

A LIMITED-TIME OFFER FOR YOUR PATIENTS

2 DIFFERENT TREATMENTS

#1 CHOSEN JUVÉDERM®
FILLER COLLECTION¹ COLLECTION OF FILLERS

#1 SELLING BOTOX®
PRODUCT OF ITS KIND² COSMETIC
onabotulinumtoxinA
injection

2X
THE POINTS
ON JUVÉDERM®

- Your patients can now earn up to 1800 AllēSM points (\$180 in savings)*
- The more points they earn now, the more they will save on future treatments

*Patients can earn up to 1600 points on JUVÉDERM® and 200 points on BOTOX® Cosmetic treatments, for a total of 1800 points.

Note: The safety and efficacy of these products for combined use have not been studied.



Actual patient treated with JUVÉDERM® VOLUMA™ XC in the cheeks and chin. Results may vary.

Double Points on JUVÉDERM® Promotion Terms and Conditions

- Members who purchase treatments with both BOTOX® Cosmetic (onabotulinumtoxinA) and at least 1 of any JUVÉDERM® Collection of Fillers products in the same visit will earn double Allē Loyalty Program points on a maximum of 4 qualifying syringes of any JUVÉDERM® product (1600 maximum JUVÉDERM® points)
- Members will earn standard Allē points on all other eligible purchases
- A healthcare provider will determine if a member is an appropriate candidate for BOTOX® Cosmetic and JUVÉDERM® Collection of Fillers treatments
- The safety and efficacy of these products for combined use have not been studied
- BOTOX® Cosmetic and JUVÉDERM® treatments must occur in the same visit and be part of the same Allē transaction to qualify for Double Points offer
- Treatments must occur between February 9, 2021, and May 9, 2021, to be eligible for Double Points offer
- Standard Allē Loyalty Program Terms and Conditions apply, including Earnings Caps
- Double Points offer cannot be applied to past transactions
- Please allow up to 48 hours for points to deposit into Allē members' accounts
- Double Points offer can be combined with other Allē, BOTOX® Cosmetic, and JUVÉDERM® offers; Allē, BOTOX® Cosmetic, and JUVÉDERM® gift cards; and Allē points

BOTOX® Cosmetic (onabotulinumtoxinA) Important Information

Indications

BOTOX® Cosmetic (onabotulinumtoxinA) is indicated in adult patients for the temporary improvement in the appearance of:

- Moderate to severe glabellar lines associated with corrugator and/or procerus muscle activity
- Moderate to severe lateral canthal lines associated with orbicularis oculi activity
- Moderate to severe forehead lines associated with frontalis activity

- If you have questions, please contact Allē Customer Support at 1-888-912-1572, Monday to Friday, 8 AM to 6 PM CT
- Allergan Aesthetics, an AbbVie company, reserves the right to alter or cancel this offer at any time

IMPORTANT SAFETY INFORMATION, INCLUDING BOXED WARNING

WARNING: DISTANT SPREAD OF TOXIN EFFECT

Postmarketing reports indicate that the effects of BOTOX® Cosmetic and all botulinum toxin products may spread from the area of injection to produce symptoms consistent with botulinum toxin effects. These may include asthenia, generalized muscle weakness, diplopia, ptosis, dysphagia, dysphonia, dysarthria, urinary incontinence, and breathing difficulties. These symptoms have been reported hours to weeks after injection. Swallowing and breathing difficulties can be life threatening and there have been reports of death. The risk of symptoms is probably greatest in children treated for spasticity, but symptoms can also occur in adults treated for spasticity and other conditions, particularly in those patients who have an underlying condition that would predispose them to these symptoms. In unapproved uses and approved indications, cases of spread of effect have been reported at doses comparable to those used to treat cervical dystonia and spasticity and at lower doses.

Please see additional Important Safety Information about BOTOX® Cosmetic on following pages.

Please see Indications and Important Safety Information for the JUVÉDERM® Collection of Fillers on following pages.

One syringe
of JUVÉDERM®

~~200~~
400

Allē points

How it works for your patients:

- 1) Receive BOTOX® Cosmetic (onabotulinumtoxinA) and JUVÉDERM® in the same visit.
- 2) Earn **double points** on JUVÉDERM® treatments*,†

*Double points earned in this offer apply only to JUVÉDERM® products. Eligible patients must be enrolled in the Allē program to earn points.

