This means our Scio/Eductor EPR TVEP is under patent protection
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CLINICAL EVALUATION

Only one device does all of the major Electrical Measures and the Electro-Stim all in a Cybernetic loop for safety and efficacy

measures
treats

Volts and Oscillations (EMG, EEG)
Amps and Oscillations (ECG)
Resistance (GSR)
Hydration
Oxidation (Redox potential)
Ph acid vs alkalinity
Reactivity evoked potential to voltammetric fields of substances (TVEP) over 228,000 measures a second of these energetic factors

Brain wave and emotions with (MCES)
Pain with (MENS) (TENS)
Trauma or wounds (EWH)
Electro Weakness Ph, Redox disorder (VARHOPE Correction)
Trickle charge the body electric

All designed to detect + reduce Electro-stress and Balance the Body Electric Automatically

If you need more information on the SCIO and purchase details please get in touch with us
web: www.qsbsubspace.com
e-mail: info@qsbsubspace.com
Introduction

On December 9, 1946, an American military tribunal opened criminal proceedings against 23 leading German physicians and administrators for their willing participation in war crimes and crimes against humanity. During World War II, German physicians conducted medical experiments on thousands of concentration camp prisoners without their consent. They shot concentration camp prisoners to test blood clotting. They infected groups of inmates with viruses, then only treated part with the test vaccines, while they observed how the course of the disease in the untreated inmates. They tested poison bullets to find more effective ways of killing; they tested prisoners to see how long they could remain alive under high altitude conditions of low air pressure and lack of oxygen.

The prosecution team submitted a memorandum outlining legitimate research to the Counsel for War Crimes, which was the basis for a section of the final verdict entitled "Permissible Medical Experiments." The ten points of the section have been subsequently referred to as "The Nuremberg Code."

In 1964, the World Medical Association established recommendations guiding medical doctors in biomedical research involving human subjects. The Declaration governs international research ethics and defines rules for "research combined with clinical care" and "non-therapeutic research." The Declaration of Helsinki was revised in 1975, 1983, 1989 and 1996 and is the basis for Good Clinical Practices used today.

Issues addressed in the Declaration of Helsinki include:

- Research with humans should be based on laboratory and animal experimentation
- Research protocols should be reviewed by an independent committee
- Informed consent is necessary
- Research should be conducted by medically/scientifically qualified individuals
- Risks should not exceed benefits

Every study that we have ever done has been compliant with all aspects of the Declaration of Helsinki and regulatory requirements. With that in mind, we will move to presenting the research we have conducted all over the world, in countries like France, Germany, Italy, Switzerland, England, China, Mozambique, South Africa, and United States of America.

For over 100 years The FDA in America has controlled manufacturing quality and claims of Food, Drugs and Cosmetics. In 1976 America started the first registration of medical devices. Thus controlling all medical device quality control and sales claims. Other countries have all followed suit. Notified bodies as independent businesses do this in Europe outside of Governmental control.

More and more validation and verification of sales claims has become a constant trend. Good science, bench studies, case studies, double blinks, peer reviewed journal publications for recognized journals and medical textbook publication in medical universities are all needed to assure validation and verification.

Accredited Medical Universities dictate the practice of medicine. The regulators do not determine what is or is not medicine. The Teaching Medical Universities and Medical Hospitals do. And they perform and supervise most research. The legal requirements of performing research have been a changing and evolving process. A gradual tightening of restrictions has made this more and more difficult.
We will review now a 40 plus year history of clinical research done at the legal requirements, informed consent and proper ethical and professional supervision. We have a constant desire to clinically investigate and validate all of our processes. Our clinical evaluation starts in Ohio, USA at Youngstown State University in 1973.

**Study done at Youngstown State University**

"The Trivector Signature of Human Beings as a Basis of Psychic Communication" 1974

Research with 26 pairs of Intimate couples showed a reactive Trivector body electric and a subspace transfer of information

---

**Early research**

We will start this presentation by going back in time more than 30 years ago, to the 1970’s, to discuss the significant study conducted at the Youngstown University by prof. William Nelson. In 1973, Prof. Nelson started studying the body electric on a group of subjects. The study was published in 1974 at the Post Graduate Department.

Prof. Nelson took 40 pairs of intimate friends (that is married people, couples going steady, very, very close friends). One of the pair was put into a dark room in a building in Youngstown State University. A stroboscope and a siren was placed into the dark room next to the ears and eyes of the person. At random times over the course of a two hour plus+ session, a one minute signal of stroboscopic burst and siren’s scream would be initiated, subjecting the patient to a fluctuation that would initiate an evoked potential brain wave response. This would provide a startle to the system, a threat to the system that, although safe, would be interpreted by the patient as possibly life-threatening. This would produce a hormonal and nerva1 reaction and would it would be discernible by their friend.

The other person in the group was placed in a separate building at Youngstown state campus. The separate building was needed to make sure that there was no electrical artifact in the electric measures that could be detected. This person was hooked up to a polygraph device capable of measuring the brainwave, the heart rate, and the galvanic skin response. Thus we were measuring voltage, amperage and resistance (VAR). During the two hour plus session the person hooked to the polygraph was to
make verbal guesses as to when they thought their friend was being subjected to the evoked potential shock. It is shown that the verbal guesses were less than chance. In other words there was no verbal ability to understand what was going on. However, there was always an electrophysiological voltammetric plus resistance reaction that could be discerned. There was over 90% correlation to a type of electrophysiological (VAR) reaction that could be determined. Thus, the article could show that there was a type of psychic communication. It was seen to be isolated from verbal, conscious pathways and was more of an unconscious cybernetic bioelectronic signal.

Prof. Desire’ Dubounet has done very significant work with Nobel Prize Winners William Fowler, Roger Sperry, Albert Szent Gyorgy and Hans Selye, one of the most recognized medical doctors in the world of stress. Selye’s work on accumulated stress as a cause of disease became a big part of Prof. Dubounet’s work.


Desiré has been a controversial provocative champion for freedom of choice and truth in medicine. In 1992 Desiré got Homeopathy to be legal real medicine in Hungary when she offered a course in Homeopathy for Semmelweis Medical University. Working with the Hungarian doctor Hans Selye to find that Stress is the main cause of disease. Her early life has been documented in a Full length movie “Intellect is the final frontier, Story of Desi’s Youth”. America was afraid of the Soviet threat.

Some of her earliest research is published in an internationally recognized, and registered peer review medical journal. She worked on the Apollo project as an electrical engineer for AC Electronics in 1969. Desiré has worked with several scientists including Nobel Prize winners and nominees Hans Selye, Albert St Gyorgi, Roger Sperry, and William Fowler. Desiré went to Pakistan in 1986 to open up natural medicine. Desiré has had a remarkable and yet controversial life of challenge and heroism.

Desiré wrote a 510k thesis to get acupuncture needles registered as medical equipment in America. Desiré was wrongfully and maliciously attacked for this. She won this battle, acupuncture needles were registered in 1996, but the FDA still harbors a grudge. The Ultra-Rich controlled media also attacks her.
Going forward to the 1980’, we will now take a look at the research conducted at the American Academy of Quantum Biofeedback Technology (AAQBT). The AAQBT, situated in Rio Rancho, New Mexico, U.S.A., has been the setting for a series of studies conducted over 4 years, from 1984 to 1988, on 935 patients, with the EPFX (Electro Physiological Feedback Xrroid System) device.

Conducted with proper IRB supervision, medical supervision, informed consent, these studies have been among the first ones to analyze the body electric and therefore represent groundbreaking scientific advances in the field of energetic medicine.

Over the 4 years, the following fields have researched:

- **Alarm Response** – in this part of the study it has been shown through statistically significant results that a stimulus that might be too much for the body provokes an alarm response.
- **Calibration** – it has been designed from EPR data to test the reaction speed of the patients
- **Electro – Acupuncture** – it showed that particular electrical signatures applied to unhealthy acu-points might cause them to improve
- **Electro – Physiological Reactivity (EPR)** – the study measured the subjects’ reactivity patterns to nosodes, allersodes, isodes, sarcodes, and classic homeopathy and proven an accuracy of approximately 71% percent to known medical conditions
- **Skin Capacitance** – the goal of this particular measurement was to further analyze the nature of the body electric by better understanding the basic skin capacity to store charges and other bio-electric measures
- **Skin Conductance** – the study describes the EPFX method for measuring the skin conductance responses
- **Skin Inductance** – this part of the study analyzed the skin’s capacity to affect inductance transfer
- **Trivector** – this review makes a comparison between skin conductance, capacitance and inductance (known as the Trivector) and SCIO Electro-Physiological-Feedback-Xrroid EPR reactivity
- **Stress Reduction through Electro Stimulation** – this part of the study showed with significant results that the EPFX treatment reduces stress; 76% of the 935 subjects tested declared they had reduced levels of stress, 14% reported they felt no different, and 10% reported they felt more stress.

Out of the 935 subjects tested over a period of 4 years, there have been no adverse events reported, proving that the device was safe to use to the indications for use.
Below we present you with the abstracts of the study discussed above.

Situated on a golf course in the Land of Enchantment, in the City of Vision the AAQBT made history.

From 1984 to 1988, 935 subjects were tested with EPFX.

With proper IRB supervision, informed consent, and medical doctor direction the AAQBT did a series of studies on the electro-physiology to validate the 1989 FDA registration of the EPFX.
The American Academy of Quantum Biofeedback Technology
Located in Rio Rancho, New Mexico since 1986

**Skin Impedance**

By William Nelson

**ABSTRACT**

Skin impedance is a measurement that involves the resistance of the skin to an electrical current. It is used to assess the integrity of the skin barrier and to monitor various physiological markers such as hydration, inflammation, and skin barrier function. This measurement can be useful in diagnosing skin conditions and evaluating the effectiveness of different treatments. Skin impedance can provide valuable insights into the skin's health and can help in personalized skincare routines and treatment plans.

*Published AAQBT Press 1988*

**Super Learning Language Stimulation with the EPFX**

By William Nelson

**ABSTRACT**

Super Learning Language Stimulation with the EPFX is a technique that utilizes biofeedback to enhance learning and language development. This method involves the use of bioelectrical signals to provide feedback to the learner, aiding in the acquisition and retention of new information. The EPFX device is designed to help individuals improve their language skills and cognitive abilities through the stimulation of specific brain wave patterns.

*Published AAQBT Press 1988*

**Stimulation of Math Skills with an Auto-Focused Cybernetic Loop Stimulation**

By William Nelson

**ABSTRACT**

Stimulation of Math Skills with an Auto-Focused Cybernetic Loop Stimulation is a method that utilizes biofeedback to improve mathematical skills. This technique involves the use of bioelectrical signals to provide feedback to the learner, helping to increase concentration, focus, and memory retention. The device is designed to enhance learning and problem-solving abilities in mathematical tasks.

*Published AAQBT Press 1988*

**Calibration**

By William Nelson

**ABSTRACT**

Calibration is a process that involves the adjustment of a device to ensure that it provides accurate and consistent measurements. In the context of biofeedback technology, calibration is crucial to ensure that the device functions correctly and provides reliable data. The EPFX device undergoes rigorous calibration procedures to ensure its accuracy and reliability, making it a valuable tool for various applications in healthcare and education.

*Published AAQBT Press 1988*
The American Academy of Quantum Feedback Technology

Located in the Americas, since 1950

A Q.F.T.

The American Academy of Quantum Biofeedback Technology

Located in Rancho, New Mexico since 1978

AAQBT

The American Academy of Quantum Biofeedback Technology

Located in the Americas, since 1978

Electro-Acupuncture

By William Nelson

AAQBT

The American Academy of Quantum Biofeedback Technology

Located in Rancho, New Mexico since 1978

Skin Conduction

By William Nelson

AAQBT

The American Academy of Quantum Biofeedback Technology

Located in Rancho, New Mexico since 1978

Electro-Pneumoplymetry

By William Nelson

AAQBT

The American Academy of Quantum Biofeedback Technology

Located in Rancho, New Mexico since 1978

AQPBT

To be Legal Requires
1. Informed Consent from all Subjects
2. Ethics Committee Supervision

Must be Voluntary and Provide Confidentiality, Safety, Consent, Feedback of Performance, Benefits of Participation, and to Society
510(k) Registration of the EPFX

This body of research led to the EPFX FDA 510(k) registration obtained on October 13, 1989. Inside the 510(k) registration we have the first appearance of the Electro-Physiological Reactivity (EPR) and the VARHOPE. This registration is still valid and can be found on the FDA website. A copy of our 510(k) is included for your reference.
The EPFX 510(k) registration was the first one in a long line of legal registrations obtained over the world.
The Long Term Pathological Findings of the Camelford Toxicity Group

A study done in Camelford, England on aluminum toxicity in 1992 has been reproduced in Budapest, Hungary in 2012.


Title
The Long-term Pathological Findings of the
Camelford Toxicity group. 1990

Subtitle
The Premature Ageing Effects of a Toxic Water Syndrome Case.

By
Dr. William Nelson LPCC, Peter Smith LCH

ABSTRACT:

In July 1988, a toxic water spilt in the Camelford water district by South West Water, the public water utility, in Cornwall England resulted in some 20,000 people being exposed to a toxic cocktail of chemicals in their drinking water. This produced a host of different physiological diseases. It also resulted in a homeopathic practitioner, together with other colleagues, launching a long-term 7 year study of the effects of this toxicity on the population.

Besides extensive case notes on 200 people, hair and nail samples, several different electrical measures, chemical measures, and psychological interventions have yielded a understanding of these patients' disease profile. Various lengthy papers have been prepared by the North Cornwall Homeopathic Project and the Lowermoor Support Group. A book is in preparation.

One of the key factors that have been observed in this population is that of premature aging. This is discussed within this article.
Hungarian XRROID analysis of the people exposed to the Aluminum Toxicity showed extremely sensitive to the same compounds as the Camelford England Aluminum accident 15 years earlier Proving the Meta-Analysis of the Xrroid technology

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<th>p-value</th>
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<td>12</td>
<td>0.001</td>
<td>SS</td>
</tr>
</tbody>
</table>

* = > 2 Standard Deviations
SS = Statistically significant
Dr. Bandics Istvan Research

Preceding the existence of the companies (Eclosion Kft., Maitreya Kft. and Eclosion ltd Kft.), but post ceding the 510(k) FDA registration, Dr. Bandics Istvan’s clinic was the location for several research projects on the SCIO, in 1995-1996, under the medical supervision of D. Istvan Bandics, who was following the work of Dr. Albert Szent-Gyorgy.

We present you a list of abstracts of the aforementioned studies:
Bio-electronic Increase of Power Lifting Performance Clinic

Details

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HIPPOCAMPUS
Egészségügyi Kereskedelmi- és Szolgáltató BT.
Adószám: 21284823-2-05
OTP BANK Rt.: 050-010767-9

RENDELO és IRODA
Budapest III.
Nánái út 67. I. em.
Tel.: 1-188-68-65
06-20-342-662

Supervising researcher: Dr Istvan Bandics MD Licensed Hungarian Medical doctor. This study was done at the Hippocampus office in Budapest in January 1994. Studies done with the supervision of a local ethics committee and all subjects gave informed consent to participate.

Abstract

This study took 18 members of a Hungarian Power lifting team and measured their performance before and after an EPFX therapy and some sport oxygen formula. Their personal best are a matter of record. Each had two sessions on the EPFX over two days and they were asked to do their best in Squats and bench press. By comparing to the personal bests most of these athletes had increases in performance after two sessions on the EPFX.

---

Electro-Physiological Stimulation of Body Charge Potentials

TELEPHELY
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Tel.: 49 / 311-026

HIPPOCAMPUS
Egészségügyi Kereskedelmi- és Szolgáltató BT.
Adószám: 21284823-2-05
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RENDELO és IRODA
Budapest III.
Nánái út 67. I. em.
Tel.: 1-188-68-65
06-20-342-662

Supervising researcher: Dr Istvan Bandics MD Licensed Hungarian Medical doctor. This study was done at the Hippocampus clinic in Budapest on 73 patients attending the clinic in 1994. Studies done with the supervision of a local ethics committee and all subjects gave informed consent to participate as part of their intake form.

Abstract

In this study we took 73 patient/subjects from the medical clinic and tested their global Voltage, Amperage, Resistance, Hydration and Oxidation body electrical parameters in per and post fashion. Each subject was treated with the EPFX device for one week after pre and before post testing. The patients had significant increases in their post electrical measures. These factors are called the VARHO.
In 1995, at the Szent Janos Hospital in Budapest, Prof. Dubouret has done work on cataract patients, showing significant results in TVEP patterns that can be helpful in detecting disease patterns.
In 1995, Semmelweis University Budapest was the location for a study that investigated the Zap capabilities of the QXCI/EPFX on Human Papilloma Virus. The results showed that overall, the therapy applied was more than 60% successful in treating papilloma virus.

