

BTL CELLUTONE™

GENERAL PATIENT RECORD

Date of birth:

Patient's name:	Age:	
Phone:	Email:	
You are scheduled for a series of non-invasive treatment	s with the BTL Cellutone device.	
Your treatment provider will discuss your specific treatme	ent needs. Recommended number of treatments is 4-6,	
with the frequency of 1-2 treatments per week. You may	need additional treatments depending on the severity of	
your condition. For optimal results, it is important to follo	ow the treatment plan that has been established for you.	
The results will typically continue to improve over the nex	ct few weeks. Initials:	
Please arrive at your appointment well hydrated. Ideally,	you should hydrate 2 days before, on the day	
of the treatment, and 4 days after the treatment. This wi	Il result in a more comfortable and efficacious treatment.	
Initials:		
On the day of the treatment, it is recommended to we	ar comfortable clothes which allow easy access to the	
treated area.		
You will be asked to remove any jewelry from the area of	interest. Initials:	
I acknowledge that successful treatment outcome can be	e affected by smoking or excessive alcohol consumption,	
as well as: eating disorders, on-going medication or inst	ufficient hydration. While no special diet is required, you	
are encouraged to eat healthy to help promote and main	tain results. Initials:	
There is typically no downtime associated with your trea	tment and there is no anesthetic required. Most patients	
describe		
the sensation of the therapy such as comfortable but inte	nse mechanical vibrations. Initials:	



<u>Ple</u>	Please answer whether you currently have or have had any of the following:				
•	Coagulation disorders, thrombosis NO		□ YES		
•	Cardiovascular disorders NO		□ YES		
•	Metal or electronic implants in the treated area, implanted cardiac pacemakers NO		□ YES		
•	Malignant or benign tumors	□ YES	□ NO		
	Diabetes mellitus	□ YES	□ NO		
	Arterial hypertension	□ YES	□ NO		
	Serious renal or hepatic insufficiency	□ YES	□ NO		
	Venous surgery on legs/sclerotherapy	□ YES	□ NO		
	ou answered YES to any of these questions, please specify:				
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For the full range of contraindications, warnings and cautions, consult your treatment provider.



•	I am aware that pregnancy and nursing are contraindicated and pregnant women can't undergo the
	treatment.
	Initials:
•	I understand there are certain risks associated with BTL Cellutone treatments and they include but are not
	limited to: local erythema, swellings, temporary loss of bodily sensation or itching, hematoma and petechiae.
	Initials:
•	I understand that the treatment may involve risks of complications or injury from both known and unknown
	causes, and I freely assume these risks. Initials:
	I agree to before and after treatment photographs, measurements and weighting, as this will help for medical
	evaluation of the results of the treatment. Information will be acquired for medical records or marketing
	purposes. Initials:
	I understand the results may vary from person to person and that an exact result cannot be predicted.
	Completing a full treatment series is necessary to maximize treatment efficacy. It is very unlikely but it is
	possible that you will not feel any recognizable result after the procedure. I acknowledge the results may not
	meet my expectations. Initials:
•	I certify that I have read this entire document and that I agree with all provisions. I certify that I have had the
	opportunity to ask questions and these questions have been answered in full to my satisfaction. I fully
	understand the treatment conditions, the procedure and possible side effects. Initials:
•	I have read the above information, and I request and give my consent to be treated with the BTL Cellutone
	procedure by the physician(s) in the below stated practice and his/her designated staff. Initials:
Му	signature below indicates that the above information is accurate and current.
Pat	ient signature: Date:
Wit	ness (in print):Signature:
Dat	e:
Pra	ctice Name: Marcos Medical Wellness

*For the full range of possible adverse effects and expected device-related treatment sequelae, consult your treatment provider.