Allē

Make the most of Allē Insights

Insights is Allē's reporting engine, built to highlight the information that matters to your practice. Personalize your communications, recommend treatments with confidence, and find opportunities for improvement using the information logged at checkout.

Stay informed with:

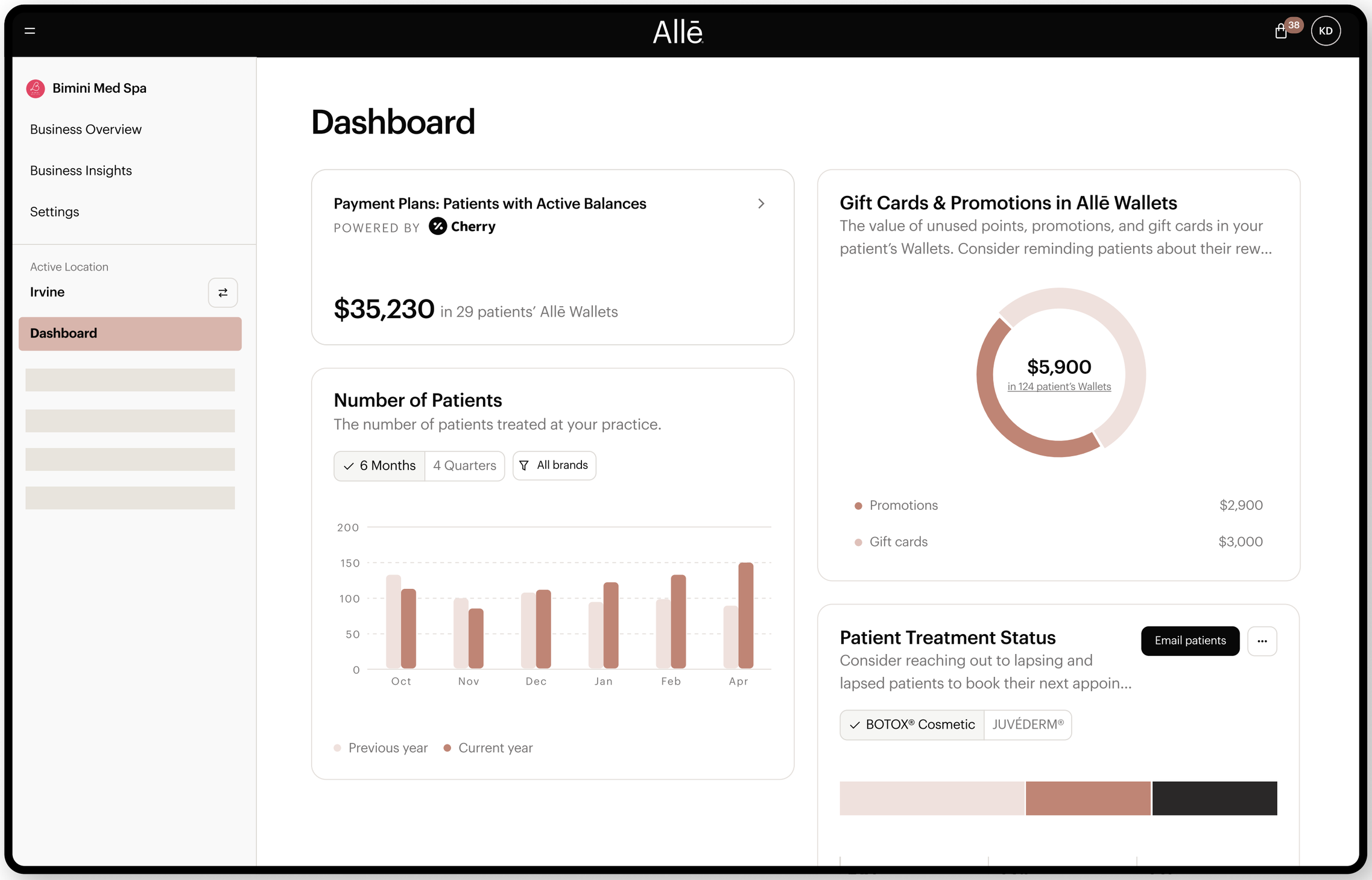
- Interactive widgets displayed on your dashboard
- Patient 360, viewable directly in Allē
- Downloadable Insights reports

Not an actual provider.

