

NEW CLIENT HISTORY

First Name:	Date:		
	:Birth Date:		
Address:			
City:State:		_Zip Code:	
Cell Phone:	Home Phone:		
Email:	Occupation:		
How did you hear about us?			as authorities and a second of the second of
MEDICAL HISTORY			
Do you have any chronic medical conditions we	should know about?	Yes	No
If so, please list:	·		
Do you have any allergies to latex, medications,	herbal or natural supplements?	Yes	No
If so, please list:			····-
Do you have, or have you had, any changes in m	edical history recently?	Yes	No
Please list any and all current/past surgeries or s	urgical procedures.		
Have you taken Accutane within the past year?		Yes	No
Are you on any anticoagulants, daily Aspirin, M	otrin, or Advil?	Yes	No
Are you a smoker?	•	Yes	No
Do you have veneers on your teeth?		Yes	No
WOMEN ONLY:			
Are you pregnant?		Yes	No
Are you currently breast-feeding?		Yes	No
Are your menstrual cycles normal?		Yes	No
Additional information you would like your tech	nician to know:		
Client Signature:		Date:	
Witness:	······································	Date	



CLIENTS RIGHTS AND RESPONSIBILITES

We are committed to serving you with compassion, care, and respect. As one of our valued clients you are entitled to the following:

You have the right:

To be treated with respect and dignity.

To know the names and professional status of the person(s) serving you.

To privacy and confidentially.

To receive accurate information about your health-related concerns.

To know the effectiveness and potential side effects of all forms of treatment.

To participate in choosing the form of treatment best suited to your skin.

To review education and counseling about treatments.

To review your medical record with you clinician.

To amend your records.

To receive any information about potential services or related services.

You have the responsibility:

To seek medical attention promptly, and to provide useful feedback.

To be honest about your medical history.

To ask questions about anything you do not understand.

To follow health advice and instructions.

To report any significant changes in your health.

To show up to appointments or cancel 48 hours in advance.

I authorize Houma Family Dental to perform the treatment or procedures recommended. I acknowledge no guarantee; either expressed or implied has been made to me regarding the outcome of any treatment or process.

I fully understand that it is impossible for anyone to make a guarantee regarding the outcome of any medical treatments or procedures.

I authorize the release of information to a licensed physician of the facility's choosing for the purpose of professional interpretation and establishment of the recommendations.

Client Signature:	Date:
Reviewed by:	Date:



Botunlinum Toxin Type A: Botox * Cosmetic & Dysport * Consent Form

BOTOX Cosmetic is indicated for the temporary improvement in the appearance of moderate to server glabellar lines associated with corrugator and/or procesus muscles activity in adult patients \leq 65 years or age.

BOTOX® Cosmetic (onabotulinumtoxinA) for injection, is a sterile, vacuum-dried purified botumlinum toxin type A, produced from fermentation of Hall strain Clostridium botulinum type A grown in a medium containing casein hydrolysate, glucose, and yeast extract, intended for intramuscular use. BOTOX® Cosmetic blocks neuromuscular transmission by binding to acceptor sites on motor nerve terminals, entering the nerve terminals, and inhibiting he release of acetylcholine. This inhibition occurs as the neurotoxin cleaves SNAP-25, a protein integral to the intramuscularly at therapeutic doses, BOTOX® Cosmetic procedures partial chemical denervation of the muscle resulting in a localized reduction in muscle activity.

Administration of BOTOX® Cosmetic is not recommended during pregnancy. There are no adequate and well-controlled studies or BOTOX® Cosmetic in pregnant women, it is not known whether this drug is excreted in human milk. Because many drugs are excreted in human milk, caution should be exercised when BOTOX® Cosmetic is administered to a nursing woman.

DYSPORT^{$^{\infty}$} (abobotulinumtoxinA) is an acetylcholine release inhibitor and a neuromuscular blocking agent indicated for the temporary improvement in the appearance of moderate to server glabellar lines associated with procerus and corrugator muscle activity in adult patients ≤ 65 years of age.

The effects of DYSPORT[™] and all botulinum toxin products may spread from the area of injection to produce symptoms consistent with botulinum toxin effects. 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, particularly in those patients who have underlying conditions that would predispose them to these symptoms.

