



JUVÉDERM® Ultra and JUVÉDERM® Ultra Plus Dermal Filler Fact Sheet

About JUVÉDERM® Ultra and JUVÉDERM® Ultra Plus

- JUVÉDERM® Ultra and JUVÉDERM® Ultra Plus, approved by the FDA in June 2006, are hyaluronic acid (HA) dermal fillers and the only HA dermal fillers FDA-approved to last one year from initial treatment (includes up to two touch ups), providing a smooth, long-lasting correction of moderate to severe facial wrinkles and folds. JUVÉDERM® dermal filler is the latest advance in HA dermal filler options from Allergan, Inc.
- JUVÉDERM® is the first and only smooth consistency gel formulation, and the only HA dermal filler developed using the proprietary HYLACROSS™ technology, a technologically advanced manufacturing process that results in a malleable, smooth gel. JUVÉDERM® flows easily into the skin and creates a smooth, natural look and feel.
 - All other HA dermal fillers currently on the market have a granular consistency gel. These granules can be seen under 2.4X magnification as opposed to the smooth consistency gel of JUVÉDERM® dermal filler.
- In addition, JUVÉDERM® contains a high concentration of non-animal, cross-linked HA. This unique attribute provides optimal results with a single treatment in the majority of patients and delivers sustained results for up to one year.
- JUVÉDERM® dermal filler is biodegradable and has demonstrated its safety and effectiveness in patients of color. Studies with JUVÉDERM® demonstrated no increased risk of hyperpigmentation or hypertrophic scarring in patients of color.
- JUVÉDERM® Ultra and JUVÉDERM® Ultra Plus can be administered in a smooth-flowing injection, providing physicians with a high level of control for individualized, tailored wrinkle treatment and for achieving a smooth and natural look.
- The U.S. Food and Drug Administration (FDA) approved the following formulations of JUVÉDERM®, providing physicians with the flexibility to tailor each treatment to the particular needs of the patient. Product formulations include:
 - JUVÉDERM® Ultra, a highly cross-linked formulation for more versatility in contouring and volumizing of facial wrinkles and folds
 - JUVÉDERM® Ultra Plus, a more highly cross-linked robust formulation for volumizing and correction of deeper folds and wrinkles

How JUVÉDERM® Works

- The key component in JUVÉDERM®, hyaluronic acid, is a naturally occurring, biodegradable complex sugar found in the human body and in all mammals. Among other things, hyaluronic acid hydrates the skin and adds volume, contributing to the overall appearance of the skin.

Please see page 3 for important JUVÉDERM® safety information



- The ability of cells to produce hyaluronic acid diminishes with age, often resulting in the formation of facial wrinkles and folds as the skin loses volume. JUVÉDERM® is administered in small doses by intradermal injection directly into moderate to severe facial wrinkles and folds, such as nasolabial folds (the “parentheses” along the side of the nose and mouth), temporarily filling and augmenting the treated area.
- JUVÉDERM® Ultra and JUVÉDERM® Ultra Plus should only be administered by a trained and qualified health care provider. Further product and patient risk information is available by visiting www.Juvederm.com.

JUVÉDERM® and the ALLERGAN® Facial Aesthetics Portfolio of Products

- Physicians often will use multiple products, depending upon the patients’ needs.
- With nearly 60 years of experience providing high-quality, science-based products, Allergan, Inc.’s Facial Aesthetics Portfolio of Products includes: BOTOX® Cosmetic (onabotulinumtoxinA); JUVÉDERM® hyaluronic acid dermal filler; and physician-dispensed products such as PREVAGE® MD anti-aging treatment, which contains idebenone, the most powerful antioxidant available in a skin care product today; VIVITÉ®, the only luxury physician-dispensed skin care regimen that works through *GLX Technology™* to help renew skin and enhance the skin’s natural production of hyaluronic acid; and most recently LATISSE® (bimatoprost ophthalmic solution) 0.03%, the first and only prescription treatment for hypotrichosis used to grow eyelashes, making them longer, thicker and darker. Eyelash hypotrichosis is another name for having inadequate or not enough eyelashes. **(Please see page 4 for important LATISSE® safety information)**

BOTOX® Cosmetic (onabotulinumtoxinA)

- BOTOX® Cosmetic, approved by the FDA in 2002, is a prescription medicine that is injected into muscles and used to improve the look of moderate to severe frown lines between the eyebrows (glabellar lines) in adults younger than 65 years of age for a short period of time (temporary). BOTOX® Cosmetic works differently – by temporarily reducing the muscle activity of the dominant frown muscles between the eyebrows (the glabellar area). This allows the two vertical lines between the brows, often referred to as the “11,” to temporarily diminish in appearance.

IMPORTANT BOTOX® COSMETIC SAFETY INFORMATION

BOTOX® Cosmetic may cause serious side effects that can be life threatening. Call your doctor or get medical help right away if you have any of these problems after treatment with BOTOX® Cosmetic:

- **Problems swallowing, speaking, or breathing.** These problems can happen hours to weeks after an injection of BOTOX® Cosmetic usually because the muscles that you use to breathe and swallow can become weak after the injection. Death can happen as a complication if you have severe problems with swallowing or breathing after treatment with BOTOX® Cosmetic.
- Swallowing problems may last for several months. People who already have swallowing or breathing problems before receiving BOTOX® Cosmetic have the highest risk of getting these problems.