Pro tip: Train your staff to use Allē at checkout.

The information revealed by Insights is only as good as the visit details your team logs in Allē—be sure to log all on-label treatments so that they appear in your reports.

Information at a glance with interactive widgets



The Insights widgets on your dashboard show high-level patient trends like treatment statuses, rewards, and brands.

- Use the filters to narrow down the results and see specific treatment trends, which can allow you to identify ways to help patients meet their aesthetic goals.

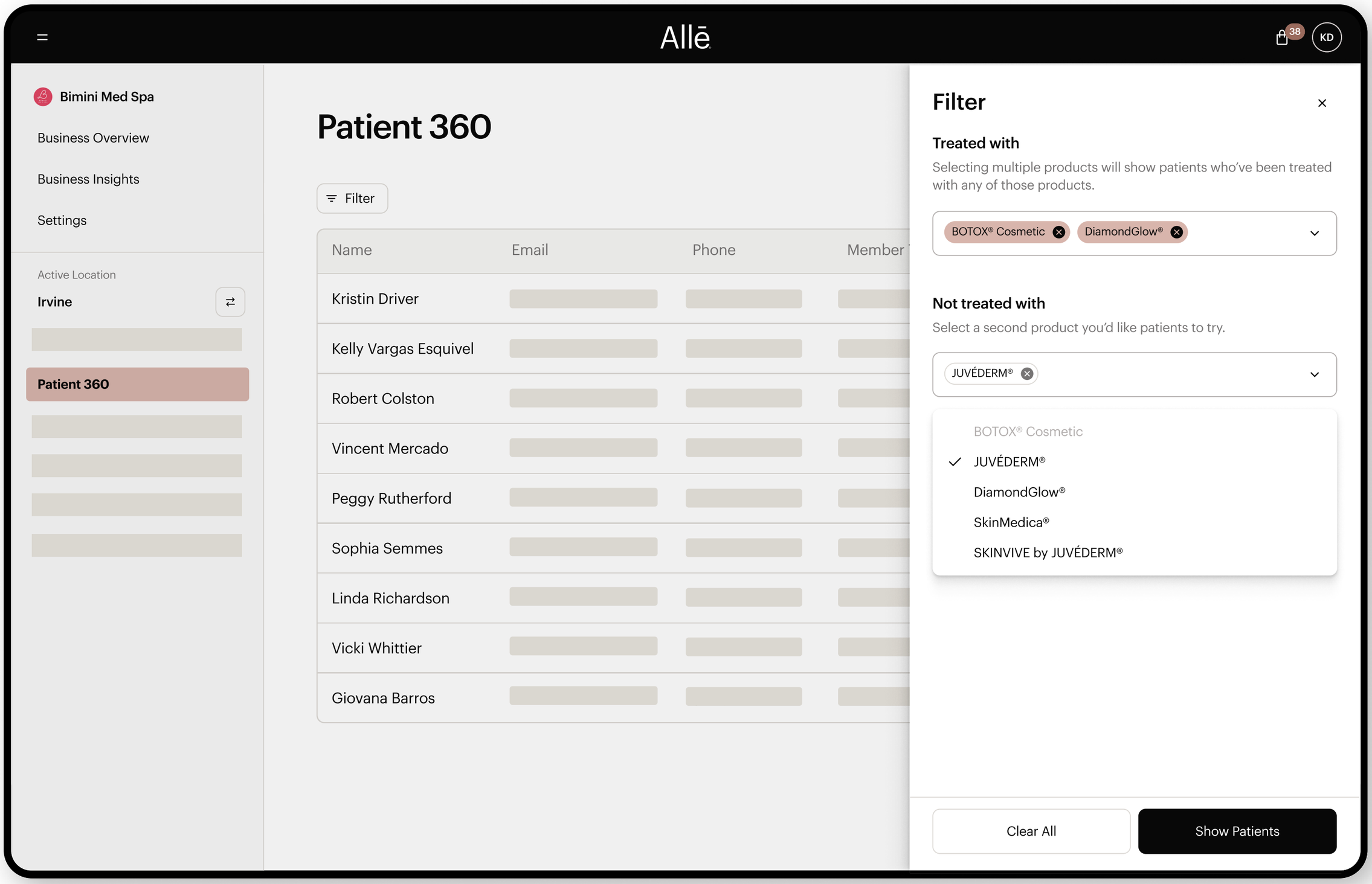
- Click the widgets to view a list of the last 50 patients who match the selected criteria.

