

## INFORMED CONSENT FOR TREATMENT OF FACIAL RHYTIDS

## REJUVENATION WITH BONTA (BOTOX® COSMETIC (OnabotulinumtoxinA), XEOMIN® (incobotulinumtoxinA), DYSPORT® (AbobotulinumtoxinA)

Patient Printed Name:		Date	
Diagnosis: Facial	wrinkles directly related to mus	scle contraction.	
Dysport® by of the following ard lines. Off-label site	with the following BoNTA pro Dr. Susan Geerlings , t eas: FDA approved sites of injects of injection may include brown nuscle, masseter muscles, or necessity	o treat lines and/or wrinkles ection of forehead lines, crow w lift, lower eyelid, bunny lin	in one, two or all w's feet, frown

NATURE AND PURPOSE OF THE PROCEDURE: The injection of BoNTA is a cosmetic procedure the FDA has approved only for wrinkle reduction in the glabellar region. Injection into any area other than the glabellar area is considered off-label use. The treatment plan is to inject an appropriate amount of BoNTA, a purified Neurotoxin produced by the Clostridium Botulinum bacteria, into a targeted facial muscle to intentionally produce weakness or temporary paralysis of the injected muscle. Relaxation of the muscle should improve lines and wrinkles that the targeted muscle action produced or improved contour of the face. Response is usually seen in 2 to 6 days after injection. It is common for the muscle's action along with its associated wrinkles to return in 3 to 6 months. Repeat injections are necessary to maintain the effects received. Lines and wrinkles present when the face is at rest may not improve with treatment of BoNTA alone, since BoNTA is designed to treat lines caused by facial muscle action. (pt intials)

**DISCLAIMER OF GUARANTEES AND EXPLANATION OF MATERIAL RISKS:** The practice of medicine is not an exact science and no guarantees or assurances have been made concerning the outcome and/or the result of this procedure. Injections with BoNTA is routinely performed without incident, however, there are some material risks. I understand that it is not possible to list every risk for this procedure and this consent form only attempts to identify the most common material risks which are headache, bruising, pain during injection, infection, asymmetry, twitching, numbness, and drooping of eyelids or eyebrows. I understand that some patients may not respond to the injection of BoNTA for unknown reasons. I understand fewer facial expressions will be possible after my injection with BoNTA. (pt intials)

**MEDICAL HISTORY:** I understand <u>Dr. Susan Geerlings</u> will provide my treatment and will rely on my documented medical history, as well as other information obtained from me in determining whether to perform this procedure. I agree to provide accurate and complete

information about my medical history and conditions. I herein state that I am **not pregnant**, **nursing or have any known neurological diseases**. If taking aminoglycoside antibiotics, Penicillin or Quinine, I understand these medications may potentiate the effect of BoNTA.

(pt initials)

Important Inquiry for Botox Cosmetic	c Treatment:		
History:		YES	NO
Are you suffering from a skin infection/disease at the			
Do you have a history of any bleeding disorders?			
Do you have a history of heart disease?			
Could you be or are you currently pregnant/breastfo			
Have you had Botox Cosmetic in the past?	•		
If yes, what was the last treatment date:			
Have you ever had any adverse reaction to Botox C	Cosmetic?		
Do You or Any Family Members Have a History o	f the Following:		
Amyotrophic Lateral Sclerosis	_		
Motor Neuropathy			
Myasthenia Gravis			
Lambert-Eaton Syndrome			
Facial nerve (Bells) Palsy			
<b>FOLLOW UP TREATMENT:</b> I agree to fol contact the office at 770-502-0350 and advise of may experience. (pt initials)			
<ol> <li>I have read or had this Consent Form re</li> <li>I fully understand the contents of this C</li> <li>I have been given ample opportunity to answered satisfactorily.</li> <li>I understand the risks and potential com</li> <li>No guarantees have been made concern</li> <li>I hereby voluntarily request and give my conse procedure described herein, injection of BoNT.</li> </ol>	and and/or explained to me consent Form. ask questions and all questions have applications of the treatments ing the results nor the outcome of the treatments of the for Dr. Susan Geerlings to perform	is pro	cedure.
PATIENT SIGNATURE:	DATE:		
WITNESS SIGNATURE:	DATE:		