†Double points can be earned per syringe on a maximum of 4 syringes of JUVÉDERM® products. If additional syringes are needed, patients earn standard Allē points.

GET STARTED TODAY



Offer valid from 2/9/21 through 5/9/21.

BOTOX® Cosmetic (onabotulinumtoxinA)
IMPORTANT SAFETY INFORMATION (continued)

CONTRAINDICATIONS

BOTOX® Cosmetic is contraindicated in the presence of infection at the proposed injection site(s) and in individuals with known hypersensitivity to any botulinum toxin preparation or to any of the components in the formulation.

WARNINGS AND PRECAUTIONS

Lack of Interchangeability Between Botulinum Toxin Products
The potency units of BOTOX® Cosmetic are specific to the preparation and assay method utilized. They are not interchangeable with other preparations of botulinum toxin products and, therefore, units of biological activity of BOTOX® Cosmetic cannot be compared to nor converted into units of any other botulinum toxin products assessed with any other specific assay method.

Spread of Toxin Effect

Please refer to Boxed Warning for Distant Spread of Toxin Effect.
No definitive serious adverse event reports of distant spread of toxin effect associated with dermatologic use of BOTOX® Cosmetic at the labeled dose of 20 Units (for glabellar lines), 24 Units (for lateral canthal lines), 40 Units (for forehead lines with glabellar lines), 44 Units (for simultaneous treatment of lateral canthal lines and glabellar lines), and 64 Units (for simultaneous treatment of lateral canthal lines, glabellar lines, and forehead lines) have been

reported. Patients or caregivers should be advised to seek immediate medical care if swallowing, speech, or respiratory disorders occur.

Serious Adverse Reactions With Unapproved Use

Serious adverse reactions, including excessive weakness, dysphagia, and aspiration pneumonia, with some adverse reactions associated with fatal outcomes, have been reported in patients who received BOTOX® injections for unapproved uses. In these cases, the adverse reactions were not necessarily related to distant spread of toxin, but may have resulted from the administration of BOTOX® to the site of injection and/or adjacent structures. In several of the cases, patients had pre-existing dysphagia or other significant disabilities. There is insufficient information to identify factors associated with an increased risk for adverse reactions associated with the unapproved uses of BOTOX®. The safety and effectiveness of BOTOX® for unapproved uses have not been established.

Hypersensitivity Reactions

Serious and/or immediate hypersensitivity reactions have been reported. These reactions include anaphylaxis, serum sickness, urticaria, soft-tissue edema, and dyspnea. If such reactions occur, further injection of BOTOX® Cosmetic should be discontinued and appropriate medical therapy immediately instituted. One fatal case of anaphylaxis has been reported in which lidocaine was used as the diluent and, consequently, the causal agent cannot be reliably determined.

Please see additional Important Safety Information about BOTOX® Cosmetic on following pages.

Please see Indications and Important Safety Information for the JUVÉDERM® Collection of Fillers on following pages.

BOTOX® Cosmetic (onabotulinumtoxinA)

IMPORTANT SAFETY INFORMATION (continued)

Cardiovascular System

There have been reports following administration of BOTOX® of adverse events involving the cardiovascular system, including arrhythmia and myocardial infarction, some with fatal outcomes. Some of these patients had risk factors including pre-existing cardiovascular disease. Use caution when administering to patients with pre-existing cardiovascular disease.

Increased Risk of Clinically Significant Effects With Pre-existing Neuromuscular Disorders

Individuals with peripheral motor neuropathic diseases, amyotrophic lateral sclerosis, or neuromuscular junction disorders (eg, myasthenia gravis or Lambert-Eaton syndrome) should be monitored when given botulinum toxin. Patients with neuromuscular disorders may be at increased risk of clinically significant effects including generalized muscle weakness, diplopia, ptosis, dysphonia, dysarthria, severe dysphagia, and respiratory compromise from onabotulinumtoxinA (see *Warnings and Precautions*).

Dysphagia and Breathing Difficulties

Treatment with BOTOX® and other botulinum toxin products can result in swallowing or breathing difficulties. Patients with pre-existing swallowing or breathing difficulties may be more susceptible to these complications. In most cases, this is a consequence of weakening of muscles in the area of injection that are involved in breathing or oropharyngeal muscles that control swallowing or breathing (see *Boxed Warning*).