Where to find them:

Log in to Allē > *Business Overview* and *Location Dashboard* Tabs

Please see Indications and Important Safety Information including Boxed Warning for BOTOX® Cosmetic (onabotulinumtoxinA) on page 5 and Indications and Important Safety Information for the JUVÉDERM® Collection of Fillers on page 5.

Patient 360: Comprehensive patient and practice reporting



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

Dig into your practice's Allē activity and connect on a more personalized level with the Allē Members who visit your practice.

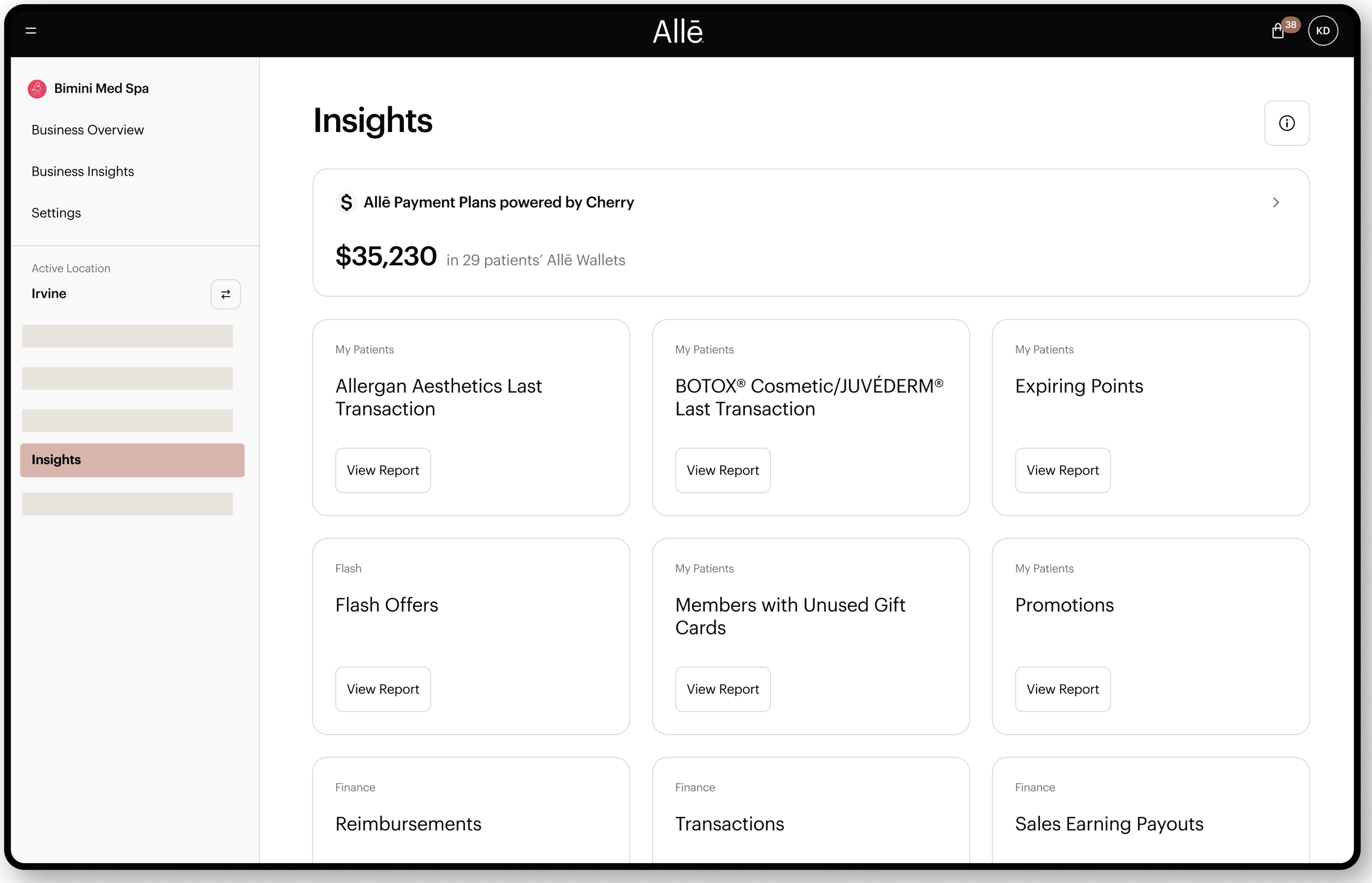
- Using the *Filter* button in your Patient 360 view, you can create a list of patients who have or have not been treated with a specified product or treatment.
- Patient 360 is viewable directly in Allē, and can also be downloaded as a report for easier distribution to your team members.
- Use the information in your Patient 360 to provide targeted treatment recommendations and help patients schedule their next appointments.