Zapping the Human Papilloma Virus

By William Nelson LPCC

At the Semmelvise Hospital in Budapest 1994

Abstract

We know of no good evidence for Dr. Kruger’s Zap therapy. The Zap technology of the QXCI/EPFX has some tested capabilities. In this study 25 women showed signs of papilloma virus spots. The papilloma virus HPV spots fluoresce in UV light if they are exposed to vinegar. A vinegar swab showed spots on all 25 with an average of 12 spots per person. The women were given three 30 min QXCI Zap therapies over the course of one week. In 2 of the subjects there was no change. All of the others had lessening of their spots in size and number. In five subjects there was complete removal of the spots. The rest had approximately 60% reduction in the number of spots. The overall average therapy was 60% effective in treating papilloma.

A SHORT STUDY OF COMPARISON FACTORS OF COPROLITH VERSUS QXCI DETECTION OF INTESTINAL PARASITES

- Chief Editor: William Nelson, N. M.D.; Independent Medical Editor: Budapest, Hungary
- Edited and Validated By: Istvan Bandics, M.D; Budapest, Hungary Györra Panszki, M.D; Budapest, Hungary Attila Kiss, M.D; Győr, Hungary
- Consultant: Dr. Simon Gütj, M.D; Hanover, Germany
- Developed By: The staff of Maitreyo; Limerick, Ireland

This study was performed in 1984 at the King Health Center in Lowellville, Ohio, USA Revalidation and further clinical testing and has been repeated by medical doctors at the Homeopathy Clinic in Budapest, Hungary and by the doctors listed above.


This work was presented at the Singapore World Congress on Sexually Transmitted Diseases. This Body of Research has led to the first registration of the QXCI in 1997. The QXCI is a follow on the original EPFX now known as the Eductor.

Royal Society of Medicine

In 1992, Dr. Nelson/Desire’ Duboune was invited to the Royal Society of Medicine, where a group of 6 doctors lectured on this technology.
Dr. Nelson, now known as Desire’ Dubounet, has been nominated for the Nobel prize, and in 2000 was invited to speak at the Nobel Prize Hospital in Stockholm, Sweden.
Dr Nelson has been nominated for the Nobel Prize in Medicine many times. In 2000 he was invited to lecture in the Nobel Prize Hospital in Stockholm, Sweden. His lecture was a Revolution in Medicine. The Nobel Prize Hospital in Sweden Place of Desire' Dubounet's lecture in 2000.
A group of doctors from the Szent Janos Hospital and Semmelweis University in Budapest have gotten together and created the International Journal of the Medical Science of Homeopathy in 1997. This journal has been unchallenged and recognized for over 16 years. Over a hundred volumes have been published with over 1,000 studies.

The International Journal of the Medical Science of Homeopathy, Naturopathy and Energetic Medicine has been a reputable, well established, publically accessible and professional medical peer reviewed journal since its conception in 1997. The original international peer reviewed library registration number was ISSN 1417-0876, and currently it is ISSN 2041-4293. The first medical doctor that supervised the peer review process as a director was a Hungarian doctor, Judith Nagy M.D. After her retirement, she was replaced by Dr. Hiľf Klára, M.D., who is still the supervisor and director of the medical peer review process.

This journal has been credentialed and recognized by over 20 respected and recognized medical doctors, all very well acknowledged and respected in their field of application. Below we present some of the doctors, professionals and lay people that have been involved with the International Journal of the Medical Science of Homeopathy, Naturopathy and Energetic Medicine over the years:
The *International Journal of the Medical Science of Homeopathy, Naturopathy and Energetic Medicine* (IJMSHNEM) has operated without a single challenge to its credibility for over 20 years. It has published hundreds of articles, double blind studies, case studies, and clinical data on various topics focused on natural, energetic and alternative medicine.
IUMSHNEM Ethics Supervisor & Institutional Director: Dr. of Sociology Brad Victor Johnson, Dean of the International Medical University of Natural Education – IMUNE

Editorial Consultant: Alejandro Arregui Henk, Argentina,

IUMSHNEM Associates:

Alejandro Arregui Henk, Argentina, – Chris King, – Dr. Igor Cetojevic, Cyprus – Dr. Bill Cunningham, USA, – Dr. Kofi Gharretey, Africa, – Dr. Amanda Vellore, South Africa, – Dr. Danis Gyorgy MD, Hungary, – Dr. Debbie Draks, M.D., Canada, – Dr. Sarca Ovidiu, M.D., Romania, – Andreea Tafian, – Dr. Steve Small, USA, – Dr. Pauline Wills, USA, – Bala Lodhia, Canada, – Anna Marie Stinton, Romania, – Dr. Gulyas Kinga, Hungary, – Dr. Aurel Bacean, M.D., Romania, – Dr. Oana Bacean, M.D., Romania, – Dr. Theron Francois, M.D., South Africa, – Turako Yui, Homeopath, Japan, – Dr. Ho, M.D., China, – Gage Tarant, – Dr. Marco Antonio, Rodriguez Infante, Mexico, – Dr. Alex Von Pelet, Germany, – Dr. Hilf Klaud MD Hungary, – Dr. Bacean Aurel MD, Romania, – Dr. Gebhard Gerhing MD, Bavaria, Dr. Hobian Veronica, Romania, – Rita Nemenyi, Biofeedback Research Technician

Our Journal was started in 1997 and for decades has looked for evidence to support Homeopathy, Naturopathy and Energetic Medicine. We are dedicated to validation of this art.

Brad Victor Johnson
This is to inform you that the Homeo Diagnostical Academy Press has published

The International Journal of the Medical Science of Homeopathy

as a peer reviewed medical journal

under Hungarian ISSN National Center 1997 apr. 15-1 and has awarded the international identification number:

ISSN 1417-0876

This number will remain unchanged for the lifetime of this publication.

You are kindly reminded that with every future publication the ISSN number will be printed on each book. Should you wish to have this publication printed internationally the identification number can be found near by the Hungarian code which would be:

HU ISSN 1417-0876

If it is your preference to have a detailed explanation of this publication available you may visit the Hungarian ISSN National Center web pages where you will find the OSZK web page called “Local services” and in this menu point you would enter your identification number. The web address can be found at:

http://www.oszk.hu/index_hu.htm

In relation to this document or in any other point please contact us at your convenience.

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The journals can be accessed online at http://ijmshem4u.com/
Large Scale SCIO Study

100,000+ Study of the Safety and Efficacy of the TVEP families in the SCIO Device

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The Centro Ricerche, University of Venice + Padova, Italy

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This study was performed in the field by practicing Biofeedback technicians. Data was collected and the study supervised by the Ethics International Institutional Review Board of Romania. The Data analysis and study presentation is done By the The Centro Ricerche, University of Venice,

Padova, Italy; © Ethics International, 2008.
Large Scale SCIO—Eductor Clinical Study

Over 2,200 Medically Supervised Therapists, Clinically Tested over 100,000 patients with over 220 diseases and over more than 350,000 patient visits

2005 to 2008

To be Legal Requires
1. Informed Consent from all Subjects and
2. Ethics Committee Supervision

Must be Voluntary and Provide Confidentiality, Safety, Consent, Feedback of Performance, Benefits of Participation, and to Society

World’s Largest Clinical Biofeedback Peer Reviewed Research Study on the SCIO

2,225 Medical Supervised SCIO Therapists saw over 100,000 patients thru over 350,000 visits and 220 diseases all with Positive results here is a list of the studies
The SCIO device is a Universal Electro-Physiological device used for stress reduction and patient treatment. Over 2,200 qualified biofeedback therapists joined our Ethics Committee study to evaluate how stress reduction using the SCIO device could help a wide variety of diseases. The device and thus the study has insignificant risk. There was a staff of medical doctors who designed and supervised the study.

Over 100,000 patients gave informed consent and participated in the study. The study would conclusively prove safety and efficacy of the SCIO Device. With over 60% of these patients having...
multiple visits. There were over 300,000 patient visits and over 225 diseases investigated. With a total record of the SCIO patient information, therapy parameters and reactivity data. No names of patients were recorded for confidentiality.

Two of the 2,200 plus therapists were given blank devices that were completely visually the same but were none functional. These two blind therapists were then given 35 patients each. This was to evaluate the double blind component of the placebo effect as compared to the device. Thus the studied groups were a placebo group, a subspace group, and an attached harness group.

This is just the first study in a long task of analysis in truly break down the data totally. This study verifies the safety and efficacy of the SCIO device as well as the validity of the TVEP family reactivity. There were small effects seen in the placebo group, larger effects in the subspace, and astounding effects in the real harness group.
Eductor study of Voltammetric stimulation of Blood testosterone
1. Treating High Blood Pressure Hypertension Naturally
2. Treating High or Low Blood Pressure Naturally
3. Treating Inflammation Naturally
4. Stuttering
5. **TENDON CALCIFICATION Bone Spur**
6. **THE PINEAL THIRD EYE**
7. Tinnitus Natural Treatment
8. Vasculitis Natural Treatment
9. **What is eating you**
10. **ADHD and Autism Treatment**
11. Smoking Cessation and Addiction Treatment
12. **ADHD Scio Eductor EEG**
13. Adrenal Fatigue Hypo-Adrenia
14. Aging and Inflammation
15. Allergy Therapy for Cure
16. Alzheimer’s Disease and Natural Energetic Medicine
17. Asthma Asthmatic Bronchitis
18. Baldness can be cured by plucking Remaining Hair
19. Banish Man Boobs (gynecomastia) with no drugs or surgery
20. Blood Clotting Diseases Natural Treatment
22. Burning mouth and Tongue Syndrome
23. Burns Natural Treatment
24. **CES for Depression**
25. Case Study Hodgkin’s
26. **Cataracts Natural and Energetic Medicine treatments prevention**
27. Causes and Treatment of Cognitive Mental Decline
28. Chronic Fatigue –ME now SEID Energetic Natural Medicine Treatment
29. Demyelination Disorders MS ALS Multiple Sclerosis Leukodystrophies Guillian Barre
30. Depression is a meaningful part of Life
31. Diabetes Natural Cure and Treatments
32. Ear ache – What to do?
33. **Energetic Medicine for Graves Disease**
34. **Energetic Medicine Treatment of spirochete Disease like Lyme and Syphilis**
35. **Energetic Medicine and Retinitis Pigmentosa**
36. **Energetic Natural Medicine for Thymus disorders including Myasthenia**
37. **Energetic and Natural Medicine for Headache**
38. Enuresis and Natural Treatment
39. Erectile Dysfunction Peyronie’s Disease
40. Extensive Clinical Study or Eductor with Pain Depression Pregnancy and Health
41. Fibromyalgia
42. Fifth Disease Scio Treatment
43. Four Decades of the Trivector Review of the Trivector till 2009
44. How to detect and treat Alzheimer’s
45. **Helicobacter Pylori infection causes disease**
46. Herpes Natural Treatments
47. Homeopathic Treatment of Prostatitis
48. Hyper Pituitary Acromegaly and Hypo Pituitary natural treatment
49. Hyperthyroid Grave’s disease
50. Hypothyroidism Scio Eductor and Natural Treatments
51. **IJMSHNEM Super Human Abilities**
52. **Impetigo Bacterial Skin Rash**
53. IMUNE on Infertility
54. LUPUS
55. Leukemia Treatment and Prevention with Scio Tech
56. List of Phobias and Simple cures
57. Lose weight with Electro Stimulation
58. Lung Fungus Mycobacteria
59. Midwifery
60. Mycology of Fungal Infections
61. NAHINGA Aids Research
62. Natural Anti-Inflammatories
63. Natural Medicine cures for AIDS
64. Natural Medicine for Baldness
65. Natural Medicine for Prostate
66. Natural Medicine for Epilepsy
67. Natural Rx for Kidney Disease especially PKD
68. Natural Statins versus Synthetic Statin
69. Natural Treatments for Hydrosalpinx Blocked Fallopian Tubes
70. Natural Trigeminal Neuralgia Treatment
71. Natural Medicine for ACNE and SCIO Study
72. Natural Therapy for Cystic Fibrosis
73. Natural Treatment of Lung Disease COPD, Asthma Bronchitis SCIO EDUCTOR study
74. Naturopathic and Energetic Medicine for Menopause
75. Pain and Quantum Biology
76. PMS Evidence for Energetic Medicine Treatment
77. Parkinson’s- How to increase Dopamine with Scio Tech and Natural Med
78. Prepare for Ebola or any new Virus
79. Processed High Glycemic Sugar causes Disease
80. Prof Desire Dubounet and the Eductor Proves ESP
81. Prostate Health
82. QXCI and the long term Pathological Findings of the Camelford Toxicity Group
83. Rheumatoid Arthritis and Amoeba
84. Relieve Constipation Naturally
85. Scio Eductor Study of Voltammetric Stimulation of Blood testosterone
86. SOC Index and the Evidence for Lifestyle Medicine
87. Static Pain Natural Treatments
88. Seeing the Future with the Eductor analysis of experiments on anomalous anticipation of random future events
89. Shaping and Hormone Streaming
90. Sjögren’s Syndrome natural remedies
91. Stones in the Body
92. Natural Medicine Stroke Treatments
Learning Enhancements with the SCIO

Part of the following:
Large Scale Study of the Safety and Efficacy of the SCIO Device

Chief Editor:
Prof. M. Vlasic

Edited and Published By:
Mediserv, Sera Dovzi
Somlja Ulica

International Ethics, Lestane 58,
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Developed By:
The Centro Ricerche di Prof. William Nelson University of Venice + Padua, Italy

This study was performed in the field by practicing Bioklochowska technicians. Data was collected and the study supervised by the Ethics International Institutional Review Board of Romania. The data analysis and study presentation is done by The Centro Ricerche, University of Venice + Padua, Italy.


This article is translated into English and Hungarian from the original Romanian.

Abstract
The SCIO device is a Universal Electro-Physiological device used for stress reduction and patient treatment. Over 2,200 qualified Bioimaging technicians joined our Ethics Committee study to evaluate how stress reduction and learning enhancement of the SCIO device. The device has insignificant risk. There was a staff of medical doctors who designed and supervised the study.

Over 100,000 subjects gave informed consent and participated in the study. The study was conducted by professional biologists and data analysts. The results showed a significant improvement in stress reduction and learning enhancement of the SCIO device. The device has an insignificant risk. The study was designed and supervised by a team of medical doctors.

Clinical Study No. 3 - VARIOPE Changes in a SCIO Session

Correction of aberrant body electric profiles such as voltage, amperage, resistance impedance, proton + electron pressure.

Part of the following:
Large Scale Study of the Safety and Efficacy of the SCIO Device

Chief Editor:
Prof. M. Vlasic

Edited and Published By:
Mediserv, Sera Dovzi
Somlja Ulica

Consultant:
International Ethics, Lestane 58,
Crnica, Romania

Developed By:
The Centro Ricerche di Prof. William Nelson University of Venice + Padua, Italy

This study was performed in the field by practicing Bioklochowska technicians. Data was collected and the study supervised by the Ethics International Institutional Review Board of Romania. The data analysis and study presentation is done by The Centro Ricerche, University of Venice + Padua, Italy.


This article is translated into English and Hungarian from the original Romanian.

Abstract
This study demonstrates the safety and effectiveness of the SCIO device used in a large scale study. A large scale study of over 97,000 patients with over 122,000 patient visits reported their diseases. The SCIO measures global electrical measurements of the body. When there are abnormal measurements of the electro-physiological factors, the device allows feedback loops between the central nervous system (CNS) of the patient and the device. The device resolves these imbalances by modifying its electrical profile.
Title: Large Scale Study of the Voltammetric SCI

**Stimulation of Blood Estrogen**

Chief Editor: Dr. David Polen DC USA

The Centro Ricerche, University of Venice + Padova, Italy

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Developed By:
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**Title: Large Scale Study of the Voltammetric SCI**

**Stimulation of Blood Testosterone**

Chief Editor: Prof. William Nelson M.D. MUNE

The Centro Ricerche, University of Venice + Padova, Italy

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- Richard Atkinson M.C.S.P., Physical Therapist, West Yorkshire England

Abstract:
A global and momentous research project was developed for the last few years. The SCI device is a universal electro-physiological device used for stress reduction and patient treatment. Over 2,200 qualified biofeedback therapists joined our ethics committee study to evaluate how stress reduction using the SCI device could help a wide variety of diseases.

The device and thus the study have insignificant risk. There was a staff of medical doctors who designed and supervised the study.

Over 90,000 patients gave informed consent and participated in the study. The study would conclusively prove safety and efficacy of the SCI Device. With over 90% of these patients having multiple visits, there were over 275,000 patient visits. With a total record of the SCI patient information, therapy parameters and reactive data, no names of patients were recorded for confidentiality.

Two of the 2,200 plus therapists were given blank devices that were completely visually the same but were now functional. These two blind therapists were then given 25 patients each. This was to evaluate the double blind component of the placebo effect as compared to the device. Thus the study groups were a placebo group, a subgroup, and an attached therapy group.