Pre-existing Conditions at the Injection Site

Caution should be used when BOTOX® Cosmetic treatment is used in the presence of inflammation at the proposed injection site(s) or when excessive weakness or atrophy is present in the target muscle(s).

Dry Eye in Patients Treated With BOTOX® Cosmetic

There have been reports of dry eye associated with BOTOX® Cosmetic injection in or near the orbicularis oculi muscle. If symptoms of dry eye (eg, eye irritation, photophobia, or visual changes) persist, consider referring patients to an ophthalmologist.

Human Albumin and Transmission of Viral Diseases

This product contains albumin, a derivative of human blood. Based on effective donor screening and product manufacturing processes, it carries an extremely remote risk for transmission of viral diseases and variant Creutzfeldt-Jakob disease (vCJD). There is a theoretical risk for transmission of Creutzfeldt-Jakob disease (CJD), but if that risk actually exists, the risk of transmission would also be considered extremely remote. No cases of transmission of viral diseases, CJD or vCJD have ever been identified for licensed albumin or albumin contained in other licensed products.

ADVERSE REACTIONS

The most frequently reported adverse reactions following injection of BOTOX® Cosmetic for glabellar lines were eyelid ptosis (3%), facial pain (1%), facial paresis (1%), and muscular weakness (1%).

The most frequently reported adverse reaction following injection of BOTOX® Cosmetic for lateral canthal lines was eyelid edema (1%).

The most frequently reported adverse reactions following injection of BOTOX® Cosmetic for forehead lines with glabellar lines were headache (9%), brow ptosis (2%), and eyelid ptosis (2%).

DRUG INTERACTIONS

Co-administration of BOTOX® Cosmetic and aminoglycosides or other agents interfering with neuromuscular transmission (eg, curare-like compounds) should only be performed with caution as the effect of the toxin may be potentiated. Use of anticholinergic drugs after administration of BOTOX® Cosmetic may potentiate systemic anticholinergic effects.

The effect of administering different botulinum neurotoxin products at the same time or within several months of each other is unknown. Excessive neuromuscular weakness may be exacerbated by administration of another botulinum toxin prior to the resolution of the effects of a previously administered botulinum toxin.

Excessive weakness may also be exaggerated by administration of a muscle relaxant before or after administration of BOTOX® Cosmetic.

USE IN SPECIFIC POPULATIONS

There are no studies or adequate data from postmarketing surveillance on the developmental risk associated with use of BOTOX® Cosmetic in pregnant women. There are no data on the presence of BOTOX® Cosmetic in human or animal milk, the effects on the breastfed child, or the effects on milk production.

Please see BOTOX® Cosmetic full Prescribing Information including Boxed Warning and Medication Guide.

JUVÉDERM® Collection of Fillers Important Information

INDICATIONS

JUVÉDERM® VOLUMA™ XC is indicated for deep (subcutaneous and/or supraperiosteal) injection for cheek augmentation to correct age-related volume deficit in the mid-face and for augmentation of the chin region to improve the chin profile in adults over the age of 21.

JUVÉDERM® VOLLURE™ XC 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) in adults over the age of 21.

JUVÉDERM® Ultra Plus XC and JUVÉDERM® Ultra XC injectable gels are indicated for injection into the mid-to-deep dermis for correction of moderate to severe facial wrinkles and folds (such as nasolabial folds).

JUVÉDERM® VOLBELLA™ XC injectable gel is indicated for injection into the lips for lip augmentation and for correction of perioral rhytids in adults over the age of 21.

JUVÉDERM® Ultra XC injectable gel is indicated for injection into the lips and perioral area for lip augmentation in adults over the age of 21.

IMPORTANT SAFETY INFORMATION

CONTRAINDICATIONS

These products should not be used in patients who have severe allergies, marked by a history of anaphylaxis or history or presence of multiple severe allergies, and should not be used in patients with a history of allergies to Gram-positive bacterial proteins or lidocaine contained in these products.

WARNINGS

- Do not inject into blood vessels. Introduction of these products into the vasculature may lead to embolization, occlusion of the vessels, ischemia, or infarction. Take extra care when injecting soft-tissue fillers; for example, inject the product slowly and apply the least amount of pressure necessary. Rare, but serious, adverse events associated with the intravascular injection of soft-tissue fillers in the face have been reported and include temporary or permanent vision impairment, blindness, cerebral ischemia or cerebral hemorrhage leading to stroke, skin necrosis, and damage to underlying facial structures. Immediately stop the injection if a patient exhibits any of the following symptoms: changes in vision, signs of a stroke, blanching of the skin, unusual pain during or shortly after the procedure. Patients should receive prompt medical attention and, possibly, evaluation by an appropriate healthcare professional specialist should an intravascular injection occur
- Product use at specific sites in which an active inflammatory process (skin eruptions such as cysts, pimples, rashes, or hives) or infection is present should be deferred until the underlying process has been controlled

PRECAUTIONS

- To minimize the risk of potential complications, these products should only be used by healthcare professionals who have appropriate training, experience, and knowledge of facial anatomy and product use in indicated areas
- The potential risks of soft-tissue injections should be discussed with patients prior to treatment to ensure they are aware of signs and symptoms of complications
- The safety and effectiveness for the treatment of anatomic regions other than the mid-face, chin, and prejowl sulcus regions with JUVÉDERM® VOLUMA™ XC; facial wrinkles and folds with JUVÉDERM® VOLLURE™ XC, JUVÉDERM® Ultra Plus XC, and JUVÉDERM® Ultra XC; and the lips and perioral area with JUVÉDERM® VOLBELLA™ XC and JUVÉDERM® Ultra XC have not been established in controlled clinical studies
- The safety for use of these products during pregnancy, in breastfeeding females, and in patients with known susceptibility to keloid formation, hypertrophic scarring, and pigmentation disorders has not been studied
- The safety for use of JUVÉDERM® VOLUMA™ XC has been established in patients between 35 and 65 years of age in cheek augmentation and for patients between 22 and 80 years of age for chin augmentation
- The safety for use of JUVÉDERM® Ultra Plus XC and JUVÉDERM® Ultra XC in patients under 18 years, and JUVÉDERM® VOLLURE™ XC and JUVÉDERM® VOLBELLA™ XC in patients under 22 years, has not been established
- As with all transcutaneous procedures, dermal filler implantation carries a risk of infection
- Use dermal fillers with caution in patients on immunosuppressive therapy

Please see additional Important Safety Information for the JUVÉDERM® Collection of Fillers on following page.

JUVÉDERM® Collection of Fillers IMPORTANT SAFETY INFORMATION (continued)

PRECAUTIONS (continued)

- Patients who are using products that can prolong bleeding (such as aspirin, nonsteroidal anti-inflammatory drugs, and warfarin) may experience increased bruising or bleeding at treatment sites
- Patients who experience skin injury near the site of implantation may be at a higher risk for adverse events
- The safety for use of JUVÉDERM® VOLUMA™ XC injectable gel in patients with very thin skin in the mid-face has not been established
- The safety of JUVÉDERM® VOLUMA™ XC with cannula for cheek augmentation has not been established in patients with Fitzpatrick Skin Types V and VI
- JUVÉDERM® VOLUMA™ XC was not evaluated in subjects with significant skin laxity of the chin, neck, or jaw in the chin augmentation study
- The effect of JUVÉDERM® VOLUMA™ XC injection into the chin on facial hair growth has not been studied
- Patients may experience late onset nodules with use of dermal fillers including JUVÉDERM® VOLUMA™ XC
- Patients may experience late onset adverse events with use of dermal fillers

ADVERSE EVENTS

The most commonly reported side effects for JUVÉDERM® injectable gels were redness, swelling, pain, tenderness, firmness, lumps/bumps, bruising, discoloration, and itching. For JUVÉDERM® VOLBELLA™ XC, dryness was also reported. The majority were mild or moderate in severity. For JUVÉDERM® VOLUMA™ XC, most resolved within 2 to 4 weeks. For JUVÉDERM® VOLLURE™ XC, JUVÉDERM® Ultra Plus XC, or JUVÉDERM® Ultra XC, most resolved within 14 days; and for JUVÉDERM® VOLBELLA™ XC, most resolved within 30 days.

To report an adverse reaction with any product in the JUVÉDERM® Collection, please call Allergan at 1-800-433-8871. Please visit JuvedermDFU.com for more information.

Products in the JUVÉDERM® Collection are available only by a licensed physician or properly licensed practitioner.

References: **1.** Data on file, Allergan, November 2020; Dermal Filler Monthly Tracker. **2.** Data on file, Allergan, December 2020; Neurotoxin Aesthetic Monthly Tracker.