

ALLERGAN® FACIAL AESTHETICS PORTFOLIO OF PRODUCTS

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 support the skin's natural production of hyaluronic acid; M.D. FORTE®, a comprehensive, personalized glycolic acid skin care system; the CLINIQUE MEDICAL skin care line, developed in collaboration with Clinique, the #1 prestige cosmetics brand in the United States, to help improve skin's receptivity to select procedures – laser skin resurfacing such as Intense Pulsed Light and fractionated laser treatments, chemical peels such as 30% TCA peels, and microdermabrasion – and help support the healing process by managing post-procedure visible excess redness and inflammation and encourage even skin tone; and, most recently, LATISSE® (bimatoprost ophthalmic solution) 0.03%, the first and only prescription treatment approved by the FDA for hypotrichosis used to grow eyelashes, making them longer, thicker and darker. Eyelash hypotrichosis is another name for having inadequate or not enough eyelashes.

Highlights of the product line include:

BOTOX® Cosmetic (onabotulinumtoxinA)

BOTOX® Cosmetic 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 temporarily reduces the muscle activity that causes frown lines to form by blocking nerve impulses that trigger wrinkle-causing muscle contractions, creating an improved appearance between the brows. BOTOX® Cosmetic has been the number one physician-administered, non-surgical cosmetic procedure in the U.S. since its U.S. Food and Drug Administration (FDA) approval in 2002, according to the American Society for Plastic Surgeons (ASPS)¹.

JUVÉDERM® Dermal Filler Family of Products

JUVÉDERM® Ultra and JUVÉDERM® Ultra Plus, approved by the FDA in June 2006, are the 'next generation' of hyaluronic acid (HA) dermal fillers and are the only hyaluronic acid dermal fillers approved to last up to one year, providing a smooth, long-lasting correction of moderate to severe facial wrinkles and folds. JUVÉDERM® contains a high concentration of non-animal, cross-linked hyaluronic acid, and has demonstrated its safety and effectiveness in patients of color. JUVÉDERM® is developed using the proprietary HYLACROSS™ technology, a technologically advanced manufacturing process that results in a malleable, smooth gel that flows easily into the skin and creates a smooth, natural look and feel. All other HA 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 the JUVÉDERM® family of products.

The 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 and JUVÉDERM® Ultra Plus, a more highly cross-linked robust formulation for volumizing and correction of deeper folds and wrinkles.

Please see page 3 for important JUVÉDERM® and LATISSE® safety information.

IMPORTANT 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.

Please see next page for additional Important Safety Information for BOTOX® Cosmetic.

PREVAGE® MD

PREVAGE® MD is an innovative anti-aging treatment representing the next generation of topical antioxidants. PREVAGE® MD contains the highest concentration (one percent) of idebenone, the most potent antioxidant currently available in a skin care product, to help correct the appearance of fine lines and wrinkles and protect against environmental factors including sun damage, air pollution and cigarette smoke. PREVAGE® MD anti-aging treatment is available exclusively through dermatologists' and plastic surgeons' offices without a prescription.

M.D. FORTÉ®

M.D. FORTÉ® is a comprehensive, personalized glycolic acid skin care system dispensed by dermatologists and plastic surgeons. The M.D. FORTÉ® system offers a regimen to cleanse, renew, hydrate and protect skin. M.D. FORTÉ® products provide effective glycolic acid concentrations to help improve the look and feel of skin.

VIVITÉ®

VIVITÉ® is an advanced skin care line that helps revitalize and shield skin to help reduce the signs of aging. VIVITÉ® is a scientific advancement in skin care and the only skin care line to include proprietary *GLX Technology™*, which creates a highly specialized blend of hydrating glycolic acid and powerful, natural antioxidants.

CLINIQUE MEDICAL

Developed in collaboration with Clinique, the #1 prestige cosmetics brand in the United States, the CLINIQUE MEDICAL skin care line is available only in skin care physicians' offices. The products in the CLINIQUE MEDICAL skin care line work together to help improve skin's receptivity to select procedures – laser skin resurfacing such as Intense Pulsed Light and fractionated laser treatments, chemical peels such as 30% TCA peels, and microdermabrasion – and aid in the recovery process by helping to manage post-procedure visible excess redness and inflammation and encourage even skin tone. All products are Allergy Tested, 100% Fragrance Free and contain no preservatives. The CLINIQUE MEDICAL products can be purchased individually or in the CLINIQUE MEDICAL Optimizing Regimen kit which consists of a Probiotic Cleanser, Skin Conditioning Treatment, Recovery Week Complex, Optimizing Treatment Cream and Daily SPF 38.

LATISSE® (bimatoprost ophthalmic solution) 0.03%

LATISSE® (bimatoprost ophthalmic solution) 0.03% is the first and only prescription treatment approved by the FDA for hypotrichosis used to grow eyelashes, making them longer, thicker and darker. Eyelash hypotrichosis is another name for having inadequate or not enough eyelashes.

LATISSE® is a once-daily treatment applied to the base of the upper eyelashes with a sterile, single-use-per-eye disposable applicator. The onset of effect with LATISSE® solution is gradual. In the clinical trial, the majority of patients saw a significant improvement by 2 months. To maintain effect, continued treatment with LATISSE® is required. If treatment is discontinued, eyelashes will gradually return to where they were prior to treatment over a period of weeks to months (average eyelash hair cycle). **Please see page 3 for important LATISSE® safety information.**

IMPORTANT BOTOX® COSMETIC (ONABOTULINUMTOXINA) SAFETY INFORMATION (continued)

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.

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.

Please see next page for additional Important Safety Information for BOTOX® Cosmetic.

**IMPORTANT BOTOX® COSMETIC
(ONABOTULINUMTOXINA) SAFETY INFORMATION**
(continued)

BOTOX® Cosmetic 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 (eg, 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.

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®**.

Please see next page for additional Important Safety Information for LATISSE®.

Important Latisse® Safety Information (continued)

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.

For additional information, please refer to:

BOTOX® Cosmetic: www.BOTOXCosmetic.com

JUVÉDERM®: www.JUVÉDERM.com

PREVAGE® MD: www.PrevageMD.com

M.D. FORTÉ®: www.MDForte.com

VIVITÉ®: www.VIVITE.com

LATISSE®: www.LATISSE.com

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ⁱ American Society of Plastic Surgeons; "Top Five Minimally-Invasive Cosmetic Procedures" 2002, 2003, 2004, 2005, 2006, 2007, 2008. Available at <http://www.plasticsurgery.org/Media/Statistics.html>