This is just the first study in a long task of analysis in truly break down the data totally. This study verifies the safety and efficacy of the SCI device. There were no adverse effects seen in the placebo group, larger effects in the subgroup, and astounding effects in the real therapy group.

In this study 35 females between 37 and 69 were evaluated for their daily estrogen cycle. The results have shown that the SCI estrogen streaming therapy of 23 a day for 5 days raises their estrogen.

**SCI EDUCATOR LARGE SCALE RESEARCH**

2,200+ Therapists, 100,000+ Subjects, 300,000 Patient Visits
CERTIFICAT CONSTATATOR

emis în temeiul art. 12 alin. (1) lit.b) din Legea nr. 359/2004
privind simplificarea formalităților la înregistrare persoanelor fizice,
asiociții Familiale și persoanelor juridice, înregistrarea fiscală a acestora,
precum și la autorizarea funcționării persoanelor juridice,
cu modificările și completările ulterioare,
eliberat în baza declarației pe propria răspundere înregistrată sub
nr. 46073 din 30/11/2006

Firma: INTERNATIONAL ETHIC SRL
Sediul social: MUNICIPIUL ORADEA, Str. MOLDOVEI, Nr. 17, Bloc AN50, Ap. 20, Județul Bihor.
Cod unic de înregistrare 18830990 din data 06/07/2006.
Număr de ordine în registrul comerțului JG5/13992006 din data 06/07/2006.

Prenumele certificat constatator aștează că s-a înregistrat declarația pe propria răspundere conform căreia firma îndeplinește
condițiile de funcționare, specifice pentru fiecare autoritate publică, pentru activitățile declarate, încadrare în clasa CAEN:
7210 Consultanță în domeniul echipamentelor de calcul (hardware);
7230 Prelucrarea informatică a datelor;
7240 Activități legate de bazele de date;
8542 Alte forme de învățământ;
la sediul principal din MUNICIPIUL ORADEA, Str. LEBEDEI, Nr. 58/A, biroul nr.4, Ap. biroul 4, Județul Bihor
(CENTRU DE PREGĂTIRE I.M.U.N.E.).

Valabilitate: până la modificarea condițiilor de funcționare sau activităților pentru care s-a dat declarația pe propria
răspundere.

DIRECTOR,
CORNELIA LUCIA GLIGOR

Emis la data: 05/12/2006
Eliberat la data: 05/DEC/2006
Following are the links to a recognized peer reviewed medical journal registered with an ISSN number for over 20 years

IJMSHNEM journal #-005 1989 AAQBT the work from 1980-1997/
IJMSHNEM journal #-009 1991 Updated Volt-Ammetry/
IJMSHNEM journal #-012 1993 Updated Biofeedback, Chaos, Fractals, Etc/
IJMSHNEM journal #-016 1 Electro Physiological Reactivity, the Xrroid Effect, Vol. I/
IJMSHNEM journal #-025 1998 5 Electroacupuncture/
IJMSHNEM journal #-028 2001 1 Biofeedback/
IJMSHNEM journal #-034 2002 2 Electro Physiological Response Advanced, Volume 2/
IJMSHNEM journal #-048 2005 4 Basic Body Electric/
IJMSHNEM journal #-051 2006 3 Sport Medicine/
IJMSHNEM journal #-052 2006 4 Quantum Electro Dynamics and Cancer/
IJMSHNEM journal #-053 2006 5 The Future of Natural Medicine - Conference of IMUNE St. Marteen 2006/
IJMSHNEM journal #-054 2007 1 Electro-acupuncture Advanced/
IJMSHNEM journal #-055 2007 2 Proofs of the Power of the Mind/
IJMSHNEM journal #-056 2007 3 Review of the Trivector/
IJMSHNEM journal #-059 2008 1 2 3 4 The Large Scale Study of the Scio/
IJMSHNEM journal #-061 2008 Special Issue on Voltammetry/
IJMSHNEM journal #-073 2010 VARHOPE and Stress/
IJMSHNEM journal #-074 2010 VASO-VAGAL Reaction what you need to know to operate the SCIO/
IJMSHNEM journal #-084 2013 4 Quantum Entwinement as a Principal of Human Communication/
IJMSHNEM journal #-089 2013 United States of America Research/
IJMSHNEM journal #-090 2013 Romanian Study of the Stimulation of Sports Ability/
IJMSHNEM journal #-091 2014 Causes and treatment of Cognitive + Mental Decline/
IJMSHNEM journal #-092 2014 Healing Research EEG to stimulate Healing Energies of People/
IJMSHNEM journal #-093 2014 Journal supplement additional articles proving the Transcranial Eductor/
IJMSHNEM journal #-094 2014 PAIN and Quantum Biology/
IJMSHNEM journal #095 2014 Reprint from 2001 The Princeton Engineering Anomalies Research and full discussion of the Powers of the mind and random number/

http://www.downloads.imune.net/medicalbooks/Activating%20the%20Vagus%20Nerve.pdf

IJMSHNEM journal #096 2015 Bio-Electro-Magnetism
IJMSHNEM journal #097 2015 Bio-Electro Magnetism Cyber Megnatics/
IJMSHNEM journal #098 2015 GSRtDCs issue
IJMSHNEM journal #099 2015 GSRtDCs issue 1/
IJMSHNEM journal #101 2015 Super Human Abilities
IJMSHNEM journal #102 2015 TVEP issue/
IJMSHNEM journal #103 2015 VIBRATION MEDICINE Frequency Listing(1)
IJMSHNEM journal #105 2015 issue on TVEP(2)
IJMSHNEM journal #106 2016 Alzheimer/
IJMSHNEM journal #107 2016 Journal on Homeopathy/
IJMSHNEM journal #108 2016 Weight Loss/

International Medical University of Natural Ed Certified Medical Textbook Library
On-Line Access for you. Validation of Natural Med

Next is easy access links to the Certified Medical University Textbooks quoting research on the Eductor
Click to see Registration History

This massive body of research has gotten registrations all over the world: Europe, United States of America, China, Mexico, Canada. All of our research has been conducted according to the regulatory requirements, everything has been done to the letter of the law, and all of our paperwork submitted to the notified bodies has always followed the most recent standards and requirements. Staff in our office have had more than 500 hours of training in regulatory standards and procedures.

In 2008 our company obtained the United Kingdom Intertek CE certificate:
In 2009, the Korean registration was obtained:
In 2010 we obtained the Mexican registration:

In 2010 we obtained the Mexican registration:

In 2010 we obtained the Mexican registration:

In 2010 we obtained the Mexican registration:

In 2010 we obtained the Mexican registration:
Indicaciones de uso: El SCIO es un sistema auxiliar de biofeedback para pre-diagnóstico y terapias de regulación del estrés que mide los cambios en voltaje, amperaje y resistencia de un organismo al aplicar microcorrientes eléctricas.

Descripción: Es un sistema que actúa motivando el potencial evocado del organismo mediante microfrecuencias eléctricas.

Sus dimensiones son 201 x 80 x 175 mm y su peso aproximado es de 760 g.

El equipo está integrado por: 1 electrodos de cabeza, 1 electrodos de extremidades, 1 cable conector USB a USB, 1 caja de interfaz, DVD & CD, de software y 1 manual de uso, su alimentación eléctrica es a través del puerto USB conectado al ordenador.

Cuenta con una entrada nominal eléctrica de 4 a 5 voltas (dependiendo del ordenador) y una salida nominal máxima a través de electrodos de 4 a 10 miliamperios.

El electrodos de cabeza pesa 200 g y sus medidas son 1.75 m de cable y 760 x 51 mm de goma. Este electrodos entra en contacto con la piel al ser aplicado alrededor de la cabeza.

Los electrodos de extremidades pesan 180 g y miden 25 m de cable y 19 x 370 x 2 mm de goma, el electrodos entra en contacto directo con la piel al ser aplicado alrededor de las muñecas y tobillos.

Presentaciones: Empaque con un equipo.

Modelo SCIO

Publicidad dirigida a: Profesionales de la salud.

COF 017543
On 29 August 2012 SFDA has approved the SCIO to be marketed in the Chinese market after a successful review process of our application.
Our latest European CE Certificate was obtained with TÜV Rheinland InterCert Kft on 23 February 2012.
Certificate

The Certification Body of
TÜV Rheinland InterCert Kft.

hereby certifies that the company

Mandelay Kft.
ÁTI-SZIGET IPARI PARK 11. ép.
H - 2310 Szigetszentmiklós, Hungary

Site: Kálvária tér 2., H - 1089 Budapest, Hungary

has established and maintains a quality management system
for medical devices for the following scope:

Design/Development, manufacturing, distribution and
servicing of Universal Electrophysiological Biofeedback
System

Proof has been furnished that the requirements of


are fulfilled. The certification is subject to periodic surveillance.

Certificate Registration No.: OX 69241776 0001
Audit report No.: 28208466 004
This certificate is valid: from 2012-02-23 to 2015-02-22

2012-02-23
Date of issue

Balázs Bozsik
Certifier

TÜV Rheinland InterCert Kft. – H-1132 Budapest, Váci út 48/A-B
Tel.: (+36/1) 461-1100, Fax: (+36/1) 461-1199, e-mail: medical@hu.tuv.com, http://www.tuv.com/hun/
In the United States the Eductor/Educator have been successfully registered in 2014 with the FDA:

The EPFX measures the Electrophysiologic Reactivity intensity of the patient to many QQC trivector voltammetry patterns. These are patterns or reactions to Sarcodes, Nosodes, Allersodes, Isodes, Nutritional, Herbs, Imponderable and Classic Homeopathics. The reaction patterns or profiles can relate disturbances of the patient. Therapists can then be arranged to develop harmonic reactions, desensitizations, biological resonance or rectification processes. All of these are applied and managed through biofeedback application. Biofeedback is the operation that allows for the cybernetic loop of systemic feedback. The only indicated use of this device and all claims related to this device are under biofeedback. The loop of measured reaction and bio-varied resonance response allow for a true feedback for self corrective ElectrophysiologicaL therapy. Hence it is called the Electro Physiological Feedback Xrroid.

Excerpt from the 510k registration of 1989

DEPARTMENT OF HEALTH & HUMAN SERVICES

Food and Drug Administration
1200 Piccard Drive
Rockville, MD 20850

Re: K92114A
Electro-Physio-Feedback-Xrroid System

Eclision Corporation
Attn: Frank B Mauro
3936-A Niagara Street
Denver, Colorado 80207

Dated: Undated
Received: July 18, 1989
Regulatory Class: II
Next Australian Register of Therapeutic Goods Certificate 2015
South African Registration 2015

DEPARTMENT OF HEALTH

Directorate: Radiation Control
Private Bag X62
BELLVILLE
7535

Tel: (021) 9486162
Fax: (021) 9461589

web: https://sites.google.com/site/radiationcontroloch/

MEG-ANN 5762 SERVICES CC
PO Box 40015
QUEENSBURGH
4070

Enquiries: X-Ray devices: Ms N.P. de Kok
Other devices: Mr J.P. Uys
Mr G. G. Diederiks

Reference: 6008851
Date: 30 January 2015

Attention: Marianne Lillian Van Rooyen

- This updated document contains the licences for electromedical devices as well as the licence conditions that are currently valid, and replaces the document dated 11 August 2014 and all previous documents.

- Apart from the other licensing considerations, the licence for each individual model is issued on the strength of the fact that the intended purpose, as stated in the application form, is considered to be in agreement with the intended purpose of the device as reflected in the manufacturer’s labelling and instructions for use (i.e. documentation required in terms of the certification process according to EC Directive 90/422/EEC or 90/384/EEC, whichever is applicable).

- The licence for each model remains valid only while the EC compliance documentation is valid.

- The safety and performance of all the licensed models remain the responsibility of the licence holder.

- Inspections may be performed to ascertain whether the licence conditions are being adhered to.

Yours faithfully

[Signature]

DIRECTOR-GENERAL: HEALTH
TUV Reviewed + Accepted Clinical Research Evaluation of Medical Claims

Path to Evidence

PHASE 1
1985 to 89 basic research at the AAQET with 935 Subjects

Basic Research on
The Body Electric and
Stress leads to 1989 FDA Registration

Our Dedication
to Validate
the Claims

Randomized Controlled Studies
Case Control Studies
Cohort Studies
Cross Sectional Studies
Meta Analysis

PHASE 2
2004 to 2008 2,200 therapists study
100,000 subjects on 220 diseases

PHASE 3
2008 to 2016 research on Sport, Intellect,
Memory etc makes Meta Analysis work


Situational judgment
meta-analysis
validity

Collective
Systemic
Review of the Vast
Research on the
SCIO-Eductor
Technology

Medical Research
Validation of
the SCIO

http://indavideo.hu/video/Evidence_Meta_Analysis_of_the_Eductor_SCIO_Technology_with_music
meta abstracts
http://www.downloads.imune.net/medicalbooks/Medical%20Research%20Validation%20of%20the%20SCIO.pdf
Chinese Olympics

Helping us to get the Chinese registration was the research done on the 2008 Chinese Olympic Team. The Beijing 2008 Olympic project was unbelievably successful project where, over a three month
period, more than 200 Chinese Olympic athletes were tested and 60 athletes were worked with closely in therapeutic trials in over 1200 sessions with the SCIO/EPFX System. All of this was under the supervision of Doctor Li Guo Ping, head of Sports Medicine China), as well as 20 doctors and nurses through the Federal Hospital Sports.

The people involved in the Chinese Olympics project were (QX China), Victor Ke, George Fang, Jeff Sutton, Dr. Jiao Li Ping and Yin Lin.

Voltammetry Stimulation of Testosterone in Gold medal Athletes in the China Olympics

Taken from Adam Mandel’s lecture in 2011 Budapest at the congress on SCIO about work done at the 2008 Olympics

Adam Mandel was sent to China to work on the Chinese Athletes with the SCIO in 2007 till 2008. He presented his work for the first time in the 2011 International conference on the SCIO held in Budapest. Titled “The Greatest Story Never Told” he showed the audience incredible awe inspiring details of wondrous results with the SCIO. This is a brief summary of his 3 part lecture now available from IMUNE.

Adam talked about testosterone boosting by voltammetry streaming therapy of the SCIO. The SCIO is a

Project Nahinga Continuation update

There have been a host of different other studies conducted on this technology, including Project Nahinga, in Mozambique and South Africa.
Nahinga – Immuno Compromised Protocol

**IMMUNO COMPROMISED PROTOCOL**

First Avoid All White Processed Sugar
This means all dextrose sugar products including candy, cola, doughnuts, etc.
You can have all of the levulose fructose products you want,
this means any fruit of fruit sugar

Second Avoid all Foods Boiled or Fried in Oil.
Use at least three tablespoons a day of uncooked natural cold processed oil such as
olive oil, safflower, sunflower, soybean oil.
Have some five servings a day of fresh and raw fruits or vegetables.

Third Reduce Stress and Enjoy Life
Do at least fifteen minutes twice a day of quiet meditation using affirmations, and
imagery of your immune system working.

Exercise for twenty minutes at least four times a week
work to a sweat and breathe deeply, use the thymus tap

Take the Hemo-A twice a day with other supplements
Hemo-A has Yeast, Santa, Phytobaica, Chinese Curcumin Sativa, Triglycerin, renning sperm
and other sacodes of Thymus adenoids tonsils and appendix
use 500 mg Vitamin C, 15, mg Zing, 4 pills of Oxygen Stimulator at bed
use Golden Seal, Aloe Vera, Lentil, Mustard, Tumeric, Curry, Paprika
Sesame seeds, and use Samboura

**AVOID ANTIBIOTICS**
use Probiotics Actimel, Activa, etc Twice a day

Electrical Reactivity as a Prescreen of HIV Infection Patients

**ELECTRICAL REACTIVITY AS A PRESCREEN OF HIV INFECTION PATIENTS**


**ABSTRACT**

Twenty-two ambulatory AIDS patients in Budapest were studied for xrode electrical reactivity readings. The electrical reactivity patterns and reactive substances that were in the highest faction of reactivity. In other words, those reactants that were statistically significant are compared in the groups of the AIDS patients taking the AZT as well as the AIDS patients that were treated with homeopathic and nutritional items. The purpose of the study was to analysis similarities and consistencies in their electrical reactivity patterns over the course of four measurements. This took place during the 4th, 5th, 6th and 7th month of 1994. During these months there was also a homeopathic and nutritional Intervention done on several of these patients to see the effect on blood chemistry profiles denoting aids risks and the homeopathic and nutritional intervention are described in the article known as the comparative results.

Immunological and Electrophysiological Reactivity of Patients with HIV Infection
IMMUNOLOGICAL AND ELECTROPHYSIOLOGICAL REACTIVITY OF PATIENTS WITH HIV INFECTION

By: Nagy K., Nelson W., Barabas E., Balazs E., Varkonyi V., Horvath A.
National Institute of Dermato-Venereology, Budapest, Hungary 1994

ABSTRACT

The diagnostic and prognostic value of electrophysiological reactivity patterns of HIV infected subjects were compared to complex immunological and virological laboratory markers.

Electrical responsiveness of 22 asymptomatic HIV infected patients were monitored monthly for a 4 month period by Quanta Med 4000, a sensitive multichannel diagnostic biofeedback machine, capable of measuring slight fluctuation of patients' brain waves and skin resistance, whilst the patients are sequentially exposed to a battery of numerous homeopathic nosodes and Isodes. This process known as the Xrodeo process.

Immunological and virological analysis included determination of CD cell count, HIV, HCV and CMV antibodies, HbsAg, and serum Beta 2-microglobulin (B2M) and Neopterin levels.

In the study - 4000 substances (items) were tested to determine which of these the patient had the most reaction to. The reactivity scores were then analyzed statistically. As a result a profile of electro-reaction is suggested, which considered characteristic of HIV disease in contrast to that of the normal subjects.

After initial evaluation a treatment protocol was designed. Half of the patients received a fatty acid blend and homeopathic medicaments throughout the test. Subjects were instructed to use the products daily and compliance was evaluated in the monthly interview. Reevaluation of electrophysiological reactivity and immunological tests were repeated every month.

H2M level was found decreased in 88 percent of those who received homeopathic treatment and in 50 percent it was found < 3 mg/L, compared to 27 percent of those untreated. Antibody level to CM was also found decreased in consequence of treatment. No changes, however could be detected in CD count and HbsAg and HBC antibody level.

The electrophysiological reactivity test provided information, which suggest that it can be used as a pre-diagnostic method, which might complete laboratory analysis. Complex homeopathy and individual nosode treatment shows some positive intervention.
ANTIBIOTICS AS A PRIMARY CO-FACTOR IN AIDS PROGRESSION


"If a Man sees a Wrong and does not Correct it, He is NOT a Man"

ABSTRACT

The world has now recognized the demise of antibiotics; iatrogenic damage, resistant strains, immunosuppression and dependency have now challenged the core of one of the prides of modern medicine. The vast marketing of antibiotics has left medicine with a severe crisis. Reductionistic research and philosophy has been used for financial reward of the chemical companies. These antibiotics have been shown to have a wide variety of deleterious side effects, including effects on the bowel flora. We also theorize about how this disruption of the bowel flora, could be a contributing cofactor to the AIDS epidemic.

The populations with the greatest antibiotic use are the highest risk for development of AIDS. A balanced bowel flora could be essential in defense against the virus propagation into the deadly disease. The antibiotics might then increase the progression of risk in the disease. This hypothesis, because of its non-reductionistic complexity is difficult to challenge in a single study. Funding of such a study would also be extremely difficult, in light of the challenge to synthetic chemistry. This brief article is but an introduction to the concept. For further information please refer to the collection of studies in the Journal of the Medical Science of Homeopathy, special issue on AIDS and vituses.
Dr. Kofi Ghartey of Project Nahinga in Ghana

We have seen six patient data files from Ghana where patients who had HIV in a Blood test were cured and the tests have come back negative for HIV after the Nahinga protocol.

In Budapest, Africa, and elsewhere we have seen several reports of cures and viral irradiation. This begs for further study and confirmation of results. But the research from project Nahinga seems interesting at least. We are continuing the search for more funds to do better studies and better controls, but till then we still seek to study and treat patients with this future orientated medicine.
Dr Amanda Velloen - Budapest Conference 2009
Update on Project Nahinga

Project Nahinga (bantu for the Angel that comes from the heavens to help us, the bantu name for Desire) has been spreading and many extreme success stories have shown fantastic results.

PROGRESS
- Results in PHASE 1 -
- Up to 100% feeling better
- Average 65% improving in Critical Measures of HI V progression
- No side effects reported
- Symptom free
- No secondary infections
- Improved quality of life

Critical Measures
- Subjects tested positive for HI Virus
- CD4 count
- Elisa test if available
- Viral Load test

Schedule for 2009
- Father John Mugaga in Kwa-Zulu Natal 1st Satellite
- October 2009 – AIDS Children’s homes
- November 2009 – Brazzaville Congo, DRC, Angola, Nigeria
- November 2009 - Kenya, Zimbabwe, Mozambique
- December 2009 – Lesotho, Swaziland

After Desire has donated over 2 million dollars of equipment for our research we have shown safety and efficacy and there appears to be a natural cure for this horrible disease on the future horizon
FEKI – freiburger etik kommission

In 2009 we received approval from FEKI, Freiburger Etik Kommission, for a body of research that was conducted in Italy, Germany, Romania, Switzerland, France and United States of America.

FEKI was the ethics committee for the following clinical studies:

- VARHOPE Improvements in a Clinical Setting
- VARHOPE Large scale study – Correction of aberrant body electric profiles such as voltage, amperage, resistance impedance, proton + electron pressure
- Verbal report of stress reduction - A double-blind placebo-controlled study of the application of Eclosion EPFX/SCIO therapy for stress reduction
- A double-blind placebo-controlled study of the application of the SCIO Universal Electrophysiological Biofeedback System for statistical evaluation of the SCIO’s ability to increase Body Wellness after one 45-minute session
- Double Blind Study of Sport Performance with the SCIO device versus Placebo control 2013 USA
- Trauma Sport Pain Electro Healing With SCIO-2013 USA
- MCES and Addiction Control a Dbl Blind Clinical Study -2013 USA
- SCIO’s Effect on Body Osmosis 2013 -USA
- Stimulating Eye Hand Coordination With SCIOVARHOPE Update 2013
- SCIO Effects on Oxidation/Oxygenation 2013
- TVEP reactivity scores to Allersode compounds measured 2013 USA
- TVEP reactivity scores to compounds measured update 2013 USA
- Voltammetric Sarcode Hormone Streaming of Testosterone Update 2013 USA
- VARHOPE and EPR Validation Of the SCIO technology -2013 USA

TUV inspected and approved our research, including our FEKI approval obtained in 2009, and gave us our CE mark in 2010. We were also given medical device approval at the time.

Need for Clinical Investigations

The following guidance documents were referenced regarding the Clinical Investigations Route:


Conduct of Clinical Investigations

The clinical study, in addition to being conducted under the above guidance, was also conducted in following these guidance documents and regulations:

a) MDD 93/42/EEC Annex X Clinical Evaluation
b) NB-MED/2.7/Rec3 Evaluation of Clinical Data
c) ISO 14155 Clinical Investigations of Medical Devices for Human Subjects
d) ICH 6 Guidance for Industry: Good Clinical Practice: Consolidated Guideline
e) ICH 8 Guidance for Industry: General Considerations for Clinical Trials
ICH 9 Guidance for Industry: Statistical Principles for Clinical Trials

Requirements

i. Identification of Relevant Documents:
   a. Copy of the letter of “no objection” (Approval Granted letter) and opinions/comments from the Ethics Committee. Note: This was the first submission to the Ethics Committee and this protocol has never been rejected.
   b. Copy of the signed and dated Final Report.
   c. Copy of letter of no objection from a European Competent Authority. The Romanian Competent Authority have received the application and have permitted the study.

ii. Information to be checked
   a. The determination of the device as non-significant risk (NSR) was approved on 16 November 2009 by the Freiburger Ethics Committee (feki).
   b. Clinical Investigation Plan (CIP): the CIP used for the Clinical Study is the same that was submitted for approvals. Evidence of this is the Case Notes stored in the Office of the Regulatory Manager at the Sponsor are the same as the Clinical Study Protocol which was approved by the Freiburger Ethics Committee (feki).
      i. The number of patients entered were: 151 TOTAL from the following testing sites (Seattle, Washington (USA), Paris (France), Speyer (Germany), Timisoara (Romania).

   ii. Objectives of Investigation (in particular which Essential Requirements are being addressed):
      ✓ (1) The devices must be designed and manufactured in such a way that, when used under the conditions and for the purposes intended, they will not compromise the clinical condition or the safety of patients, or the safety and health of users or, where applicable, other persons, provided that any risks which may be associated with their use constitute acceptable risks when weighed against the benefits to the patient and are compatible with a high level of protection of health and safety;
      ✓ (2) The solutions adopted by the manufacturer for the design and construction of the devices must conform to safety principles, taking account of the generally acknowledged state of the art.
      In selecting the most appropriate solutions, the manufacturer must apply the following principles in the following order:
      ▪ eliminate or reduce risks as far as possible (inherently safe design and construction),
      ▪ where appropriate take adequate protection measures including alarms if necessary, in relation to risks that cannot be eliminated,
      ▪ inform users of the residual risks due to any shortcomings of the protection measures adopted,
      ✓ (3) The devices must achieve the performances intended by the manufacturer and be designed, manufactured and packaged in such a
way that they are suitable for one or more of the functions referred to in Article 1 (2)(a), as specified by the manufacturer.

✓ (6) Any undesirable side effect must constitute an acceptable risk when weighed against the performances intended.

✓ (10.1) Devices with a measuring function must be designed and manufactured in such a way as to provide sufficient accuracy and stability within appropriate limits of accuracy and taking account of the intended purpose of the device. The limits of accuracy must be indicated by the manufacturer.

- The Duration of the Investigation, per the approved protocol, was one 45-minute Treatment with immediate follow-up within 3 hours after treatment. This was completed on all 150 patients and documented in the Case Notes.
- The end points in terms of diagnostic tools and patient assessment are stated in the approved Protocol under “Expected Results.”
- The Inclusion and Exclusion criteria, as stated in the approved protocol, were all met so that the total 150 subjects all met the inclusion criteria.

- All parameters were followed as set up in the approved, original CIP.
- There were no changes to the parameters as set up in the approved, original CIP.
- One testing site (Seattle, Washington, USA) was outside the EU and the population, as shown in the inclusion/exclusion criteria, is equivalent to those for which the device will be used within the EU and other parts of the world were regulatory status is sought.
Dear Mr. Lloyd,

please find enclosed the original version of the following documents for the a. m. study without an invoice:

- Grants Approval (x2)
- Amendment #1 (x1)

The feci wish the study great success and thank you for the confidence you have shown us.

Yours sincerely,

Karin-A. Graf
CERTIFICATE

Study Title: A Double-Blind Placebo Controlled Study of the application of the SCIO Universal Electrophysiological Biofeedback System for Statistical evaluation of the SCIO’s ability to increase Body Wellness after one 45-minute session
Clinical Study Protocol
Final Version
Maitreya Kft.
Revision 2.0 19 August 2009 CT-103-01

Study Code: CT-103-01
fcii Code: 09/2120
Sponsor: Maitreya Kft.

Date of meeting: August 24, 2009 grants conditional approval
Place of meeting: Mozartstrasse 21, 79104 Freiburg, Germany

The proposed clinical study was reviewed on August 24th, 2009 with conditional approval. The freiburg ethics commission international (fcii) has completed a careful review of the study protocol, the informed consent and other submitted documentation (see Review Request Form Documents page 2), in particular from ethical and legal points of view and with impartial expertise. The regulations of the German Medical Device Law (MPG) § 20 Abs. 8 (MPG § 23) are reviewed and the bylaw about protection against damages caused by X-rays or radioactive material/ ionizing rays (§ 28g RöV and § 92 StRöSchV) have also been reviewed. (The sum insured stated in the documents fulfills the demands of risk assessment according to MPG).

The fcii requests the submission of an interim report after one year (should the study last longer than one year) and a brief final report upon completion of the study.

In your letter (E-Mail) dated November 6th, 2009 you substantiate that all conditions have been fulfilled.

With regard to proposed clinical study, the fcii hereby

grants approval

Prof. Hans-Peter Graf , Md PhD Freiburg, November 16, 2009
CERTIFICATE

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☑ grants approval

Prof. Hans-Peter Graf, Md PhD
Freiburg, November 16, 2009
CERTIFICATE
to an
Amendment 1
of the clinical study

Study Title: A Double-Blind Placebo Controlled Study of the application of the SCIO Universal Electrophysiological Biofeedback System for Statistical evaluation of the SCIO’s ability to increase Body Wellness after one 45-minute session Clinical Study Protocol Final Version Maitreyka Kft. Revision 2.0 19 August 2009 CT-103-01

Study Code: CT-103-01

feci Code: 09/2120

Sponsor: Maitreyka Kft.

Date of meeting: August 24, 2009 grants conditional approval November 16, 2009 grants approval November 16, 2009 amendment 1

Place of meeting: Mozartstrasse 21, 79104 Freiburg, Germany

The freiburg ethics commission international (feCi) has reviewed the protocol amendment # 1 – Clinical Study Protocol Revision 2.2, dated October, 2009 (Patient Informed Consent Form English included) – under consideration of the relevant protocol and accompanying documentation according to ethical, legal and medical-scientific points of view, and with impartial expertise. The regulations of the German Medical Device Law (MPG) § 20 Abs. 8 (MPG § 23) and the bylaw about protection against damages caused by X-rays or radioactive material/ionizing rays (§ 28g RöV and § 92 StrlSchV) have also been reviewed. (The sum insured stated in the documents fulfils the demands of risk assessment according to MPG).

With regard to proposed amendment, the feci hereby

grants approval

Prof. Hans-Peter Graf MD PhD

Freiburg, November 16, 2009
VARHOPE (Voltage, Amperage, Resistance, Oxidation, Hydration, Proton and Electron pressure, the body electric's vital signs) Clinical Journal

The VARHOPE Clinical Journal (ISSN 2041-4293) published a series of studies sponsored by Maitreya Kft./Eclosion Ltd Kft. on various topics concerning charging the human battery, charged stability, stress reduction with the SCIO System.

VARHOPE Improvements in a Clinical Setting

The first study we are going to discuss has taken place in Toronto, Canada, where the investigator, Jonathan Sargent, a SCIO therapist, did an evaluation of 100 patients of before and after VARHOPE readings. The results (published in this journal) showed significant changes after a two week therapy session.

This study shows three cases and an overview of one hundred cases.

VARHOPE Improvements in a Clinical Setting

By Jonathan Sargent

ABSTRACT: We are made up of atoms that are mostly electrons and protons. The outer electrons of any atom or molecule never touch. The outer electrons of any atom or molecule never touch another set of electrons. The entire interaction is through electro-magnetic-static, quantic, or other interactive fields.

There is Electrical energy in the human body. The most simple factors of anything electrical are the volts amps and resistance. This makes up Ohms law of electronics, where Volts = Amps times Resistance. This is a correlation not an exact law. Oscillations of the volts and amps give us the frequency of a current. Fluctuations of these calculations can be used in virtual or mathematical ways to calculate other biological factors. There are norms of the body electric variables relative to age and lifestyle.

Thus the key Bio-electric factors of Volts Amps Resistance Hydration Oxidation and Proton versus Electron charge stability are measurable with the SCIO. Then device then using a cybernetic biofeedback loop can help to stabilize the VARHOPE of a client.

In this study we assay the VARHOPE scores of several clients. In all cases there is a significant improvement in the bio-electric after a simple SCIO therapist lead session.

This study shows three cases and an overview of one hundred cases.

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Case Study 1
Charts for Case Study 1
Case Study 2
Charts for Case Study 2

Case Study 3

Charts for Case Study 3

Discussion

the patient

A student coming through the beginner program brought this client to me. The student was living two hours north of the patient and was not comfortable working with his situation. The patient was diagnosed with HIV (Human Immunodeficiency Virus) in late 2004. When the patient came to me, it was mid March and he was exhibiting and feeling many symptoms.

The symptoms were ranging from skin lesions, skin discolouration, yellow eyes, and chronic fatigue. The patient was taking an AIDS (Aquired Immune Deficiency Syndrome) prescription drug, prescribed by his doctor, as well as seeing a nutritionist and having colonic sessions, 1-2 times a week. He was not feeling any results and felt that he was on his death bed. He was therefore in a fragile emotional state, which needed to be balanced immediately. As we were talking and unraveling the deep emotional core issues, the patient made a decision that his mother’s lack of love for him was the strongest negative resonance. He believed that the tension was created due to his choice in sexual orientation and belief.

On February, 23rd 2006 the patient’s HIV count was 327,021 copies/mL.

(data collected by his doctor).

Demographic

Male

Born – 1960

Age - 46

Diagnosed with HIV and AIDS

Prescription drug for AIDS

Lifestyle – altered and healthy

Water consumption – fair

Employment – High school teacher

Case

With this particular client it was a life and death situation right from the beginning. The patient’s digestive system was compromised due to years of unhealthy eating, without the use of digestive enzymes. On our first session together, we helped each other to balance the emotional issues and help to build the immune system. Through stress management and organ stimulation, we worked with the lymphatic system, adenoids, appendix, tonsils, thymus, liver and spleen. These organs were gently stimulated through biofeedback therapy and the spiritual healing was balanced. NLP (Neuro Linguistic Programming) techniques were used to work with guilt and self esteem issues and in addition, I added Auto zaps for AIDS and “feel good”. I told the patient that if I were in his situation I
would take a “detox bath”, containing baking soda, Epsom salts, and hydrogen peroxide 3%, to help detox the skin. After a bath I would take a shower and rub my skin with coconut oil, to help bring back the moisture and oils to my skin.