DYSPORT[™] is contraindicated in patients with known hypersensitivity to any botulinum toxin preparation or to any of the components in the formulation. The product may contain trace amounts of cow's protein. Patients known to be allergic to cow's milk protein should not be treated with DYSPORT[™]. DYSPORT[™] is contraindicated for use in patients with infection at the proposed injection site(s).

There are no adequate and well-controlled studies in pregnant women. DYSPORT[™] should be used during pregnancy only if the potential benefit justifies the potential risk to the fetus. It is not known whether DYSPORT[™] is excreted in human milk.



Dermal Fillers: Consent Form

A. PURPOSE & BACKGROUND

As my patient, you have requested my administration of Dermal Filler; used in the correction of moderate to severe facial wrinkles and folds. All medical and cosmetic procedures carry risks and may cause complications. The purpose of this document is to make you aware of the nature of the procedure and its risks in advance so that you can decide whether to proceed with the procedure.

B. PROCEDURE

- 1. This product is administrated via syringe, or injection, into the areas of the face sough to be filled with dermal filler to eliminate or reduce the wrinkles and folds.
- 2. As anesthesia, numbing medicine used to reduce the discomfort of the injection, may or may not be used.
- 3. The treatment site(s) is washed first with an antiseptic (cleansing) solution.
- 4. Dermal fillers are to be injected under your skin into the tissue of your face using a thin gauge needle.
- 5. The depth of the injections will depend on the depth of the wrinkles and their location.
- 6. Multiple injections may be made depending on the site, depth of the wrinkle and technique used.
- 7. Following each injection, the injector should gently massage the correction site to conform to the contour of the surrounding tissues.
- 8. If the treated area is swollen directly after the injection, ice may be applied on the site for a short period.
- 9. After the first treatment, additional treatments may be necessary to achieve the desired level of correction.
- 10. Periodic touch-up injections help sustain desired level of correction.

C. RISK/DISCOMFORT

- 1. Although a very thin needle is used, common injection related reactions could occur. These could include some initial swelling, pain, itching, discoloration, bruising or tenderness at the injection site. You could experience increased bruising or bleeding at the injection site if you are using substances that reduce blood clotting such as aspirin or non-steroidal anti-inflammatory drugs as Advil.
- 2. These reactions generally lessen or disappear within a few days, but may last for a week or longer.
- 3. As with injections, this procedure carries the risk of infection. The syringe is sterile and standard precautions associated with injectable materials have been taken.

- 4. Some visible lumps may occur temporarily following the injection.
- 5. Some patients may experience additional swelling or tenderness at the injection site and on rare occasions, pustules may form. These reactions might last for as long as two weeks and in appropriate cases, may need to be treated with oral corticosteroids or other therapies.
- 6. Dermal fillers should not be used in patients who have experienced hypersensitivity, those with severe allergies to latex or xylocaine products (including but not limited to: xylocaine, nonacaine, zylocaine, benzocaine, prilocaine, or tetracaine) and should not be used in areas with active inflammation or infections (e.g., cysts, pimples, rashes or hives).
- 7. If you are considering laser treatment, chemical peels or any other procedure based on skin response after dermal fillers, or if you recently had such treatments and the skin has not healed completely, there is a possible risk of an inflammatory reaction at the implant site.
- 8. Most patients are pleased with the results of dermal fillers. However, like any cosmetic procedure, there is no guarantee that you will be completely satisfied. There is no guarantee that wrinkles and folds will disappear completely, or that you will not require additional treatments to achieve the results you seek. While the effects of dermal fillers can last longer than other comparable treatments, the procedure is still involving additional injections for the effect to continue.
- 9. After treatment, you should minimize exposure of the treated area to excessive sun or UV lamp exposure and extreme cold weather until any initial swelling or redness has gone away.

D. ALTERNATIVES

This is strictly a voluntary cosmetic procedure. No treatment is necessary or required. Other alternative treatments include, but are not limited to Botox, Laser Skin Modalities and Cosmetic Surgery.

