MEDICAL QUESTIONNAIRE & CONSENT FORM

prior to treatment



Subject to Medical Secrecy

YOUR DETAILS			
First Name Surname Address Postcode GP's Name, Address & Tel	Date of Birth Phone Number Email Address How did you hear about us? Do you use sun beds? Yes No Do you smoke? Yes No Do you drink alcohol? Yes No Do you drink alcohol?		
MEDICAL INFORMATION			
Have you suffered from any of the following? If yes, please tick			
Heart Disease/Angina Auto Immu High/Low Blood Pressure Stomach Ul Depression Thyroid Pro Glaucoma/Cataract Diabetes Bell's/Facial Palsy Asthma/Bro Other	Arthritis Skin Disease/Acne Facial Cold Sores Convulsions		
Do you practice sport? Yes No If yes, please specify			
PREVIOUS TREATMENT			
	Iin Toxin Other Date of Treatment (month/year) Treatment Areas		
Name of Products	Name of Products		

INFORMATION SHEET

Before the treatment please read this document carefully. 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

- 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 auto immune 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. 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.

GENIUST

CONSENT FORM
l agree to receive Genius dermal filler injections.
The area to be treated is
I hereby authorise to treat me using Genius 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 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
 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 Name Client Signature Date

Practioner Name.....

Practioner Signature

TREATMENT RECORD..... **PRODUCT STICKER** I confirm that I have been provided aftercare following my treatment. PRINT _____ DATE____ **PRODUCT STICKER**

I confirm that I have been provided aftercare following my treatment.

PRINT ______SIGN _____ DATE____

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PRINT ______SIGN _____ DATE_____

TREATMENT RECORD		
	A DO	PRODUCT STICKER
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PRINT	SIGN	DATE
		PRODUCT STICKER
Date		
I confirm tha	at I have been provide	ed aftercare following my treatment.

PRINT ______ DATE____