INFORMED CONSENT & RELEASE FORM FOR BIOREGULATION (BioReg) SESSIONS

Bioregulation (BioReg) sessions are provided using Lenyosys LuxPro and Fractal devices and the Lenyosys CellCom biofeedback device. These devices are not designed to diagnose, treat, mitigate, prevent or cure any health condition. They provide general benefits recognized for biofeedback and PEMF (pulsed electromagnetic field) equipment, including promoting relaxation.

These are European CE certified medical devices and meet those regulatory requirements. The Lenyosys CellCom (LCC) is a non-invasive biofeedback device that captures and feeds back the body's bioelectrical signaling to the individual. Per Lenyosys, the manufacturer, the theory is that this may help stimulate the body's own inner cell signaling to support its recovery to help in balancing and calming the nervous system.

The LCC is a Biofeedback Class II Medical Device registered with the FDA (Product Code: HCC). The Lenyosys LuxPro (LLP) and Fractal are also non-invasive and feature extremely low intensity PEMF (pulsed electromagnetic fields) signals with complex frequency signatures. These are registered as Class I medical devices with the FDA. If you have a serious medical condition, the use of this technology should not replace any qualified medical advice you are currently receiving.

They Lenyosys Company indicates that the technology is designed to improve cell-to-cell communication which can allow the nervous system to "settle" or to "function more efficiently." Although the Center for Brain Training has many positive client reports from utilizing this technology, there is no research to bear out the described mechanism of action.

I agree that I have read the above information. I also understand I may experience an initial increase in my symptoms as the nervous system adapts, or I could have other non-typical symptoms, including being tired or achy. We recommend drinking more water than normal for 2-3 days after a session to help "clear" the system. You may choose at any time to stop the treatment. Our clinical experience is that the response for most people is neutral or positive. However, everyone is a unique individual, and responses can vary. Not everyone will notice a response to this technology.

I understand it is my responsibility to disclose accurately all information requested by the Center for Brain Training and to report any change in symptoms or any change to my medications. I agree NOT to hold CENTER FOR BRAIN TRAINING liable for any activities or results associated with the use of the Lenyosys devices. If using the Lenyosys Luxpro, I understand that I cannot have a pacemaker, defibrillator or potentially any other implanted stimulator in my body; the CellCom can be used with all of these devices.

I am not a minor (under 18 years of age). These devices have never been tested with pregnant women for safety. I have been given satisfactory answers to my questions concerning the use of these devices, and I give my full consent to their use.

I further acknowledge that I am fully aware that the LCC/LSC practitioner is not a licensed medical practitioner, but rather, a Certified LCC/LSC practitioner. I acknowledge that he or she has not made any promises of any kind to diagnose, treat, cure or otherwise address any medical issues I might be undergoing.

Signature: _____ DOB: _____

Printed Name: _____ Date: _____

Name of Dependent *(if applicable)*: _____