He saw the benefit this would have for him. the patient mentioned that he was getting colonics done 1-2 times a week, but I noticed that he was neglecting to replace to vital digestive enzymes (probiotics) that his body was desperately searching for, this information was concluded from the extreme mal absorption issues appearing on the device. From the beginning of his first session it was clear there was an issue with blood, environmental, immune, inflammation, liver, and fungus.

It was determined at that time that we would have a 2 hour session once a week and if there was a concern of any kind, he could call and be seen right away. What I noticed after the first session with the patient is that his skin colour returned and he had a look in his eye of confidence, as if he found his path to wellness.

Continuing

the patient returned once a week for 7 months. Over this time we worked through the layers of stressors on the body and helped supply the proper nutrients needed. I had the patient bring several types of supplements from his nutritionist and checked the resonance according to his meridian flow and reaction. It was not long after the first session that the patient’s skin began to clear up and he could return to work. His digestion and bowel movements improved greatly. This was a great relief to him, for the goal seemed to be in sight! As we proceeded through the process his body was now ready to start detoxing. At first I told him in this situation, I would do a lymphatic cleanse for 2 weeks, and then begin a liver cleanse that was supplied from a Hulda Clark website. Over 100 stones were released within his stool and lots of internal bowel congestion, that was blocking his main detox outlet.

Throughout this entire time I noticed that his VARHOPE scores were improving greatly per session, with the odd slip. The more we worked together the longer the varhopes would stay in tune. This whole process was very exciting for both of us, since at the outset of our meeting he had been sent home by his doctor to die. This alone was a traumatic event. It was a coaching and confidence building conversation, which was critical throughout the entire process.

Since I am not a doctor, I was strongly recommending that he see his doctor for check ups and routine blood checks. During this time his doctor would “forget to get his blood test done” and would not disclose his information. We couldent figure it out. Then the patient became angered by this and demanded that another blood test be taken and reported to him immediately. This alone was a feat of emotional confidence that was not the same the patient I first met, frail and scared of the unknown and what the future held. Two weeks passed and the patient arrives in my office with a giant smile on his face. I ask him what he was so happy about. He handed me his blood test, not only from the recent blood test but also the “lost” blood results.

It read:

HIV count –

327,021 copies/mL, T-cell count 30 February, 2006
200,000 copies/mL, T-cell count 120 May, 2006
70,000 copies/mL, T-cell count 449 July, 2006
This was a very exciting time for both of us since the marker for HIV activity is 50,000.

It meant that we have almost reached our goal! We then continued throughout the next couple months with more biofeedback sessions.

**Review**

In reviewing this case I feel very confident in the direction of the therapy sessions. Other than the minor set backs, which is natural in the healing process. The body needs time to heal and to detox as well. Over the 7 months that I was working with the patient, I saw a great improvement in his spiritual, physical and emotional state. This was a great joy to see that we can help with such a fatal dis-ease.

I watch as his confidence and feeling of self worth improved. This was a very good thing, since now he was able to enjoy life to the fullest. I am looking forward to seeing the patient again in the future and see if there is any decline in overall state.

My advice to all new practitioners coming into the field of biofeedback is that we must listen to the body and not the “standard protocols”. In my experience it is relevant that balancing the person instead of the dis-ease. This alone causes for less therapy sessions and more productive results. We must understand that every individual is different and special in there own way and it would be unfair to treat every person the same as the last.

Our job and duty as biofeedback therapist is to assist the body in balancing stressors, in turn making it easier for the body to heal its self. We make suggestions of lifestyle changes and coach them through their emotional issues, so that they can make their own decisions. I find it much easier to work with clients when you have them create their own conclusions and help them to learn about themselves on an even deeper level. We need to educate, so that when our clients leave us, they are able to maintain a state of health and not fall back into their previous lifestyle that brought them to us.

March 22, 2006
March 25, 2006
May 10, 2006

Before

May 15, 2006

Before

Chart
May 17, 2006

May 29, 2006
July 20, 2006

July 24, 2006
July 28, 2006

August 8, 2006
the patient

This particular client was sent by a past client that found very good results working with me and the system. The patient was diagnosed with lung cancer at the age of 74.

He had decided that chemo therapy and radiation were not the answer for him. Over the course of five months we only saw the patient five times. This is not a lot of sessions, but this was the only time that the patient could fit into his very busy schedule.

the patient was very hard to communicate with due to his small understanding of the English language. He was always accompanied by his daughter, who translated everything in to his native tongue. (Italian)

Demographics

Born – 1931
Male
Diagnosed with lung cancer
Six different Prescription drugs
Water Consumption – fair
Employment – Retired

Case

As I looked into the patient’s case, it was prevalent that virus, fungus and parasite activity was high within his body. One of the first things I noticed with the patient besides the large amount of pathogens present, was his Ph levels were acidic. Making it a perfect home for all pathogens to mutate and duplicate. These pathogens were highly concentrated within the lungs and bowels of the client. This is also where he was diagnosed with cancer. Was it possible the parasites were mistaken for tumors? Very possible, I then decided to work with this concern because cancer was not in high levels, compared to pathogen activity.

the patient’s organs were also highly under stress due to his age and lack of previous care to his body. Degeneration was present, and he had a high reactance speed. His high reactance speed was telling me that his body did not recognize the pathogens as a threat; another good sign there was a possible wrong diagnoses.

Continuing

Over the five months that I saw the patient I worked with what his body needed for that day, balancing all stresses one by one. In turn this would allow his body to heal itself much quicker, and understand what was happening. The sessions were also mostly centered around Rife therapy and zapping for pathogens.
Since there were a lot of emotional issues coming to the surface, it was clear that there were some things that need to be balanced within the NLP program as well. It was very exciting to see as the weeks and months pass, the patient responding very well and balanced on all levels.

Near the end of month five the patient went missing. I didn’t hear from him for at least three (3) to four (4) weeks. It was making me worried, I decided to call his daughter and see what was happening. I was hoping for the best.

The answer was, “we didn’t tell you?” my father went to get some radiation done and before the treatment they did another MRI scan. The “cancer” was gone! Gone!

**Review**

To sum it all up, the patient was pleased with the sessions he received. It was amazing to see the traumatic things that were told to this particular client from his medical doctor without the proper research being gathered. Maybe if the medical world would be more accepting of biofeedback we could put an end to this “medicate or operate” system.

With a few lifestyle changes and a coaching process to deal with unspoken emotions, the results are amazing a significant. This technology of biofeedback is truly a leap towards a more productive medical system.

A few weeks after seeing the patient last, I decided to call his daughter. She had a tear in her eye and a frog in her throat. Her father went to the hospital for radiation. Before they perform the treatment it is standard protocol to administer another MRI, to find… THE TUMOR WAS GONE! How exciting is that! I was very impressed. Another person helped from biofeedback.

November 2\textsuperscript{nd}, 2005

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January 19th, 2006

March 3rd, 2006
The patient

The patient was recommended to me by his brother. His brother found us on the internet. The patient had been diagnosed with Parkinson disease. He had experienced troubles with gambling and money management due to the side effects from the medications that he was currently taking and he had “the shakes”.

When I first met the patient, he was in a very negative state. He could not hold still. Kicking and hand twitches were common. He was on three (3) different medications and was emotionally stressed. the patient is a retired hardware store owner, and had work very hard with several highly toxic chemicals on a regular basis.

Demographics

the patient
Male
Born – 1942
Age - 65
Diagnosed with Parkinson’s
Prescription drug for Parkinson’s
Lifestyle – fragile
Water consumption – fair
Employment – retired hardware store owner

Case

The patient had been diagnosed with Parkinson’s when I first met him. Sent to me by his brother, the patient did not know what to expect on his first visit. His thoughts were scattered and his emotions were high.

I did my best to guide the patient through the initial process in explaining what biofeedback was and how it works. the patient seemed to catch on quickly, and he was ready to try it out.

There was a high amount of heavy metals present, which told me that the nerves would be damaged and the myelin sheath would be gone for the most part. Chemical toxins were also high with in his blood. On the first session I always like to go through what the body is trying to say, nutrition, spinal, NLP, nelson report, all charts and patient super conscious reduction. This gives me a good idea of what is causing the most stress within the body and what the body is ready to deal with first.

Once I had my information it was very easy to find the path to wellness and the thread to dis-ease. With the patient it was all about the toxins running everywhere in his body. The homotoxicology screen was filled with red and endless supplies of toxins were present.
Continuing

With this information I asked the patient how often his bowels movements were and if there was any pain. Once I had that information it was evident that his bowels needed to be detoxed before any toxins were released.

Most sessions were centered on detoxing the body, repairing the nerves and bringing back mental clarity, although every session was different then the last. After our first session with the patient, he held out his hands. Which were not shaking! He then said “I haven’t been able to do this in 20 years”! I was amazed.

With more talking and coaching it was obvious the 30 years of working with toxic materials was the cause of his diagnosis of Parkinson’s. And the side effects of his medications were the cause of his gambling addiction. Working with his doctor, the patient was able to come completely off his medications and the nutrition profile allowed the patient to have a nutritious lifestyle. Increasing the amount of water in his diet and using a cilantro pesto for heavy metal detox was essential to the patient regaining health and reversing the aging process.

Review

In conclusion I would like to say that over 6 months and 11 sessions that the patient had we reached our goal! the patient was able to return back to the things he loved most. His sense of humor was back and he was able to play golf again with his brother. It was amazing to see the patient start as an individual sent home by his doctor to be a vegetable, into a strong willed, funny, and loving husband and brother. the patient was truly blessed to have come in contact with biofeedback and to receive the results he did.

the patient is currently living out his retirement as he should, playing golf, laughing lots and loving life even more. His self-confidence has returned and his gambling has stopped.

November 3rd, 2005
November 7\textsuperscript{th}, 2005

November 11\textsuperscript{th}, 2005
Before
January 3rd, 2006

January 10th, 2005
Discussion

The SCIO measures global electrical measures of the body. When there are abnormal measures of the electro-physiological factors, the device allows a feedback loop between the Central Nervous System (CNS) of the patient and the device. We have seen hundreds of clients in our Clinic. this report shows 3 cases intimately. But in the other cases there is almost always some improvement in the VARHOPE indices are even one therapy. Of our clients visits, some 50 plus % show very low electro-physiological factors, some all below 30% normal. The cybernetic electro-physiological feedback loop is used to help the client reduce stress and thus improve the electro-physiological factors. There was improvement in over 95% of the electro-physiological measures at the end of the session (post test) versus the pre test. The average improvement in electro-physiological VARHOPE factors is 5% per session. These clients report stress reduction and improved well being as well.

Norms of patient voltage, amperage, resistance, capacitance, inductance, reactance, impedance, and proton electron balance, have all been established.

When we intake air and breathe we oxidate. This produces a shift up and down of our amperage. Oxidation is measured as the average shift in amperage ove a measured set of time. Hydration is the shift of voltage over a set of time.

Proton pressure versus electron pressure is the measure of which is more plentiful in the body. Excess protons means an acid condition, excess electrons means alkainity. By measuring the electrical imbalance from the various electrodes of the SCIO we can measure the proton versus electron pressure. A balance form is reported at 70 where there are equal numbers of electrons and protons. Below 70 is excess protons, above excess electrons. A variation of the ph scale.

The SCIO software will allow the Central Nervous System (CNS) of the patient to guide to stabilize electrical and vibrational divergence in your body. This is the cybernetic loop or biofeedback component of the system.

Important Note (This study had the following contributors:

Institution : International Medical University NE
Ethics Committee Int IRB : Ethics International
Medical staff has supervised the overal study
Peer review committee for the International Journal of the Medical Science of Homeopathy has reviewed and accepted this study for publication.

The purpose of this study was to assay the safety and efficacy of a visit to a trained SCIO therapist, exacting statistics are not assayed beyond the simple questions of reported success. Thus this study points to further scientific studies of more refined statistics. )

VARHOPE Large scale study – Correction of aberrant body electric profiles such as voltage, amperage, resistance impedance, proton + electron pressure

This study demonstrates the safety and effective qualities of the SCIO device used in a large scale study. A large scale study of over 97,000 patients with over 320,000 patient visits reported their diseases. The SCIO measures global electrical measures of the body. When there are abnormal measures of the electro-physiological factors, the device allows a feedback loop between the
Central Nervous System (CNS) of the patient and the device. Of the patient visits listed some 50 plus % (55,921) showed very low electro-physiological factors, below 30% normal. The cybernetic electro-physiological feedback loop was then used to help the patient reduce stress and thus improve the electro-physiological factors. There was improvement in over 95% of the electro-physiological measures at the end of the session (post-test) versus the pre-test. The average improvement in electro-physiological VARHOPE factors is 5%. These patients reported stress reduction and improved well-being as well.

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This study was performed in the field by practicing Biofeedback technicians. Data was collected and the study supervised by the Ethics International Institutional Review Board of Romania. The Data analysis and study presentation is done By the The Centro Ricerche, University of Venice + Padova, Italy © Ethics International, 2007.

Correction of aberrant body electric profiles such as voltage, amperage, resistance impedance, proton + electron pressure,

Part of the Following:

Large Scale Study of the Safety and Efficacy of the SCIO Device

Chief Editor:

Prof N. Vilmos

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and the study supervised by the Ethics International Institutional Review Board of Romania. The Data analysis and study presentation is done By the The Centro Ricerche, University of Venice + Padova, Italy © Ethics International, 2007. (This article is translated into English and Hungarian from the original Romanian)

Abstract

This study demonstrates the safety and effective qualities of the SCIO device used in a large scale study. A large scale study of over 97,000 patients with over 320,000 patient visits reported their diseases. The SCIO measures global electrical measures of the body. When there are abnormal measures of the electro-physiological factors, the device allows a feedback loop between the Central Nervous System (CNS) of the patient and the device. Of the patient visits listed some 50 plus % showed very low electro-physiological factors, below 30% normal. The cybernetic electro physiological feedback loop was then used to help the patient reduce stress and thus improve the electro-physiological factors. There was improvement in over 95% of the electro-physiological measures at the end of the session (post test) versus the pre test. The average improvement in electro-physiological VARHOPE factors is 5%. These patients reported stress reduction and improved well being as well.

Introduction

Overview

This Large scale research was designed to produce a extensive study of people with a wide variety of diseases to see who gets or feels better while using the SCIO for stress reduction and patient monitoring. The SCIO is a biofeedback Universal Electro-Physiological Medical apparatus. The device is registered in Europe, America, Canada, S Africa, Australia, S. America, Mexico and elsewhere.

The traditional software is fully registered. Some additional functions where determined by the manufacturer to be worthy of evaluation. Thus a study was necessary to determine safety and efficacy. (As a result of these studies these additional functions are now registered within the EC) An European ethics committee was officially registered and governmental permission attained to do the insignificant risk study. Qualified registered and or licensed Biofeedback therapists where enlisted to perform the study. Therapists were enrolled from all over the world including N. America, Europe, Africa, Australia, Asia, and S. America. They were trained in the aspects of the study and how to attain informed consent and transmit the results to the ethics committee or IRB (Institutional Review Board).

2,569 therapists enlisted in the study. There were 98,760 patients. 69% had more than one visit. 43% had over two visits. There were over 320,000 patient visits recorded. The therapists were trained and supervised by medical staff. They were to perform the SCIO therapy and analysis. They were to report any medical suspected or confirmed diagnosis. Therapists personnel are not to diagnose outside of the realm of their scope of practice. Then the therapist is to inquire on any reported changes during the meeting and on follow-ups any measured variations. It must be pointed out that the Therapists were free to do any additional therapies they wish such as homeopathy, nutrition, exercise, etc. Therapists were told to not recommend synthetic drugs.
Thus the evaluation was not reduced to just the device but to the total effect of seeing a SCIO therapist.

- Part 1. The emphasis was on substantiating safety followed by efficacy of the SCIO.
- Part 2. Proving the efficacy of the SCIO on normalizing the electro-physiological factors

**Methods and Materials:**

**SCIO Device:**

The SCIO is an evoked potential Universal Electro-Physiological Medical device that measures how a person reacts to items. It is designed to measure electro-physiological factors. Biofeedback is used for stress reduction and muscular re-education.

The SCIO measure global electrical measures of the body. Norms of measurement can only be assayed on an individual piece of equipment. The nature of the electrodes, the size or surface area of measurement, age and personal demographics, the reading mechanism etc. The SCIO has been registered in many countries and has been legally sold for over two decades. After thousands of patients tested with this device in the late 1980s, norms of the electrical parameters have been set. Norms of patient voltage, amperage, resistance, capacitance, inductance, reactance, impedance, and proton electron balance, have all been established.

