Informed Consent:

The EPFX, SCIO, EDUCTOR Biofeedback Medical device is registered in the USA, Europe, S Africa, Mexico, China etc. It is a TVEP Biofeedback device that measures how a person reacts to items. It is designed to measure VARHOPE TVEP reactions for allergy, homeopathy, nutrition, sarcodes, nosodes, vitamins, minerals, enzymes and many more items. Biofeedback is used for pre-diagnostic or therapy. These functions are registered in all of the above regions. Eclosion manufactures the hardware. Eclosion assists distribution of the EPFX, SCIO, Indigo and EDUCTOR software.

This software offers no risk and is completely safe. We recognize that this new type of system needs to be tested experimentally. The USA allows us to use an Institutional Review Board, Ethics Committee in Europe and operate an Investigational Device Testing for this software. To do any study in the world today we need to get informed consent from the patients or persons who are tested. Informed consent must be signed, implied, or understood.

The registered EPFX, SCIO, Indigo, EDUCTOR software and hardware uses a micro current medically safe pulse applied to the wrists, ankles and forehead. We safely measure some of the electrical aspects of the body. A variant micro current is then adapted to the patient to feedback the signal.

There is insignificant risk and the only discomfort is sitting still for the 30 or 40 min evaluation. The patient name will be held completely confidential in the study. Participation is always purely voluntary. There is no penalty for withdraws. And confidentiality is always guaranteed.

The results of the studies are to be published on the International Journal of the Medical Science of Homeopathy. http://ijmshnem4u.com/

The device and the study are always voluntary, confidential and safe. There are a wide amount of benefits already displayed by the thousands of users and millions of patients. A millions of people have already been helped. Results of the study and answers to your questions are available to the participants upon completion of the study.

I am informed of the experiment on the QX ltd software. I willingly give my consent to participate in the study. I give my consent for any children under my supervision or custody. I am to be guaranteed confidentiality of the data. I will be allowed to see the results of the publication in roughly one year. I recognize that there is no firm diagnosis resulting from the software. We are diagnosing and treating only Stress via Biofeedback.

I give my full and informed consent to partake in this research.

Name of Subject ___________________________________________________________

SIGNATURE________________________________________

DATE______________________________________________

THERAPIST OR WITNESS______________________________________________