



JUVÉDERM™ injectable gel Important Treatment Considerations for Patients

In the US, JUVÉDERM™ injectable gel is indicated for injection into the mid-to-deep dermis for correction of moderate to severe facial wrinkles and folds (such as nasolabial folds) and is generally well tolerated. The most commonly reported side effects are temporary injection-site redness, swelling, pain/tenderness, firmness, lumps/bumps, and bruising. Exposure of the treated area to excessive sun and extreme cold weather should be minimized until any initial swelling and redness have resolved. If laser treatment, chemical peel, or any other procedure based on active dermal response is considered after treatment with JUVÉDERM™ injectable gel, there is a possible risk of an inflammatory reaction at the treatment site.

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. You should inform your physician before treatment if you 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.

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. 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.

www.juvederm.com

1-877-345-5372

BOTOX® Cosmetic (Botulinum Toxin Type A)

BOTOX® Cosmetic is approved for the temporary treatment of moderate to severe frown lines between the brows in people ages 18 to 65.

Important Safety Information: Serious heart problems and serious allergic reactions have been reported rarely. If you think you're having an allergic reaction or other unusual symptoms, such as difficulty swallowing, speaking, or breathing, call your doctor immediately. The most common side effects following injection include temporary eyelid droop and nausea. Localized pain, infection, inflammation, tenderness, swelling, redness, and/or bleeding/bruising may be associated with the injection. Patients with certain neuromuscular disorders such as ALS, myasthenia gravis, or Lambert-Eaton syndrome may be at increased risk of serious side effects.

Full prescribing information has been provided to your doctor.

www.BotoxCosmetic.com

1-800-BOTOXMD