When we intake air and breathe we oxidate. This produces a shift up and down of our amperage.

Oxidation is measured as the average shift in amperage over a measured set of time. Hydration is the shift of voltage over a set of time.

Proton pressure versus electron pressure is the measure of which is more plentiful in the body. Excess protons means an acid condition, excess electrons means alkainity. By measuring the electrical imbalance from the various electrodes of the SCIO we can measure the proton versus electron pressure. A balance form is reported at 70 where there are equal numbers of electrons and protons. Below 70 is excess protons, above excess electrons. A variation of the ph scale. The SCIO software will allow the Central Nervous System (CNS) of the patient to guide to stabilize electrical and vibrational divergence in your body. This is the cybernetic loop or biofeedback component of the system. For complete functional details and pictures, see appendix.

**Basic Software Design:** The SCIO software is designed for electro-physiological connection to the patient to allow electro-physiological and rectification of subtle aberrance of the body electric. The feedback loop is established by measuring the electro-physiological factors, feeding them back to the CNS and re-measuring the changes, feeding them back to the CNS, and on and on till a satisfied result is attained for the day.

**VARHOPE Scores**

As previously described, there are norms set for the Body electro-physiological measures. Since these measure are relative to age, sex, and other demographics, a percentage of the norm is used on the report. There is a degree of inaccuracy as well. norms are reported loosely as above 80%. In this series of tests when the percentages of the global VARHOPE are all below 30%, the device patient records were used to report the post test or after therapy results.

**SOC Index (Library Function stressor questionnaire):**

The SCIO interview opens with a behavioral medicine interview. This is called the SOC Index. Named after the work of Samuel Hahneman the father of homeopathy, he said that the body heals itself
with it’s innate knowledge. But the patient can suppress or obstruct the healing process with some behavior. Hahneman said that the worst way to interfere with the healing natural process was allopathy or synthetic drugs. These upset the natural healing process by unnatural intervention and regulation disturbance. Other ways to Suppress or Obstruct the Cure are smoking, mercury amalgams, stress, lack of water, exercise and many others. This behavioral survey then gives an index of SOC.

The scores relate to the risk of Suppression and Obstruction to the natural Cure. The higher the scores the more the Suppression and or Obstruction. The scores of 100 or lower are ideal. A copy of the SOC index questions appear in the appendix.

Study Technicians:

The study technicians were educated and supervised by medical officers. The study technicians were to execute the SCIO therapy and analysis. All were trained to the standards of the International Medical University of Natural Education. Therapists from all over the world including N. America, Europe, Africa, Australia, Asia, S. America and elsewhere were enlisted to perform the study according to the 1951 Helsinki study ethics regulations, since modified several times.

They were to chronicle any medical suspected or confirmed diagnosis. Therapists personnel are not to diagnose outside of the realm of their scope of practice. Then the study technician is to inquire on any disclosed observations during the test and on follow-ups report any measured changes.

To test the device as subspace against the placebo effect, two of the 2,500+ therapists were given placebo SCIO devices that were totally outwardly the same but were not functional. These two blind therapists were then assigned 35 patients each (only 63 showed). This was to assess the double blind factor of the placebo effect as compared to the device. Thus the studied groups were

A. Placebo group, and B SCIO group.

Cross placebo group manipulation was used to further evaluate the effect.

Important Questions : these are the key questions of the study

1. Define Diseases or Patient Concerns
2. Percentage of Improvement in Symptoms
3. Percentage of Improvement in Feeling Better
4. Percentage of Improvement Measured
5. Percentage of Improvement in Stress Reduction
6. Percentage of Improvement in SOC Behavior
7. What Measured+How (relevant measures to the patient’s health situation)
8. If Patient worsened please describe in detail involving SOC_

After the patient visit is was complete the data was e-mailed to the Ethics Committee or IRB for storage and then analysis. This maneuver minimized the risk of data loss or tampering. Case studies were reported separately in the disease analysis.

Results:
Before we review the direct disease improvement profiles, we need to review the overall results.

The first most basic of question in the results is the basic feedback of the generic patient conditions.

1. Percentage of Improvement in Symptoms
2. Percentage of Improvement in Feeling Better
3. Percentage of Improvement Measured
4. Percentage of Improvement in Stress Reduction
5. Percentage of Improvement in SOC Behavior

The SOC index gives us great insight to this study. Each disease has a different cut off where the ability of the SCIO to help was compromised. As a general index scores of 200 + where much less successful.

The electro-physiological aberant group total number of patients was 55,921

SCIO Harness Patients, 55,921 Patients

OVERALL ASSESSMENT

A. Placebo Group- 63 cases with a Dbl Blind System and no Treatment

There were no cases of patients who reported a negative Improvement.

There were

- 19 cases reporting no improvement of Symptoms, 30% of group
- 12 cases reporting no improvement in feeling better, 19% of group
- 13 cases reporting no improvement in stress reduction 20% of group
- 12%--- Percentage of Improvement in Symptoms
- 15%--- Percentage of Improvement in Feeling Better
- 2%---. Percentage of Improvement Measured
- 12%-- Percentage of Improvement in Stress Reduction
- 3%----Percentage of Improvement in SOC Behavior

B. SCIO Treatment 163,870 patient visits

There were 658 cases of patients who reported a negative Improvement.

There were

- 512 cases reporting no improvement of Symptoms, .003% of group
- 759 cases reporting no improvement in feeling better, .004% of group
- 460 cases reporting no improvement in stress reduction .002% of group
- 65% -- Percentage of Improvement in Symptoms
- 56% -- Percentage of Improvement in Feeling Better
• 24%---Percentage of Improvement Measured
• 53%--Percentage of Improvement in Stress Reduction
• 20% ---Percentage of Improvement in SOC Behavior

There was an overall 43% average improvement in the VARHOPE score from the therapy on each visit. There was an additional improvement on pursuing visits.

**Pre and Post SCIO Therapy Electro-Physiological percentages**

<table>
<thead>
<tr>
<th>SCIO Treatment 163,870 patient visits</th>
<th>Pre</th>
<th>Post</th>
</tr>
</thead>
<tbody>
<tr>
<td>Voltage</td>
<td>25</td>
<td>33</td>
</tr>
<tr>
<td>Amperage</td>
<td>21</td>
<td>27</td>
</tr>
<tr>
<td>Resistance</td>
<td>10</td>
<td>15</td>
</tr>
<tr>
<td>Hydration</td>
<td>27</td>
<td>36</td>
</tr>
<tr>
<td>Oxidation</td>
<td>28</td>
<td>35</td>
</tr>
</tbody>
</table>

SCIO Treatment 100,834 patient visits

Below

| Proton vs Electron                  | 50  | 57   |

SCIO Treatment 63,036 patient visits

above

| Proton vs Electron                  | 79  | 77   |

**Discussion**

The results show significant improvement in symptoms and feeling better. The Collective results show a dramatic benefit to the SCIO therapist visit.

The Electro-physiological factors were slightly improved during the SCIO therapy. This is reported by most of our therapists. The Cybernetic Feedback loop of the SCIO to the CNS seems to be helpful in stabilizing the VARHOPE.

There is much more to the body electric than resistance. It is impossible to measure a frequency with a resistance device such as the Listen, Voll, Vega or other simple resistance devices. This makes for a very complicated fractal system that can be difficult to analyse. With a computer a vast amount of electrical data can be collected and analyzed. This can then allow for the beginning of a true energetic medicine. Below is an abbreviated list of electrical variables and their corresponding components that our SCIO can analyze in the short space of minutes in a clinical setting. The time of ionic exchange is approximately one hundredths of a sec. Thus it would be impossible for a person who wants to test a patient with a manual device.

*articles in Promorphus
As we pointed out in the Promorpheus, electricity as an electrical entity travels in the direction of, for example, your right thumb. Then for conduction of the electron, there is a magnetic field produced at 90°, and a static field will be produced at another 90°. This electromagnetic and electrostatic combination and its effect on conductance and from conductance is the basis for understanding electrical phenomena.

The factors of the electrolyte in the body greatly affect the electrical nature of the body. The amount of minerals, liquids, oxygen, amino acids, fatty acids and others effect the nature of the electrolyte. So our total energetic medicine (beyond simple resistance variables) can offer us great insights into many factors of health. Since so much of energetic medicine is fixated in one channel resistance point probetechniques it is time for a quantum leap in the technology. In this article we will outline some basic aspects of energetic medicine for electroencephalographs electro cardiology and energetic medicine.

This article will outline the electron and its action. The photon link is outlined in the Promorpheus. French physicist Coulomb laid out a law, which states: "The force of attraction or repulsion between two charged bodies is directly proportional to the product of the charges and inversely proportional to the square of the distance between them."

Thus the force can be allowed in the following equation

\[ F \approx \frac{Q_1 \times Q_2}{D^2} \]

The inverse square law is a dictum of four-dimensional physics. Our ten-dimensional model questions its pervasiveness.

Here Q represents the force of the charges, D is the distance, and F is the force in dynes. A coulomb of charge, C, is nearly 3 times 10^9 esu. The strength of an electrical field will have the equation

\[ E \approx \frac{9 \times 10^9 \times q}{R^2} \]

This is called the electrical potential. The potential at a point is equal to the work needed to bring one coulomb charge to the point from an infinite distance away. Biology will need to monitor this effect very closely.

An electric potential is thus work per unit of charge. Kinetic energy, which is equivalent to work, is measured in a relationship of force to distance. A gram that is moved at one centimeter per second of velocity is an erg. A kilogram that is moved at one meter per second is known as a joule.

When we have a joule per coulomb, this is known as a volt. One volt equals one joule divided by one coulomb. The volt is often a measure of potential energy. It is the difference between two points, between positive and negative charge; thus a six-volt battery with a potential difference of 6 joules or coulombs that can flow from one terminal to the other. Potential difference, thus, is an integral measurement of profound importance in biology and medicine.

If the surface of an item has a charge that is stored as potential energy, the ratio of charge to potential is called the capacitance of the body. The basic unit of capacitance is known as the farad, which is one coulomb per volt. If one coulomb of charge added to a body gives it potential of one volt, it has the capacitance of one farad. In a capacitor current is proportional to the rate of change of voltage.

Thus capacitance can be measured as a fluctuation in voltage (DV) over a qualitative time.

\[ 1 \text{ Farad} \approx \frac{1 \text{ Coulomb}}{1 \text{ Volt}} \quad \text{BOLD} \quad \text{Capacitance} \approx \frac{\text{dV}}{\text{dT}} \approx \text{Amps} \]
The farad is a very large unit, measuring a lot of potential. Often in electronics we use micro-farads, or even pico-farads; a micro-farad being 10^-6 farads and a pico-farad being 10^-12 farads. By having two sheets of a high conductor, such as metal, with an insulating material between them, we can produce a condenser or capacitor. In biology cellular forces will invoke pico-farads. Organismic forces must relate to and control micro-farads.

The capacitance of the capacitor is the amount of the electrical charge on its plate divided by the potential difference between its plates. This depends on several factors, such as the area of the plates. If the plates are made larger, greater charge can be put on them. The thickness of the insulating layer is important. The closer the plates are to one another, the greater the amount of charge that is held. It is the strength of the electric fields of the electric plates as they are brought closer together. In biology organs, cells, organ systems, and organisms must store charge to deal with metabolism and growth.

The material between the plates will have an influence on the capacitor. These insulators, or non-conductors between the plates, are also known as dielectrics. Biology is filled with membranes that act as storage entities. We have only to review neuronal axon transfer to see biocapacitance at work.

The dielectric constant of an insulating material is a relationship between the effect of the material and that of a vacuum between the plates. The dielectric constant of water is 80; the dielectric constant of air is 1.001, as compared to a vacuum. The dielectric constant of rubber is 2.5.

Water has such an enormous dielectric constant because the water molecule is already polarized, even if it is not in an electric field. One end of the water molecule is positive and the other negative, because of the dipole magnetic effect. Biology uses this concept of water to store and use energy.

The molecules can now rotate easily in the liquid state, and in response to the electric forces on them can readily produce strong layers of induced charge on its surfaces. Capacitance action is of extreme importance to biology.

When we move one coulomb of charge per second, this is known as an ampere. An amp is movement or quantity of charge. Movement of charge, amps, is the most important criteria of biology. This correlates to life force and indolamine production.

\[ 1 \text{ Amp} = 1 \text{ Coulomb per second} \]

\[ 1 \text{ Volts} = \frac{\text{Inductance}}{\frac{\text{d Amps}}{\text{d Time}}} \]

\[ \text{Amps} = \frac{\text{Capacitance}}{\frac{\text{d Volts}}{\text{d Time}}} \]

Dr. Ohm, a German physicist, found that electric current in a conductor is directly proportional to the potential difference between its ends. Thus he generated Ohm's law, finding that the resistance of one ohm is generated in a conductor if the potential difference of one volt between its ends will cause a current of one ampere to flow through it. Thus we have generated and found

Ohm's law, which is

\[ \text{Ampere} = \frac{\text{Volts}}{\text{Ohms}} \]

\[ \text{or Volts} = \text{Amps} \times \text{Resistance} \]

Ohm's law is not strictly adhered to in electrolytes, discharge of gasses, and semiconductors; nor is it followed perfectly applicable to biology, for there are many different factors that can affect it. Changing potentials over time causes an instability in Ohm's law for biology. But in knowing an
electrical system we must know the amperage, the voltage, and the resistance in order to be able to
calculate variables more accurately. Ohm’s law, when involved in quantic systems, is not precise, but
still shows the tendencies of electromotive force. For biology Ohm’s law offers an invaluable systemic
measuring system for easy bio force analysis.

Now let us look at some of the basic components and relationships of magnetic fields.

When strongly polarized molecules align, they induce stronger and stronger magnetic poles. An
electric current flowing through a wire will also generate a magnetic field of 90 (right-hand rule).

The strength of the magnetic field created by a current is directly proportional to the strength of the
current and inversely proportional to the distance from the wire. The formula for this will show that
Magnetic~ Fields~ left ( {Amp} over {2 pi d} right )

Thus a magnetic field strength can be measured in units of amperes per meter. Inductance is the
factor measured for biological significance. Magnetic and paramagnetic forces can have strong
implications in the long- and short-range forces of biology (see PROMORPHEUS).

A magnet near a stationary electric charge will not have an effect on it. If there is movement, then
they have a natural influence on each other. Biology will need to be dynamic, and move constantly to
use magnetic properties. The force of this influence is at right angles to both the velocity of the
charge and the direction of the field. Stagnation is a magnet’s enemy.

The magnitude of this force is

Force = Charge in Coulombs x Velocity in meters per secondand Magnetic Force of Amperes per
meter x the Permeability Factor through which the Magnetic Field permeates.

This permeability factor times the magnetic factor, which is amperes per meter, is known as the
magnetic flux density, or the magnetic induction, and is expressed in Webers per square meter. In an
inductor the voltage is proportional to the rate of change in the current.

Inductance~ TIMES~ {d~ Amps} over {d~ Time}~ =~ Volts #phantom x #

1~ Henry =~ 1~ Volt/~ Second/Amp

These permeability factors are rated between that of the material and that of permeability of a
vacuum. Materials that are high-ratio (that increase the flux density) are called ferro-magnetic; such
as iron, cobalt, and nickel. Substances that are close to the ratio of 1, or other substances (which are
very near to the relationship of the vacuum) are para-magnetic, and will contribute weakly, such as
aluminum. There are substances like bismuth that are actually detrimental to the magnetic field.
These are called diamagnetic, and their ratio is actually less than 1. Items which are non-magnetic
will have no influence, and thus have a ratio of 1. Bismuth will have a place in biology, and is used in
several homeopathics for energetic stability. Magnetic induction can be measured by changes in
amperage over a qualitative analysis, such as the QXCI* machine test.

This might be used to infer magnetic interaction, and thus, involvement of geopathic stress.

Thus we have outlined the concept of magnetic, static, and conductive forces, which are used to our
understanding of the electrical nature of our homeopathic pharmaceuticals. By measuring the
inductance, the dielectric constant and the conductance relationship, we can find an electrical profile
for these various substances. This makes up an electrical fingerprint that allows us to calculate and
plot its electrical nature. The trivector analysis is born. The long-range implications on energetic medicine are profound.