E. CONSENT

Your consent and authorization for this procedure is strictly voluntary. By signing this consent form, you hereby grant authority to your physician's office/authorized medical spa facility to preform Facial Augmentation and/or Filler Therapy injections using the Dermal Filler of your choice for any related treatment as may be deemed necessary or advisable in the treatment areas you choose.

The nature and purpose of this procedure, with possible alternative methods of treatment as well as complications, have been fully explained to my satisfaction. No guarantee has been given by anyone as to the results that may be obtained by this treatment.

I have read this informed consent form and certify that I understand its contents in full. I have had enough time to consider this information from my physician's office/authorized medical spa facility, and I feel that I am sufficiently advised to consent to this procedure. I hereby give my consent to this procedure and have been asked to sign this form after being fully informed of the risks and benefits involved.

PRINT NAME:	DATE:
PATIENT SIGNATURE:	
INJECTOR:	DATE:

I authorize and direct		to perform the following procedure of
Botox ® Cosmetic and Dysp	ort [™] injections on	
(patient name) for the treatn	nent of (areas to be treated:	
□□Glabella	Initials:	
		nitials:
□□Crows F		nitials:
□ Other:	midais.	
Please initial the following:		
The deta	ails of the procedure have been e	xplained to me in terms I understand.
Alternat	ive methods and their benefits a	nd disadvantages have been explained to me.
	tand that the FDA has only approved the brows. Any other cosm	oved the cosmetic use of Botox® Cosmetic and etic use if considered off label.
Dysport [™] injections.	tand and accept the most likely r	isks and complications of Botox ® Cosmetic and
Including but not lin	nited to:	
•	Paralysis of a nearby muscle that could interfere with opening of eye(s).	
•	Local numbness	
•	 Headache, nausea, or flu-like symptoms 	
•		
•		
•	Disorientation and double vision	on
•	Temporary asymmetrical appe	arance
•	Abnormal or lack of facial exp	ression
•	Inability to smile when injected	in the lower face
•	Facial pain	
•	Product ineffectiveness	
I underst	and and accept that the long-tern	n effects of repeated use of Botox ® Cosmetic and
	——————————————————————————————————————	cations that have been identified but are not limited
to:	•	
•	Muscle Atrophy	
3	31	
•	Production of antibodies with	inknown effect to general health
Lundami	and and accept the less sommer	complications including
serious disability that exists		complications, including remote risk of death or

complications.	I am aware that smoking during the pre and post-operative period could increase chances of
The state of the s	I have informed the doctor or nurse of all my known allergies, including allergies to latex.
	I have informed the doctor or nurse of all medications I am currently taking including remedies, herbal therapies, and any other.
surrounding the pro	I have been advised whether I should take any or all of the medications on the days ocedure.
made or implied.	I am aware and accept that no guarantees regarding the result of the procedure have been
	I understand that I will receive my treatment in a training environment and the medical ming this treatment is being supervising by an experienced injector. (not necessary to initial ments)
recommendation, I	I understand that the medical professional supervising my injector will recommend the that he/she believes is appropriate for the results that I desire. If I chose not to accept that understand that I may not achieve the desired result and any further treatments to achieve the equire full payment.
treatment. Any add	Prices are subject to change. The pricing I receive during this treatment is only for today's litional treatment, products or services will be billed at rates in effect at time of the ts.
procedures I can de	I have been informed of what to expect post-treatment, including but not limited to if I wish to maintain the appearance that this procedure provides me.
while using Botox	I am not currently pregnant or nursing, and I understand that should I become pregnant ^D Cosmetic or Dysport [™] there are risks including fetal malfunction.
I understand that th	If pre and post-treatment photos and/or video are taken of the treatment for record purposes, ese photos will be the property of the attending doctor or nurse.
-	The doctor and/or nurse has answered all of my questions regarding this procedure.
respiratory disorder	I have been advised to seek immediate medical attentions if swallowing, speech, or s arise.
filled in PRIOR to	I certify that I have read and understand this agreement and that all spaces for initials were my signature.
Patient Signature: _	Date:
alternatives of the punderstands what I	I certify that I have explained the nature, purpose, benefits, risks, complications, and proposed procedure to the patient. I have answered fully, and I believe that the patient fully have explained.
Doctor Signature:	Date: