INFORMED CONSENT FOR BOTULINUM TOXIN TREATMENT

Patient
Date of Birth
Address
Phone#
The purpose of this informed consent form is to provide written information regarding the risks, benefits and alternatives for the procedure named above. This material serves as a supplement to the discussion you have with your doc tor/healthcare provider. It is important that you fully understand this information, so please read this document thoroughly. If you have any questions regarding the procedure, ask your doctor/healthcare professional prior to signing the consent form.
THE TREATMENT Botulinum toxin (Botox and similar agents) is a neurotoxin produced by the bacterium Clostridium A. Botulinum toxin can relax the muscles on areas of the face and neck which cause wrinkles associated with facial expressions or facial pain. Treatment with botulinum toxin can cause your facial expression lines or wrinkles to be less noticeable or essentially disappear. Areas most frequently treated are: a) glabellar area of frown lines, located between the eyes; b) crow's feet (lateral areas of the eyes); c) forehead wrinkles; d) radial lip lines (smokers lines), e) head and neck muscles. Botox is diluted to a very controlled solution and when injected into the muscles with a very thin needle, it is almost painless. Patients may feel a slight burning sensation while the solution is being injected. The procedure takes about 15-20 minutes and the results can last up to 3 months. With repeated treatments, the results may tend to last longer. Initial
RISKS AND COMPLICATIONS Before undergoing this procedure, understanding the risks is essential. No procedure is completely risk-free. The following risks may occur, but there may be unforeseen risks and risks that are not included on this list. Some of these risks, if they occur, may necessitate hospitalization, and/or extended outpatient therapy to permit adequate treatment. It has been explained to me that there are certain inherent and potential risks and side effects in any invasive procedure and in this specific instance such risks include but are not limited to: 1. Post treatment discomfort, swelling, redness, and bruising, 2. Double vision, 3. A weakened tear duct, 4. Post treatment bacterial, and/or fungal infection requiring further treatment, 5. Allergic reaction, 6. Minor temporary droop of eyelid(s) in approximately 2% of injections, this usually lasts 2-3 weeks, 7. Occasional numbness of the forehead lasting up to 2-3 weeks, 8. Transient headache and 9. Flu-like symptoms may occur. Initial
PREGNANCY, ALLERGIES & NEUROLOGIC DISEASE I am not aware that I am pregnant and I am not trying to get pregnant, I am not lactating (nursing), I do not have any significant neurologic disease including but not limited to myasthenis gravis, multiple sclerosis, lambert-eaton syndrome, amyotrophic lateral sclerosis (ALS), and parkinson's. I do not have any allergies to the toxin ingredients, or to human albumin. Initial

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ALTERNATIVE PROCEDURES Alternatives to the procedures and options that I Initial	have volunteered for have been	fully explained to me.	
PAYMENT I understand that this is an "elective" procedure a that time of time of treatment. Initial	and that payment is my responsi	bility and is expected at	
RIGHT TO DISCONTINUE TREATMENT I understand that I have the right to discontinue to	reatment at any time. Initia	al	
PUBLICITY MATERIALS I authorize the taking of clinical photographs and marketing purposes. I hold McDowell Dentistry h waive my rights to any royalties, fees and to inspendent of the property of the propert	armless for any liability resulting ect the finished production as we	from this production. I	
RESULTS I am aware that when small amounts of purified weakness or paralysis of that muscle. This appear be shorter or longer. In a very small number of in for as long as usual and there are some individua be able to use the muscles injected as before while after a period of months at which time re-treatmerect posture and that I must not manipulate the period. Initial	rs in 2-10 days and usually lasts u dividuals, the injection does not Is who do not respond at all. I ur le the injection is effective but the ent is appropriate. I understand	up to 3 months but can work as satisfactorily or nderstand that I will not nat this will reverse that I must stay in the	
I understand this is an elective procedure and I hereby voluntarily consent to treatment with botulinum toxin injections for facial dynamic wrinkles, TMJ dysfunction, bruxism and types of orofacial pain including headaches and migraines. The procedure has been fully explained to me. I also understand that any treatment performed is between me and the doctor/healthcare provider who is treating me and I will direct all post-operative questions or concerns to the treating clinician. I have read the above and understand it. My questions have been answered satisfactorily. I accept the risks and complications of the procedure and I understand that no guarantees are implied as to the outcome of the procedure. I also certify that if I have any changes in my medical history, I will notify the doctor/healthcare professional who treated me immediately. I also state that I read and write in English.			
Patient Name(Print) Pati	ent Signature	Date	

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am the treating doctor/healthcare professional. I have discussed the above risks, benefits, and alternatives with the patient. The patient had an opportunity to have all questions answered and was offered a copy of this informed consent. The patient has been told to contact my office should he/she have any questions or concerns after this treatment procedure.				
Doctor Name (Print)	Doctor Signature	Date		