Where to find it:

Log in to Allē > Navigation Menu > *Patient 360* Tab

Please see Indications and Important Safety Information including Boxed Warning for BOTOX® Cosmetic (onabotulinumtoxinA) on page 5 and Indications and Important Safety Information for the JUVÉDERM® Collection of Fillers on page 5, DiamondGlow® on page 5, SKINVIVE by JUVÉDERM® on page 6 and SkinMedica® on page 6.

Insights reports provide a deeper look



Access a library of reports with a designated focus.

- View your practice's Allē transactions and reimbursements, patient treatment statuses, promotions, gift cards and Payment Plans available to your patients, and more.
- Stay on top of patient trends, make stronger data-driven decisions, and support your patients' full treatment journeys by regularly reviewing your Insights reports.

Where to find it:

Log in to Allē > Navigation Menu > *Insights* Tab

Please see Indications and Important Safety Information including Boxed Warning for BOTOX® Cosmetic (onabotulinumtoxinA) on page 5 and Indications and Important Safety Information for the JUVÉDERM® Collection of Fillers on page 5.

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
- Moderate to severe platysma bands associated with platysma muscle activity

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.

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 Equivalency Between Botulinum Toxin Products

The potency Units of BOTOX® Cosmetic are specific to the preparation and assay method utilized. BOTOX® Cosmetic is not equivalent to 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 preexisting 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 a reaction occurs, discontinue further injection of BOTOX® Cosmetic and immediately institute appropriate medical therapy. 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.

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 preexisting cardiovascular disease. Use caution when administering to patients with preexisting cardiovascular disease.

Increased Risk of Clinically Significant Effects With Preexisting Neuromuscular Disorders

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*). Monitor individuals with peripheral motor neuropathic diseases, amyotrophic lateral sclerosis or neuromuscular junction disorders (eg, myasthenia gravis or Lambert-Eaton syndrome) when given botulinum toxin.

Dysphagia and Breathing Difficulties

Treatment with BOTOX® and other botulinum toxin products can result in swallowing or breathing difficulties. Patients with preexisting 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*).

Preexisting Conditions at the Injection Site

Use caution 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 a 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), which would also be considered 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%).

The safety profile of BOTOX® Cosmetic treatment of platysma bands is consistent with the known safety profile of BOTOX® Cosmetic for other indications.

DRUG INTERACTIONS

Coadministration 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 injectable gel is indicated for deep (subcutaneous and/or supraperiosteal) injection for cheek augmentation to correct age-related volume deficit in the mid-face, for augmentation of the chin region to improve the chin profile, and for supraperiosteal injection to augment the temple region to improve moderate to severe temple hollowing in adults over the age of 21.

JUVÉDERM® VOLUX® XC injectable gel is indicated for subcutaneous and/or supraperiosteal injection for improvement of jawline definition in adults over the age of 21 with moderate to severe loss of jawline definition.

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® VOLBELLA® XC injectable gel is indicated for injection into the lips for lip augmentation and correction of perioral rhytids, and for the improvement of infraorbital hollowing 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® Ultra XC injectable gel is also 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, after insertion of the needle and just before injection, the plunger rod can be withdrawn slightly to aspirate and verify the needle is not intravascular, 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 are knowledgeable about the anatomy and the product(s) for use in indicated area(s), and who have appropriate training in facial anatomy, vasculature, safe injection techniques, and identification and management of potential adverse events, including intravascular complications
- 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 indicated areas for each product have not been established in controlled clinical studies
- The safety for use of these products in patients with known susceptibility to keloid formation, hypertrophic scarring, and pigmentation disorders has not been studied
- The safety for use during pregnancy and in breastfeeding females has not been established
- The safety for use of JUVÉDERM® VOLUMA® XC has been established in patients between 35 and 65 years of age for cheek augmentation, 22 and 80 years of age for chin augmentation, and 32 and 82 years of age for improvement of temple hollowing
- The safety for use of JUVÉDERM® Ultra Plus XC and JUVÉDERM® Ultra XC in patients under 18 years, and the safety for use of JUVÉDERM® VOLUX® XC, JUVÉDERM® VOLLURE® XC, and JUVÉDERM® VOLBELLA® XC in patients under 22 years, has not been established
- Dermal filler implantation carries a risk of infection. Follow standard precautions
- Dermal fillers should be used with caution in patients on immunosuppressive therapy
- Patients taking medications 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
- If laser treatment, chemical peel, or any other procedure based on active dermal response is considered after treatment, or before skin has healed from a procedure prior to treatment, there is a possible risk of eliciting an inflammatory reaction at the injection site
- 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 using a cannula with JUVÉDERM® VOLUMA® XC for cheek augmentation in patients with Fitzpatrick Skin Types V and VI or to improve temple hollowing has not been established
- 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 adverse events with injectable gel implants, and late-onset nodules with use of JUVÉDERM® VOLUMA® XC
- Based on preclinical studies, patients should be limited to 20 mL of any JUVÉDERM® injectable gel per 60 kg (132 lb) body mass per year. The safety of injecting greater amounts has not been established
- Injection of more than 9 mL of JUVÉDERM® VOLUX® XC for improvement of jawline definition has not been studied

ADVERSE EVENTS

The most common 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.

To report an adverse reaction with any product in the JUVÉDERM® Collection, please call the Allergan® Product Support at 1-877-345-5372. Please visit rxabbvie.com for more information.

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

DiamondGlow® Important Information

Indication and Use

The DiamondGlow® device is indicated for general dermabrasion of the skin and also delivers topical cosmetic serums onto the skin.

Important Safety Information

DiamondGlow® is contraindicated in patients who have compromised skin quality including but not limited to, sunburned, chapped, irritated or broken skin, open wounds, active, weeping acne, cold sores, or herpetic ulcers. Ask your patient if they are pregnant, lactating, or have any medical conditions, including allergies, and usage of topical medication on the area to be treated.

Typical side effects include a scratchy, stinging sensation during the treatment and temporary tightness, redness or slight swelling after the treatment. Rare serious side effects may also occur and include severe skin irritation and allergic reactions. Cease use of the device immediately if any of these serious side effects are observed.

Patients should be advised to use a sunscreen with a sun protection factor of 30 or higher following treatment.

Consult the DiamondGlow® User Manual for a complete list of Contraindications, Warnings, Precautions, and potential side effects.

SkinMedica® Pro-Infusion Serums Disclaimer

SkinMedica® Pro-Infusion Serums are intended to meet the FDA's definition of a cosmetic product, an article applied to the human body to cleanse, beautify, promote attractiveness, and alter appearances. These products are not intended to be drugs that diagnose, treat, cure, or prevent any disease or condition. These products have not been approved by the FDA and the statements have not been evaluated by the FDA.

Allē®

SKINVIVE by JUVÉDERM® Important Information

INDICATIONS
SKINVIVE by JUVÉDERM® injectable gel is indicated for intradermal injection to improve skin smoothness of the cheeks in adults over the age of 21.

IMPORTANT SAFETY INFORMATION

CONTRAINDICATIONS
This product 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 this product.

