

BOTULISM TOXIN TYPE A CONSENT FORM BOTOX, DYSPORT, XEOMIN

Botox, Dysport and Xeomin is made from Botulism Toxin Type A; a protein produced by the bacterium Clostridiu, botulinum. For the purpose of improving the appearance of wrinkles, small doses of the toxin are injected into the affected muscles blocking the release of a chemical that would otherwise signal the muscle to contract. The toxin thus paralyzes or weakens the injected muscle. The treatment begins to work in 24 to 48 hours with results lasting up to 3-4 months. The Food and Drug Administration (FDA) have approved the cosmetic use of Botulinum Toxin Type A for the temporary relief of moderate to severe frown lines between the brow recommends that the procedure be performed no more frequently than once every three months. Botox has also been approved for crossed eyes (strabismus), eyelid spasm (blepharospasm), cervical dystonia (spastic muscle disorder of the neck), motor disorders of the facial nerve (VII Cranial nerve) and Hyperhidrosis (severe under arm sweating).

- Paralysis of nearby muscle that could interfere with opening the eye(s)
- Local numbness
- Headache, nausea, or flu like symptoms
- Swallowing, speech or respiratory disorders
- Swelling, bruising or redness at injection site
- Disorientation, double vision or past pointing
- Temporary asymmetrical appearance
- Abnormal or lack of facial expression
- Inability to smile when injected into the lower face
- Facial pain
- Product ineffectiveness

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Patient Consent: I certify that I have read an signature,	d unders	tand this treatment a	agreement and that all blanks are filled in prior to my
Patient Signature and date			Witness signature/date
Print Patient Name			Print Witness Name
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Physician signature			Date
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Original placed in chart	 Date	 Initials	