- **Spread of toxin effects.** In some cases, the effect of botulinum toxin may affect areas of the body away from the injection site and cause symptoms of a serious condition called botulism. The symptoms of botulism include: loss of strength and muscle weakness all over the body, double vision, blurred vision and drooping eyelids, hoarseness or change or loss of voice (dysphonia), trouble saying words clearly (dysarthria), loss of bladder control, trouble breathing, trouble swallowing.

These symptoms can happen hours to weeks after you receive an injection of BOTOX® Cosmetic.

These problems could make it unsafe for you to drive a car or do other dangerous activities. See "What should I avoid while receiving BOTOX® Cosmetic" in Medication Guide.

Please see next page for additional Important Safety Information.



IMPORTANT BOTOX® COSMETIC SAFETY INFORMATION (continued)

There has not been a confirmed serious case of spread of toxin effect away from the injection site when BOTOX® has been used at the recommended dose to treat severe underarm sweating, blepharospasm, or strabismus, or when BOTOX® Cosmetic has been used at the recommended dose to treat frown lines.

Tell your doctor about all your medical conditions, including if you have: a disease that affects your muscles and nerves (such as amyotrophic lateral sclerosis [ALS or Lou Gehrig's disease], myasthenia gravis or Lambert-Eaton syndrome).

Tell your doctor about all the medicines you take, including prescription and nonprescription medicines, vitamins and herbal products.

BOTOX® Cosmetic (onabotulinumtoxinA) can cause serious side effects. 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, swelling of your eyelids, and dry eyes. Symptoms of an allergic reaction to BOTOX® Cosmetic may include: itching, rash, red itchy welts, wheezing, asthma symptoms, or dizziness or feeling faint. Tell your doctor or get medical help right away if you are wheezing or have asthma symptoms, or if you become dizzy or faint.

Tell your doctor if you have any side effect that bothers you or that does not go away.

These are not all the possible side effects of BOTOX® Cosmetic. For more information, ask your doctor or pharmacist.

For additional information refer to Medication Guide. This Medication Guide summarizes the most important information about BOTOX® Cosmetic. If you would like more information, talk with your doctor.

Please see the accompanying full Product Information, including Medication Guide, for BOTOX® Cosmetic.

A Brief Description of Relevant Indications for Use, Contraindications, Warnings, Precautions, and Adverse Events for JUVÉDERM® Injectable Gel

Indication: In the United States, JUVÉDERM® injectable gel is indicated for correction of moderate to severe facial wrinkles and folds (such as nasolabial folds).

Contraindications: JUVÉDERM® injectable gel should not be used in patients who have severe allergies marked by a history of anaphylaxis or history or presence of multiple severe allergies. JUVÉDERM® injectable gel should not be used in patients with a history of allergies to Gram-positive bacterial proteins.

Warnings: JUVÉDERM® injectable gel should not be injected into blood vessels. If there is an active inflammatory process or infection at specific injection sites, treatment should be deferred until the underlying process is controlled.

Precautions: The safety of JUVÉDERM® injectable gel for use during pregnancy, in breastfeeding females, or in patients under 18 years has not been established. The safety and effectiveness of JUVÉDERM® injectable gel for the treatment of areas other than facial wrinkles and folds (such as lips) have not been established in controlled clinical studies. Patients who are using substances that can prolong bleeding, such as aspirin or ibuprofen, as with any injection, may experience increased bruising or bleeding at injection site. Patients should inform their physician before treatment if they are using these types of substances. As with all skin-injection procedures, there is a risk of infection. JUVÉDERM® injectable gel should be used with caution in patients on immunosuppressive therapy, or therapy used to decrease the body's immune response, as there may be an increased risk of infection. The safety of JUVÉDERM® injectable gel in patients with a history of excessive scarring (e.g., hypertrophic scarring and keloid formations) and pigmentation disorders has not been studied. If laser treatment, chemical peel, or any other procedure based on active dermal response is considered after treatment with JUVÉDERM® injectable gel, or if JUVÉDERM® injectable gel is administered before the skin has healed completely after such a procedure, there is a possible risk of an inflammatory reaction at the treatment site.

Adverse events: The most commonly reported side effects are temporary injection-site redness, swelling, pain/tenderness, firmness, lumps/bumps, and bruising. Most side effects are mild or moderate in nature, and their duration is short lasting (7 days or less).

Important: For full safety information, please visit www.JUVEDERM.com or call Allergan Product Support at 1-877-345-5372.

CAUTION: This device is restricted to sale by or on the order of a physician.

Important Latisse® Information

Latisse® solution is a prescription treatment for hypotrichosis used to grow eyelashes, making them longer, thicker and darker.

Eyelash hypotrichosis is another name for having inadequate or not enough eyelashes.

Important Latisse® Safety Information

If you are using prescription products for lowering eye pressure or have a history of eye pressure problems, only use Latisse® under close doctor supervision.

May cause eyelid skin darkening which may be reversible, and there is a potential for increased brown iris pigmentation which is likely to be permanent.

For further information please contact Allergan, Inc.:

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There is a potential for hair growth to occur in areas where Latisse® solution comes in repeated contact with skin surfaces.

If you develop or experience any eye problems or have eye surgery, consult your doctor immediately about continued use of Latisse®.

The most common side effects after using Latisse® solution are an itching sensation in the eyes and/or eye redness.

For more information on Latisse®, please refer to accompanying prescribing information or visit www.LATISSE.com and www.allergan.com.