LIMITED TIME OFFER!

If You Have Ever Considered Breast Augmentation, *Now* is the Time to See if It's Right for You!

KATELYN
Mother of Four
Natrelle® Style 20

Choose *Natrelle*® Gel and Get a FREE BOTOX® Cosmetic Treatment or LATISSE®!*†

*For a friend too!^*

*Only a physician can determine who is an appropriate patient for these products.

† For qualified augmentation patients only.

Call for a consultation today!

BOTOX® Cosmetic (onabotulinumtoxinA)
Important Information

Indication
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 people 18 to 65 years of age for a short period of time (temporary).

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 any time (hours to weeks) after injection of BOTOX® Cosmetic:

- **Problems swallowing, speaking, or breathing,** due to weakening of associated muscles, can be severe and result in loss of life. You are at the highest risk if these problems are pre-existing before injection. Swallowing problems may last for several months

- **Spread of toxin effects.** The effect of botulinum toxin may affect areas away from the injection site and cause serious symptoms including: loss of strength and all-over muscle weakness, 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. **If this happens, do not drive a car, operate machinery, or do other dangerous activities**

The dose of BOTOX® Cosmetic is not the same as, or comparable to, another botulinum toxin product.

There has not been a confirmed serious case of spread of toxin effect when BOTOX® Cosmetic has been used at the recommended dose to treat frown lines.

Serious and/or immediate allergic reactions have been reported. They 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.

Do not take BOTOX® Cosmetic if you: are allergic to any of the ingredients in BOTOX® Cosmetic (see Medication Guide for ingredients); had an allergic reaction to any other botulinum toxin product such as *Myobloc*® (rimabotulinumtoxinB), *Dysport*® (abobotulinumtoxinA), or *Xeomin*® (incobotulinumtoxinA); have a skin infection at the planned injection site.

Tell your doctor about all your muscle or nerve conditions, such as amyotrophic lateral sclerosis (ALS or Lou Gehrig's disease), myasthenia gravis, or Lambert-Eaton syndrome, as you may be at increased risk of serious side effects including severe dysphagia (difficulty swallowing) and respiratory compromise (difficulty breathing) from typical doses of BOTOX® Cosmetic.

Tell your doctor about all your medical conditions, including: plans to have surgery; had surgery on your face; weakness of forehead muscles, such as trouble raising your eyebrows; drooping eyelids; any other abnormal facial change; are pregnant or plan to become pregnant (it is not known if BOTOX® Cosmetic can harm your unborn baby); are breast-feeding or plan to breast-feed (it is not known if BOTOX® Cosmetic passes into breast milk).

Human albumin and spread of viral diseases. BOTOX® Cosmetic contains albumin, a protein component of human blood. The potential risk of spreading viral diseases [eg Creutzfeld-Jakob Disease (CJD)] via human serum albumin is extremely rare. No cases of viral diseases or CJD have ever been reported in association with human serum albumin.

Tell your doctor about all the medicines you take, including prescription and nonprescription medicines, vitamins, and herbal products. Using BOTOX® Cosmetic with certain other medicines may cause serious side effects. **Do not start any new medicines until you have told your doctor that you have received BOTOX® Cosmetic in the past.**

Especially tell your doctor if you: have received any other botulinum toxin product in the last 4 months; have received injections of botulinum toxin, such as *Myobloc*®, *Dysport*®, or *Xeomin*® in the past (be sure your doctor knows exactly which product you received); have recently received an antibiotic by injection; take muscle relaxants; take an allergy or cold medicine; or take a sleep medicine.

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.

For more information refer to the Medication Guide or talk with your doctor.

You are encouraged to report negative side effects of prescription drugs to the FDA. Visit www.fda.gov/medwatch or call 1-800-FDA-1088.

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

Important *Natrelle*® Breast Implant Safety Information
Indications:
Breast Augmentation. Breast augmentation includes primary surgery to increase the breast size, as well as revision surgery to correct or improve the result of a primary breast augmentation surgery.
NOTE: The FDA has determined that patients seeking breast augmentation with silicone-filled implants must be at least 22 years of age.

Contraindications
Breast implant surgery should NOT be performed in: Women with active infection anywhere in their body, Women with existing cancer or pre-cancer of their breast who have not received adequate treatment for those conditions, Women who are currently pregnant or nursing.

Warnings
Every effort should be made to avoid damage to the breast implants during surgery. Care should be taken to avoid the use of excessive force and to minimize handling of the implant during surgical insertion. Care should be taken when using surgical instruments in proximity with the breast
implant, including scalpel, sutures, and dissection instrumentation. Use care in subsequent procedures such as open capsulotomy, breast pocket revision, hematoma/seroma aspiration, and biopsy/lumpectomy to avoid damage to the implant. Do not contact the implant with disposable, capacitor-type cautery devices. Do not alter the implants or attempt to repair or insert a damaged prosthesis. Do not immerse the implant in Betadine solution. Do not re-use or resterilize any product that has been previously implanted. Breast implants are intended for single use only. Do not place more than one implant per breast pocket. Do not use the periumbilical approach to place the implant. Do not use microwave diathermy in patients with breast implants, as it has been reported to case tissue necrosis, skin erosion, and implant extrusion.

Precautions
Safety and effectiveness have not been established in patients with the following: Autoimmune diseases (for example, lupus and scleroderma), A weakened immune system (for example, currently taking drugs that weaken the body's natural resistance to disease), Conditions that interfere with wound healing and blood clotting, Reduced blood supply to breast tissue, Radiation to the breast following implantation, Clinical diagnosis of depression or other mental health disorders, including body dysmorphic disorder and eating disorders. Please discuss any history of mental health disorders with your surgeon prior to surgery. Patients with a diagnosis of depression, or other mental health disorders, should wait for resolution or stabilization of these conditions prior to undergoing breast implantation surgery.

Adverse events
The most commonly reported adverse events for *Natrelle*® Silicone Filled Breast Implants are: reoperation, implant removal with replacement, grade III/IV capsular contracture, implant malposition, and asymmetry. Other potential complications include: rupture, unsatisfactory results, pain, changes in nipple sensation, infection, hematoma/seroma, breastfeeding difficulties, calcium deposits in tissue around the implant, necrosis, delayed wound healing, chest wall deformity, and lymphadenopathy.

Important: For full safety information please visit www.natrelle.com or call Allergan Product Support at 1-800-433-8871. Caution: Rx only.

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