By charting the resistance, inductance, and dielectric constant of various homeopathic items we can get a trivector analysis of their electromagnetic fields. This trivector analysis gives us three vectors, which we will be able to apply to a three-dimensional space. Thus a variety of homeopathics have been analyzed for their trivector analysis. The dimension of time gives us a four-dimensional relation that with some superb mathematics we can extrapolate the six virtual dimensions using a trinary logic system.**

Here we can see some of the effects that sarcodes, nosodes, allersodes and classic herbals have in their relationship to each other. This trivector analysis gives us a quality control factor for the electric field of a homeopathic item. In analyzing patients we can analyze serum in blood or personal field in a similar fashion. We can measure body pH from urine, blood, breath, etc., as well as redox capacity and body fluid resistance. Skin resistance readings can be taken at several points and easily averaged. Body voltage can be easily measured by dissimilar metals creating potential across the electrolyte capacity of the body, just as in a battery. Most proficient instruments choose to use silver and zinc (zinc because of its equi-potential for giving or receiving electrons, silver because of its great medicine history). Amperage is a correlate of voltage and resistance by placing similar metals in contact with the body (two silver probes contacting the frontal eminences). We can get an amperage reading. For our device Carbon electrodes where chosen for their ability to accept and donate electrons. Capacitance is measured by changes in voltage during a scheduled interview. Inductance can be calculated through changes in amperage over the same interview.

Resonant frequencies of the body can be calculated from the equation

\[ \text{Resonance} \sim \text{Freq.} \sim 10^6 \text{ over sqrt} \left(1^2 + (\text{CAP})^2 + (\text{IND})^2\right) \]

From these readings we can now calculate a true metabolism chart to define a patient's overall health and energetic well-being. .

The preliminary work has shown that where patients have valleys, or dips, in their fields, homeopathic peaks will be helpful. Work on this is just starting; more work, funding and time will be needed before we can find out if this is a viable technique for quality control and/or for homeopathic utilization. Now, with the help of the computer, matching remedies is high-tech and easy.

Another factor that we can use with this trivector analysis is that once we know the first three vectors, and the vector of time, we might be able to extrapolate the other six virtual dimensions.

If we know the four factors of conductance, capacitance, inductance and time, we might be able to extrapolate other dimensional effects from this four-dimensional type of field.

Biology needs to not only look into quantum physics but also needs to embrace an energetic philosophy as well. This seems complicated at first, but is easy with today's tools. Applying our right-hand rule and Ohm's Law to energetic medicine represents a dramatic quantum leap in energetic medicine which is significant to the field.

Bibliography

Books

Verbal report of stress reduction - A double-blind placebo-controlled study of the application of Eclosion EPFX/SCIO therapy for stress reduction clinical study protocol

As we reported in the introduction, many people overvalue and overrate the verbal mind. They might even think that it is all. At any rate a second test was designed to evaluate stress reduction over a three month period by using a verbal assay of the stress of the patient. The verbal mind is only aware of very little of the body's activities and is easily confused and prone to suggestion.

A study of 240 patients, over a three-month period started with a verbal questionnaire using the Beck's Stress inventory. A placebo group was designed and the test sites of Speyer, Germany and 5 test sites in Italy were chosen. Under medical doctor supervision the intervention was a series of treatments with the SCIO or the placebo device.

This study had two main flaws. 1. There was no on-site study supervisor to properly determine that the placebo device was fully disabled from the active software. Later analysis determines that there was confusion and perhaps no placebo was used. 2. The study used only verbal subjective measures and was not properly designed to measure electro-physiological or medical data.
The study results showed no difference from placebo group to treatment group. There was 62.5 percent improvement in both groups. This is a significant improvement over no intervention. Most people do not improve such after 3 months of stress therapy. So there were results. But since there was no guarantee of an established placebo group a fourth study was conducted.

A double-blind placebo-controlled study of the application of Eclosion EPFX/SCIO therapy for stress reduction clinical study protocol

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Giuseppe Mauger  
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Dr. Rainer Mutschler  
Germany  
Kathrin Sollner,  
Germany  
2008 Ethics International

PURPOSE OF STUDY

The purpose of this clinical study is to determine the efficacy of the ECLOSION Electro Physiological Feedback Xrroid (EPFX)/Scientific Consciousness Interface Operations System (SCIO) device, manufactured by ECLOSION KFT (the Company), in stress reduction by introducing low-level electromagnetic frequencies into an individual’s body through electrodes attached to the person’s wrists, ankles, and forehead to balance or harmonize and return to normal the optimal frequencies at which the body’s cells and organs should resonate. This enables the body to strengthen, heal and expel the pathogens that propagate stress and its associated ‘unwellness’, consequently reducing stress and improving general health and function.

EXPECTED RESULTS

Following completion of the treatment phase with the ECLOSION EPFX/SCIO, it is anticipated that the subjects in the test group, relative to subjects in the control group, will show, where applicable:

• a reduction in systolic and/or diastolic blood pressure reading at rest.

• decreased resting heart rate (fewer beats per minute)

• a decreased score on the Perceived Stress Scale (PSS), implying a reduced level of overall stress.

• decreased scores on one or more of the six mood factors of the Profile of Mood States (POMS)
Standard questionnaire, implying improved mood states.

• decreased scores on the State and/or Trait Anxiety scales on the Spielberger’s State-Trait Anxiety Inventory (STAI), implying reduced anxiety levels and/or improved reactions to anxiety.

• a decreased score on the Beck Depression Inventory-II® (BDI-II®) implying reduced levels of depression.

• some degree of satisfaction with the overall study outcome.

• maintenance in improved outcome measure ratings at the one-month post-treatment phase measurement time point.

For subjects in the control group, it is expected that there will likely be some improvement in measured variables. That is, subjects in the control group will likely report some of the positive changes listed above for test group subjects. However, on average, any positive change in post-treatment measures for control subjects is expected to occur to a significantly lesser degree than for subjects in the test group.

This study will be a double-blind, placebo-controlled, randomized clinical trial designed to demonstrate safety and effectiveness of the Eclosion EPFX/SCIO.

**TREATMENT GROUPS**

There will be two subject groups in this clinical study, with as close as possible to an equal number of subjects assigned to each of the two groups, as follows:

**Test group:**

Subjects in the test group will receive the actual study treatment with an active, operational harness.

**Control group:**

Subjects in the control group will receive a ‘fake’ study treatment with a placebo harness that does not contain any active electrodes.

Apart from the distinction of whether or not the subject receives the study treatment with the true or placebo harness attached to the Eclosion EPFX/SCIO device, all subjects will adhere to all phases of the entire protocol design.

**BLINDING**

This clinical study will be a double-blind design, such that neither the subject nor the investigator will be aware of to which group a subject has been assigned (test or placebo) until after the clinical study is complete.

Subjects will be randomly assigned to either Group A or to Group B, by the independent study Monitor. Subjects assigned to Group A will be treated with the EPFX/SCIO device A using Harness A and subjects assigned to Group B will be treated with the EPFX/SCIO device B using Harness B. Only the study Sponsor will know which label (‘A’ or ‘B’) corresponds to the actual (test) device and harness and which label corresponds to the sham (placebo) device and harness until the study is complete. The Sponsor will ensure this information is stored and maintained confidentially at the Sponsor’s work site. This knowledge will not be shared with the investigators, subjects, or study Monitor until the final subject data file of the study has been completed and submitted for analysis.
The sham (placebo) equipment will be designed to have the same external physical appearance as the actual equipment. The difference is that the placebo harness will not be equipped internally with functional electrodes and the programming for the placebo device will output only blank matrices. Neither the actual (test) nor the sham (placebo) harness produces any detectable noise, heat, light or other sensation output, so this also won’t be a distinguishing factor for subjects or the investigator between the test and placebo devices.

STUDY PROCEDURE - STUDY QUALIFICATION EVALUATION

SIGNING OF INFORMED CONSENT FORM

The investigator will start by presenting and reviewing in detail the items in the informed consent form with the individual and answer any questions he or she may have. To proceed further in the study, the individual must willingly sign the informed consent form at this time.

INCLUSION/EXCLUSION CRITERIA EVALUATION

After voluntarily signing the informed consent form, the subject will undergo the study qualification inclusion/exclusion criteria evaluation as follows.

INCLUSION CRITERIA

To be considered eligible for participation in this clinical study, a subject must satisfy each of the following “Inclusive Conditions” criteria.

Population: Individuals with Elevated Levels of Perceived Stress

Individuals in this study will be males and females who present with elevated levels of perceived stress as indicated by a total score of 25 or greater on the Perceived Stress Scale.

The Perceived Stress Scale (PSS) is a global measure of perceived stress that assesses the degree to which situations in an individual’s life are appraised as stressful. The subject is asked to indicate how often he or she felt or thought a certain way regarding 14 items, following a 5-point Likert scale from 0 to 4, as follows: 0=never, 1=almost never, 2=sometimes, 3=fairly often, 4=very often.

The PSS total score is obtained by reversing the scores on seven positive items and then summing across all 14 items, for a possible total of 56.

The PSS was designed for use with samples with at least a junior high school education. The items are easy to understand and the response alternatives are simple to grasp. The questions are general in nature such that they are relatively free of content specific to any sub-population group.

Validation data for the 14-item PSS was collected from three samples: two groups of college students and one group of individuals enrolled in a smoking-cessation program. Mean scores on the PSS complete samples ranged from 23.18 to 25.0. There was no statistically significant difference in mean PSS score between males and females, and age was found to be unrelated to PSS in all three samples.

Statistical evaluations found the PSS to have adequate internal and test-retest reliability and to be correlated in the expected manner with a range of self-report and behavioral criteria.

Additional information, including the complete PSS tool, can be found in Appendix C of this clinical study protocol. This includes the original article evaluating the scales, as follows: “Cohen, S.,

• Able and willing to maintain regular and consistent diet, exercise and lifestyle regimens throughout the study.

• Able and willing to maintain current medication regimes throughout the study.

• Able and willing to abstain from partaking in treatments – conventional or alternative (such as hypnotherapy, acupuncture, massage therapy, etc.) - or over-the-counter or prescription medications, including herbal remedies, designed to reduce stress throughout the study, other than the EPFX/SCIO treatment that is part of this study.

• Between 18 and 65 years of age.

• Male or female.

• Females on adequate birth control or not of child-bearing years.

EXCLUSION CRITERIA

A subject will be considered ineligible for participation in this clinical study if he or she satisfies any one or more of the following exclusive conditions criteria.

• Total score of less than 25 on the Perceived Stress Scale.

• Stage 2 Hypertension (elevated blood pressure), defined by a systolic blood pressure level of 160 mmHg (millimeters of mercury) or higher OR a diastolic blood pressure level of 100 mmHg or higher, measured using a sphygmomanometer and averaged across three seated (resting) blood pressure readings taken at 10-minute intervals. The first measurement will be recorded after the subject has been at rest (seated) for about 10 minutes. The source for the Stage 2 Hypertension criteria is the Seventh Report of the Joint National Committee on Detection, Evaluation, and Treatment of High Blood Pressure, American Heart Association Subjects taking antihypertensive (blood pressure lowering) drugs.

• Tachycardia, Bradycardia or Irregular Resting Heart Rate, defined as follows:

• Tachycardia: rapid or increased resting heart rate of greater than 100 beats per minute.

• Bradycardia: abnormally slow resting heart rate of less than 60 beats per minute.

• Irregular Resting Heart Rate: Irregular pattern of beats wherein beats are consistently missed across a 60-second period.

Resting Heart Rate - the number of times the heart beats per minute - will be measured at the wrist (radial artery), using the manual palpation method to feel the pulse - the rhythmic expansion and contraction (or throbbing) of an artery as blood is forced through it by the regular contractions of the heart. It is a measure of how hard the heart is working.

Heart rate through measurement of the pulse at the wrist will occur as follows:

1. The palm side of the subject’s right hand is faced upwards.

2. The investigator places his or her index and middle fingers on the wrist, approximately ½-1 inch below the base of the hand.
3. The investigator presses his or her fingers down in the groove between the middle tendons and the outside bone until a throbbing sensation - the radial pulse – is felt.

4. The investigator counts the number of beats that occur in 60 seconds, using a watch with a second hand or digital second counter for accuracy.

Resting Heart Rate will be taken after the subject has been seated for 10 minutes. The subject’s final recorded pre-treatment heart rate will be the average of three consecutive measurements, each taken about 5 minutes apart.

- Generalized Obesity, defined by a Body Mass Index (BMI) of 30 kg/m² or greater, according to the World Health Organization (WHO) and Center for Disease Control (CDC) criteria.
- Significant major stressful life events in the past 3 months likely to impact not only emotional but also physical health and wellness, defined by a score of 200 or greater on the Life Events Questionnaire (LEQ). The LEQ is contained in Appendix D.
- Significant major stressful life events known or anticipated to occur during the course of the study (i.e. the upcoming 3 months), defined by a score of 200 or greater on the Life Events Questionnaire, answered for known upcoming events such as a wedding, retirement, home move, etc.
- Type 1 diabetes.
- Any known heart condition(s), such as cardiac arrhythmias, congestive heart failure disease, myocardial infarction.
- Prior cardiac surgeries such as cardiac bypass, heart transplant surgery, pacemakers.
- Seizure disorder or family history of seizure disorder.
- Serious medical illness or condition: cancer; HIV, anorexia/bulimia.
- Serious head trauma
- Pregnant, breast feeding, or planning pregnancy prior to the end of study participation.
- Serious mental health illness such as dementia or schizophrenia; psychiatric hospitalization in past two years.
- Excessive use of any illicit drug or alcohol on a regular basis.
- Infection or wound or any other external trauma in the areas to which the electrode bands of the EPFX device are to be attached.
- Developmental disability or cognitive impairment that would make it difficult for the subject to partake in the clinical study, including adequate comprehension of the informed consent form and ability to record the necessary measurements.
- Involvement in litigation and/or a worker’s compensation claim and/or receiving disability benefits because of a stress-related or involved condition.
- Participation in a clinical study or other type of research in the past 30 days

TREATMENT PROTOCOL ADMINISTRATION PROCEDURE
PRE-TREATMENT PHASE
The purpose of the pre-treatment phase is to record baseline measures against which post-treatment changes will be assessed, and to record demographic subject variables.

The following measures will be recorded during the pre-treatment administration phase:

**Physiological Measures**

1. **Blood Pressure:** Systolic and diastolic blood pressure will be measured in millimeters of mercury (mm Hg) using a sphygmomanometer. If the pre-treatment phase occurs on the same day as the study qualification evaluation phase, then blood pressure does not need to be measured again.

If the pre-treatment phase occurs on a different day to the study qualification evaluation phase, then the subject’s blood pressure will be measured again at this time, three consecutive seated measurements, each ten minutes apart (as during the study qualification evaluation phase). Also as during the study qualification evaluation phase criteria, if the subject’s three-measurement blood pressure average falls into the category of Stage 2 elevated high blood pressure (defined by a systolic blood pressure level of 160 mmHg or higher OR diastolic blood pressure of 100 mmHg or higher), then the subject shall be disqualified from further participation in the study at this time.

Else, the subject’s blood pressure reading will be classified as follows:

<table>
<thead>
<tr>
<th>Category</th>
<th>Systolic (mm Hg)</th>
<th>Diastolic (mm Hg)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Normal</td>
<td>less than 120</td>
<td>and</td>
</tr>
<tr>
<td></td>
<td></td>
<td>less than 80</td>
</tr>
<tr>
<td>Prehypertension</td>
<td>120–139</td>
<td>or</td>
</tr>
<tr>
<td></td>
<td>80–89</td>
<td></td>
</tr>
<tr>
<td>Hypertension</td>
<td>140–159</td>
<td>or</td>
</tr>
<tr>
<td>Stage 1</td>
<td>90–99</td>
<td></td>
</tr>
</tbody>
</table>

Source: The Seventh Report of the Joint National Committee on Detection, Evaluation, and Treatment of High Blood Pressure, American Heart Association

N.B.: When a person’s systolic and diastolic pressures fall into different categories, the higher category is used to classify the blood pressure status.

2. **Resting Heart Rate:** Resting heart rate - the number of times your heart beats per minute - will be measured at the wrist (radial artery), using the manual palpation method to feel the pulse. The precise methodology is detailed in the study qualification evaluation section.

If the pre-treatment phase occurs on the same day as the study qualification evaluation phase, then Resting Heart Rate does not need to be measured again.

If the pre-treatment phase occurs on a different day to the study qualification evaluation phase, then the subject’s Resting Heart Rate will be measured again at this time, three consecutive seated measurements, each five minutes apart, with the first measurement occurring after the subject has been seated at rest for about 10 minutes (as during the study qualification evaluation phase). Also as during the study qualification evaluation phase criteria, if the subject’s three-measurement Resting Heart Rate average falls into the bradycardia, tachycardia or irregular categories, then the subject shall be disqualified from further participation in the study at this time. Else, the subject’s Resting Heart Rate will be recorded as the number of beats per minute. A Resting Heart Rate in the range of 60 - 90 beats per minute is considered in the normal range. The average Resting Heart Rate for a male is 70 beats per minute, and for a female is 75 beats per minute.
Quality of Life Assessment Measures

1. The Perceived Stress Scale (PSS): The 14-item PSS questionnaire will be administered during the pre-treatment assessment phase ONLY IF the study qualification evaluation phase has occurred on a different day. Else, the PSS score attained during the study qualification evaluation administration will hold at this time. If the PSS is re-administered during the pre-treatment assessment phase, also as per the study qualification evaluation phase criteria, if the subject’s PSS total score is 25 or greater, indicative of excessively elevated levels of perceived stress, then the subject shall be disqualified from further participation in the study at this time.