WARNINGS
- Do not inject into blood vessels. Introduction of this product into the vasculature may lead to embolization, occlusion of the vessels, ischemia, or infarction. Take extra care when injecting soft-tissue fillers; for example, after insertion of the needle and just before injection, the plunger rod can be withdrawn slightly to aspirate and verify the needle is not intravascular, 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
- Injection site responses consist mainly of short-term inflammatory symptoms and generally resolve within 1 week. Refer to the ADVERSE EVENTS

PRECAUTIONS
- To minimize the risk of potential complications, this product should only be used by healthcare professionals who have appropriate training, experience, and who are knowledgeable about the anatomy at and around the site of injection
- Discuss all potential risks of soft tissue injections with patients prior to treatment and ensure patients are aware of signs and symptoms of potential complications
- Limit patients to 20 mL of any JUVÉDERM® injectable gel per 60 kg (130 lbs.) body mass per year. The safety of injecting greater amounts has not been established
- This product is intended for improving skin smoothness and fine lines of the cheeks. The safety and effectiveness of use in other areas of the body have not been established
- Injection of more than 6.0 mL of this product (initial and touch-up treatment combined) for improvement of skin smoothness and fine lines of the cheeks has not been studied
- As with all transcutaneous procedures, injections of the product carry a risk of infection
- The safety for use during pregnancy, in breastfeeding females, and in patients under 22 years has not been established
- The safety in patients with known susceptibility to keloid formation, hypertrophic scarring, or pigmentation disorders has not been studied
- This product should be used with caution in patients on immunosuppressive therapy
- Patients taking medications that can prolong bleeding (such as aspirin, nonsteroidal anti-inflammatory drugs, and warfarin) may experience increased bruising or bleeding at treatment sites
- Patients may experience late onset AEs with use of injectable gel implants, including this product
- This product should only be used by healthcare professionals who have appropriate experience and who are knowledgeable about the anatomy and the product for use in the face
- If laser treatment, chemical peel, or any other procedure based on active dermal response is considered after treatment, or before skin has healed from a procedure prior to treatment, there is a possible risk of eliciting an inflammatory reaction at the injection site

ADVERSE EVENTS
In clinical studies, injection site responses (ISRs) observed in >5% of treated subjects included redness, lumps/bumps, swelling, bruising, pain, tenderness, firmness, discoloration, and itching. Most ISRs were mild. Adverse events reported through postmarketing surveillance outside of the United States included inflammatory reaction, inflammatory nodule, unsatisfactory result, loss/lack of correction, allergic reaction, anxiety, vascular occlusion, infection, dry skin, increase/decrease in sensation, and abscess.

To report an adverse reaction with SKINVIVE by JUVÉDERM®, please call the Allergan® Product Support Department at 1-877-345-5372. Please see Directions for Use or visit SKINVIVEDFU.com for more information.

SKINVIVE by JUVÉDERM® is available only by a licensed physician or properly licensed practitioner.

SkinMedica® Important Information

CAUTION: Do not use Retinol Complex and Even & Correct Dark Spot Cream if you are pregnant, lactating, or planning to become pregnant. Mild redness, peeling, and irritation are expected when using Retinol Complex. Use a sunscreen and limit sun exposure while using this product and for a week following discontinuation.

Sunburn alert: AHA/BHA Exfoliating Cleanser, AHA/BHA Cream, Even & Correct Dark Spot Cream, and Even & Correct Brightening Treatment Pads contain an alpha-hydroxy acid (AHA) that may increase your skin's sensitivity to sunburn. Use a sunscreen and limit sun exposure while using this product and for a week following discontinuation.

Use a sunscreen with an SPF of 30 or higher during any sun exposure when using Even & Correct Advanced Brightening Treatment.

Most SkinMedica® products are intended to meet the FDA's definition of a cosmetic product, an article applied to the human body to cleanse, beautify, promote attractiveness, and alter appearances. These SkinMedica® products are not intended to be drug products that diagnose, treat, cure, or prevent any disease or condition. These products have not been approved by the FDA and the statements have not been evaluated by the FDA.

SkinMedica® Total Defense + Repair Broad Spectrum Sunscreens (SPF 34 and SPF 34 Tinted) and Essential Defense Broad Spectrum Sunscreens (Everyday Clear SPF 47, Mineral Shield Tinted SPF 32, and Mineral Shield SPF 35) are over-the-counter drug products which are formulated and marketed pursuant to FDA's governing regulations set forth at 21 CFR Part 352.

The PA rating System is used in Japan to classify UVA protection and is not an FDA requirement on sunscreens sold in the U.S.

SkinMedica® Purifying Foaming Wash and Acne Clarifying Treatment are over-the-counter drug products which are formulated and marketed pursuant to FDA's governing regulations set forth at 21 CFR Part 333 Subpart D.

Not all products are available in Canada. Subject to change at any time without prior notice.