSENT PROCESS |
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| Patient or Person Authorized to Sign for Patient | Date |
| Witness | Date |