

MEDICAL QUESTIONNAIRE PRIOR TO TREATMENT

Subject to medical secrecy.

Your Details			
First Name		Date of Birth	
Last Name		Phone Number	
Address		Email Address	
Postcode		How did you hear about us?	
GP's name, address and tel:		Do you use sunbeds? Yes No	
		Do you smoke? Yes No	
		Do you drink alcohol? Yes No	
Medical Information Have you suffered from any of the follo	wing? If yes inlease tick		
Heart Disease/Angina	Auto-immune disea	se HIV/Hepatitis	
High/Low Blood Pressure	Stomach Ulcer/Coli	<u> </u>	
		<u> </u>	
Depression	Thyroid Problems	Skin disease (Acne)	
Glaucoma/Cataract	Diabetes	Facial Cold Sores	
Bell's/Facial Palsy	Asthma/Bronchitis	Convulsion	
Other			
Do you practice sport? Yes No if yes, please specify			
Do you have a blood clotting disorder/ require anti-coagulant treatment? Yes No			
Are you pregnant, planning a pregnancy or breast feeding? Yes No			
Are you currently taking medication? Yes No No			
Have you had any surgery in the past 3 months? Yes No			
Do you have any allergies? Yes No			
If you have answered YES to any of the above, please provide details			
Previous Treatment			
Have you already had aesthetic treatment? Yes No			
What was it with? HA Dermal Filler Botulin Toxin Other			
Date of Treatment (month / year)	Treatment Are	as Name of products	
	INFORMATION S	 HEET	
Refore:	the treatment please read t		

Don't hesitate to ask questions if you feel the information is not clear
Your practioner, who is trained in the treatment techniques, will be available to answer your questions
Take the time you need before making your decision

1. PRODUCTS AND INDICATIONS

The product used includes cross-linked and non-cross-linked HA gels (cross-linking is a process which can transform a liquid gel into a viscoelastic gel) as well as gels with or without anaesthetic (lidocaine).

The product used is designed for filling wrinkles and lines, contouring the face/body, skin rejuvenation or increasing lip volume. This product has a 6 to 18-month duration, depending on several factors; skin type, the severity of the wrinkles to be corrected, the injection zone and the volume injected. Your practioner will help you choose the product for injection according to your desired results.

2. PRECAUTIONS FOR USE AND CONTRAINDICATIONS

- Pregnant or breast-feeding women
- Sports persons have to be alerted on the fact that this product contains an active compound which may lead a positive reaction to doping testing
- History of hypersensitivity to one of the components of the products tested (hyaluronic acid, lidocaine, vitamins) or of anaphylactic shock or serve allergy
- History of autoimmune disease or disease affecting the immune system (type 1 diabetes, polyarthritis, rheumatoid arthritis, ankylosing spondylitis, psoriasis, thyroid disorder, scleroderma, inflammatory intestinal disease, lupus, multiple sclerosis, ulcerative colitis)
- Pathology (herpes, acne, rosacea) or unhealed skin alteration
- Complications after surgery during the past 5 years
- Previous injection of permanent products (silicone, acrylic, polymers, dextran)
- Untreated infectious periodontitis, cellulitis or dental or ENT origin, dental abscess untreated or treated less than one week ago
- In association with a peeling, a laser or ultrasound treatment.

3. PRECAUTIONS FOR USE AND CONTRAINDICATIONS

The filler used has been available commercially for many years, with several million syringes injected. Based on current data, there is no reason to suspect any unknown risks. According to international literature and health authorities, hyaluronic acid-based products may potentially have side effects. Indeed, although hyaluronic acid is a natural constant of the dermis, an injection of hyaluronic acid is likely to cause a skin reaction as if this molecule was a foreign body. These reactions are usually temporary but influenced on the one hand, by many external factors (type of product, technique, site, number of injections and quantity of product injected), on the other hand, by factors specific to the person being injected (injection tolerance, nervousness at the time of injection, medical history).

- Dissatisfaction with the expected aesthetic results
- Redness, bruising, ecchymosis, haematoma, oedema, itching, mild pain at the injection point which may occur after the injection and is resorbed after 24 hours to 8 days (on average 72 hours)
- Indurations or nodules which may occur at the injection point 15 days to 3 months after the injection
- Discolouration of the injection zone.

I have also been informed that rare cases of medical device vigilance have been described in the literature, necrosis in the glabellar region, abscess, granuloma and hypersensitivity following injections of hyaluronic acid, however if you notice a side effect after an injection, you must contact your practioner immediately.

I agree to receive MyFiller dermal filler injections.
The area to be treated is
I hereby authorise to treat me using MyFiller dermal filler. I understand that the effects may not be 100% and that multiple treatments may be necessary to achieve the best results.
I understand that there are certain risks associated with MyFiller dermal filler. I certify that I have read the entire informed consent and I agree to all its provisions. I certify that I have had the opportunity to ask questions and those questions have been answered to my satisfaction. I fully understand the treatments conditions and procedure.
I agree to pay \pm for the above-mentioned services and understand there will be no refund for any performed services. This consent form and cost covers above mentioned treatments only. Additional treatments can be added to this consent form and will be charged for as per clinic price list.
I have been made award of the risk and I accept these terms and conditions as part of my treatment. My practioner will not accept liability for any of the above side effects. By signing, I agree to the terms and conditions and in the event of any of the above, I or any of my representatives, will not pursue the practioner in any means of compensation.
The objectives and methods of the injection/treatment procedure have been clearly explained to me by the practioner
 I have received, read and understood the information supplied by the practioner prior to the injection/treatment I have had the opportunity to ask any necessary questions
 I understand the pre and post injection recommendations and I agree to follow them
 I acknowledge that I had the time required for consideration and to make my decision I acknowledge that I have been clearly informed of the side effects and the rare cases of medical device vigilance I freely and voluntarily consent to receiving injections/treatment
Client's Name
Client's Signature
Date
Practioner Name
Practioner Signature
Date
TREATMENT RECORD