Additional information about the PSS can be found in the study qualification evaluation section as well as in Appendix C of this clinical study protocol.

2. The Profile of Mood States (POMS) Standard: The POMS Standard is a factor-analytically derived inventory that measures six identifiable mood or affective states. The POMS is easy and quick to administer and score.

The POMS Standard is a self-report inventory that contains 65 items and takes about 10 minutes to complete. The items pertain to a series of mood states and the subject responds to each item based on how well each item describes his or her mood at the present time (right now). Each item is rated on a 5-point scale ranging from ‘Not at all’ to ‘Extremely.’ The complete POMS inventory is contained in Appendix E of this clinical study protocol document.

The POMS measures six identified mood factors:

• Tension-Anxiety
• Depression-Dejection
• Anger-Hostility
• Vigor-Activity
• Fatigue-Inertia
• Confusion-Bewilderment

The POMS Standard includes psychiatric norms derived from a sample of 100 individuals, college student norms derived from 856 undergraduates, adult norms derived from a group of 400 volunteers aged 18-94, stratified by age, gender and race according to the 1990 U.S. census.

Since 1971, many research studies have provided evidence for the predictive and construct validity of the POMS Standard. Alpha coefficient and other studies have found the POMS Standard to exhibit a high satisfactory level of internal consistency, while product-moment correlations indicate a reasonable level of test-retest reliability. Factor analytic replications provide evidence of the factorial validity of the 6 mood factors, and an examination of the individual items defining each mood state supporting the content validity of the factor scores. Many recent studies continue to add to and affirm the validity of POMS normative sample. A bibliography of published research of almost 3000 research studies from 1964-2002 utilizing the POMS adds to and affirms the validity of the POMS normative sample and is available upon request.

3. Spielberger’s State-Trait Anxiety Inventory (STAI):

The State-Trait Anxiety Inventory (STAI) provides a reliable measure of both temporary and dispositional anxiety in adults. First developed by Charles D. Spielberger in the 1960s, the STAI was
later revised in 1983. The revised STAI is typically referred to as the STAI-Y. The STAI is a self-administered test and it is the most widely used measure of anxiety worldwide, used in both clinical and research settings. It is suitable for adults at a 6th grade reading level or above.

The STAI consists of 40 items divided into two subscales or domains: State Anxiety and Trait Anxiety: State Anxiety assesses an individual’s current level of anxiety – a more temporary state. The 20 items measuring State Anxiety ask subjects how they feel “right now, at this moment,” and reflects situational factors that may influence anxiety levels. Subjects rate their feelings about each statement on a four-point intensity scale of 1=Not at all, 2=Somewhat, 3=Moderately So, and 4=Very Much So.

Trait Anxiety assesses an individual’s anxiety proneness – a more general and long-standing quality of how an individual typically responds to stress. The 20 items measuring Trait Anxiety ask subjects how they “generally” feel. Subjects rate themselves on a four-point frequency scale of 1=Almost Never, 2=Sometimes, 3=Moderately So, and 4=Very Much So). Examples of items Trait Anxiety scale items are “I feel at ease;” “I feel upset;” “I lack self-confidence.”

Scoring: State and trait anxiety are scored separately. Each item is scored from 1-4, for a total inventory score range of 20 to 80, where 20 equals ‘not feeling like that at all (state anxiety) or ever (trait anxiety)’ and 80 equals ‘feeling like that very much (state anxiety) or always (trait anxiety).’ Essentially, the higher the score, the greater the level of anxiety.

Both percentile ranks and standard (T) scores are available for male and female working adults and stratified by age.

Statistical data:

Statistical analysis was conducted on a sample of almost 5,000 adults. For the Trait-anxiety scale, reliability coefficients ranged from .65 to .86, whereas the range for the State-anxiety scale was .16 to .62. This low level of stability for the State-anxiety scale is expected since responses to the items on this scale are thought to reflect the influence of whatever transient situational factors exist at the time of testing.

Regarding validity, correlations between the STAI and other common measures of trait-anxiety are as follows: the Taylor Manifest Anxiety Scale: .80; the IPAT Anxiety Scale: .75; and the Multiple Affect Adjective Check List: .52.

The STAI is contained in Appendix F of this clinical study protocol document. The STAI Manual is available upon request.

4. Beck Depression Inventory®—II (BDI®—II):

Aaron T. Beck, Robert A. Steer, Gregory K. Brown

The Beck Depression Inventory®—II (BDI®—II) is in line with the depression criteria of the Diagnostic and Statistical Manual of Mental Health Disorders—Fourth Edition (DSM–IV). This new edition of the Beck Depression Inventory® is the most widely used instrument for detecting depression.

It takes about five minutes to complete and is demonstrated to be highly clinically sensitive to measurement and change.
The BDI-II consists of 21 items that assess the intensity of depression in clinical and normal patients. Each item is a list of four statements arranged in increasing severity about a particular symptom of depression, evaluated over the period of the past two weeks. It has been validated for samples aged 13-80 years.

Reliability: Internal consistency (Cronbach’s alpha) is .92 for clinical patients and .93 for non-clinical individuals. Test-retest reliability is .93.

Validity: Concurrent validity: two comparisons between BDI-II and its previous version resulted in correlations of .93 and .84, the latter using the take-home form. Other tests found BDI-II to be correlated with the Beck Hopelessness Scale (.68), Scale for Suicide Ideation (.37), Beck Anxiety Inventory (.60), Hamilton Psychiatric Rating Scale for Depression-Revised (.71), and Hamilton Rating Scale for Anxiety - Revised (.47).

Scoring: Most items on the BDI-II are rated on a 4-point scale ranging from 0 to 3. Several items have seven response options to discern differences in behavior or motivation. The BDI-II is scored by adding the ratings for the 21 items. The maximum total score is 63.

Clinical interpretation of total scores uses the following guidelines: 0 to 13 (minimal depression), 14 to 19 (mild depression), 20 to 28 (moderate depression), and 29 to 63 (severe depression). The BDI-II is contained in Appendix G of this clinical study protocol document.

TREATMENT PHASE MEASUREMENTS

The following measurements, using the tools and protocols established during the study qualification evaluation and pre-treatment assessment phases of the study, will occur at each of the following specified time points during the treatment course of the study.

There will be three during-study assessment time points, as follows:

• End of Month 1 (after the 8th study treatment)
• End of Month 2 (after the 12th study treatment)
• End of Month 3 (after the 14th study treatment)

At end of Months 1 and 2 assessment time points, the following measures will be recorded:

• Blood pressure: three-reading average
• Resting Heart Rate: three-reading average
• Perceived Stress Scale (PSS)

At end of Month 3 assessment time point, all of the measures recorded during the pre-treatment phase will again be recorded, as follows:

• Blood pressure: three-reading average
• Resting Heart Rate: three-reading average
• Perceived Stress Scale (PSS)
• Profile of Mood States (POMS) Standard
• Spielberger’s State-Trait Anxiety Inventory (STAI)
• Beck Depression Inventory-II (BDI-II)

• Analyzing Stress in the Body Subject Questionnaire

• Revision of applicable drug, treatment and food/exercise behavior and history variables

• Satisfaction with overall study outcome rating: The subject will be asked to indicate how satisfied he or she is with any overall change in perceived level of stress attained following the treatment administration period with the ECLOSION EPFX/SCIO, using the following five-point scale:
  • Very Satisfied
  • Somewhat Satisfied
  • Neither Satisfied nor Dissatisfied
  • Not Very Satisfied
  • Not at All Satisfied

• Subject perceived group assignment: The subject will indicate whether he or she believes to have been assigned to the treatment or placebo group, and why.

• Investigator perceived group assignment: The investigator will indicate whether he or she believes the subject to have been assigned to the treatment or placebo group, and why.

At any time that is warranted:

At any time that is warranted during the course of the study treatment administration phase, the subject and/or investigator may record the following:

• Adverse Reactions and Events: Any belief or perception that the subject may have experienced an adverse reaction or event as a result of the treatment with the ECLOSION EPFX/SCIO device.

A subject adverse reactions and events sheet will be completed by the Principal Investigator. Any necessary action will be taken. More detailed information on this process can be found in the section below titled: “REPORTING OF ADVERSE REACTIONS AND EVENTS.”

• Additional Comments: A subject or investigator may record any comments related to study participation at any time, as desired.

POST-TREATMENT PHASE

The post-treatment phase will occur two weeks (14 days) following the final treatment administration with the ECLOSION EPFX/SCIO at the end of month three.

The purpose of the post-treatment assessment phase is to gain a sense of duration of treatment effect beyond the cessation of the treatment administration period. At the end of the two-week follow-up period, the following measures will be recorded:

• Blood pressure: three-reading average

• Resting Heart Rate: three-reading average

• Perceived Stress Scale (PSS)

• Profile of Mood States (POMS) Standard
A double-blind placebo-controlled study of the application of the SCIO Universal Electrophysiological Biofeedback System for statistical evaluation of the SCIO’s ability to increase Body Wellness after one 45-minute session

Gage Tarrant, Bart Keough, Jane Summers, Jill Caravalho, Gene Helton, Lynn Smith and Julie Craker in Seattle, U.S.A.;
Jacqueline Jacques, Jean-Pierre Turblin, Adrian Muresan and Anne Préau in Paris, France;
Dr. Codruta Bacean and Dr. Onut Bacean in Timisoara, Romania;
Dr. Rainer Mutschler and Kathrin Sollner in Speyer, Germany.
Dr. Danis in Budapest, Hungary

In this study there were conclusive results of electro-physiological improvement. This study was conducted at five sites of approximately 40 subjects in each.

Under medical supervision the study was conducted in Budapest, Hungary, Timisoara, Romania, Seattle Washington, Speyer Germany, and Paris, France. Thus there were over 200 patients in this medical study.

A host of wellness tests and electro-physiological tests were performed. The results showed significant results with the electro-physiological factors of the VARHOPE. Thus the SCIO was proven to have significant action in improving the electro-physiological field of a patient.

In a review of the results. Patients with dysfunctional wellness of flexibility, blood pressure, and other factors can be improved with just one session. This was a tendency that was not statistically shown at a 5 alpha level, but was seen in the data.

So after this series of experimental tests and evaluation we can conclusively conclude that the SCIO is

1. Safe (no report of any significant risk)
2. Effective at long term stress reduction
3. Effective at short term electro-physiological, charge stability and stress reduction
4. Effective at making short term wellness changes


Introduction

BACKGROUND:
Stress is known to have many negative effects on multiple aspects of an individual’s life. Stress can affect an individual’s physical, cognitive, emotional and social well-being. We hypothesized that within one 45 minute biofeedback session, a measurable improvement in Body Wellness indicators can be achieved.

OBJECTIVE:
To evaluate whether a 45 minute session with the SCIO biofeedback device affects an individual’s Body Wellness.

DESIGN: Randomized, double-blind, placebo controlled trial.

SETTING AND PATIENTS: 192 individuals with awareness of levels of perceived stress as well as injuries/pain, between 18 and 65 years of age, male or female, randomized into placebo and SCIO test groups, at private clinics at multi centered sites.

INTERVENTION:
Subjects were randomized to the test group (a 45 minute SCIO biofeedback system session) or placebo (SCIO, Maitreya Kft. www.qxsubspace.com)

MEASURES:
Pre and post measures as follows: Quality of Life Questionnaire, Energy Index Factor (systolic blood pressure left arm sitting + the diastolic blood pressure left arm sitting x pulse), Grip Strength Test (measured in kilograms), Oxygenation Test, Flexibility Test (measured in inches in The USA and centimeters everywhere else), Memory Test, pH Test, VARHOPE scores (electrical measures within the device, as follows: voltage (V), amperage (A), resistance (R), hydration (H), oxygenation (O), proton pressure (P) and electron pressure (E)).

RESULTS:
Patients in SCIO group had greater VARHOPE scores than those in placebo group (p<0.005). The other indicators of Body Wellness were not statistically significant, but there are trends in the improvement levels between the Test and Control (Placebo) groups.

CONCLUSIONS:
The electrical parameters of VARHOPE can be improved by a 45 minute SCIO biofeedback session. However, it may require more SCIO biofeedback sessions for the other Body Wellness Indicators to be increased. Further studies are suggested.

Stress is known to have many negative effects on multiple aspects of an individual’s life. Stress can affect an individual’s physical, cognitive, emotional and social well-being. Different applications of biofeedback have been shown to be effective on stress management and health. Biofeedback is usually combined with a relaxation technique, applied before, during or after the biofeedback training. Studies have evaluated the effectiveness of biofeedback combined with a relaxation technique, EEG Biofeedback, EMG Biofeedback, and HRV Biofeedback and found them to have positive results on reducing the stress burden and alleviating the conditions. In the fifth grade we learned that our bodies are made of atoms. And atoms are made mostly of protons, neutrons and electrons. There are great spaces between these electrons and protons and other atoms. Our bodies are made up mostly of electrons and their electromagnetic fields.
In Hydrogen if the protons are like marbles, the electron is over a kilometer away the next atom’s electron is over 2 kilometers away, the next proton is over 4 kilometers away. So there is more than 99.99999999999999% empty space. This space is filled with energetic fields. The electrons never touch each other so what we are made of is interacting electromagnetic fields.

Atoms are 99.999999999999% empty space and the empty space between atoms is just as or emptier 99.9999999999999999% Electrons repel of course so the atoms with outer electrons repel each other. Why don’t things pass right through things?

Things don’t fall through other things because they are levitating on an energetic electrostatic fields. When you sit on a chair, you are not really touching it. You see, every atom is surrounded by a shell of electrons. This electron cloud presents a rather negative face to the world. Remember that like charges repel each other. When two atoms approach each other, their electron shells push back at each other, despite the fact that each atom’s net charge is 0.

When two atoms come together and have empty spaces in their electron quantum shells, they will share electrons to fill in the spaces in both of their shells. The electrons really do go back and forth between atoms and they do so pretty fast. Outer Electrons tend to be kind of mobile, which is also a very nice feature of nature, since without it your walkman would not work or you would not be alive. It is the free electrons and protons in the body that allow life. Once both atoms’ outer shells are full due to this electron sharing, they go back to their usual repulsive behavior. This, by the way, is how we get molecules, hormones, enzymes etc and the secret to understanding Chemistry, Biology, Medicine, Physiology etc. It’s all about the electrons and protons, charged particles and vibration.

The electrons and atoms of our complex Fractal body obey quantum, QED, photonic, electromagnetic-static laws. This is a mouthful so we abbreviate and since these are all energy let’s say ENERGETIC.

There is undeniably a body electric and there is indeed an Energetic Medicine. Only a presumptive fool would assume otherwise. There is pressure from the chemical companies and their vast wealth and pervasive influence to view the body as a set of chemicals. But these chemicals are all made of energetic fields and they obey energetic laws like quantum, electro-magnetic, static, quantum electro-dynamic photonic laws.

This study was designed for a biofeedback device (SCIO) that combines all known applications of Biofeedback and a unique protocol that addresses specific Body Wellness Indicators.

The SCIO is designed to correct the manifestation of stress and/or electro-stress patterns within the individual at the most primal of physiological levels. The device works on the theory that stress disrupts the inherent electromagnetic frequencies at which the body’s cells, organs, etc. resonate and that by returning these frequencies to their natural state, the stress and any subsequent illness that occurred because of the disruption can be corrected.

The objective of the study was to determine if one 45-minute treatment with the SCIO would show a change, and hopefully an improvement, on a person’s Body Wellness indicators.

Methods

• Study design

The study was designed as a randomized, double blind, placebo controlled study. The goal of the study was to analyze the changes in Body Wellness Indicators after one 45-minute SCIO session, so there was no follow up necessary days or weeks after the treatment.
All study personnel and participants were blinded to treatment assignment for the duration of the study. Only the study statisticians and the data monitoring committee saw unblinded data after the study sites are completed, but none had any contact with study participants, nor will they ever have contact with future participants.

Randomization was assured at each testing site by the one person organizing the subject scheduling prior to the study. As people called to schedule their appointments based on the Subjects availability, the Scheduler would randomly assign the subject to one group or the other depending on which room was available. None of the staff members involved in the clinical trial process were aware of what group/device was placebo and what group/device was real. Each testing site was given two pieces of equipment - placebo equipment and actual equipment. Testing sites were chosen on the basis that the staff was knowledgeable of the functions of the device and are well trained and supervised to conduct the study. Either the Clinical Investigator or an Independent Monitor supervised the study.

Subjects assigned to Testing Group A were treated with the SCIO device A using Harness A and subjects assigned to Testing Group B were treated with the SCIO device B using Harness B. Only the study Sponsor knew which label (‘A’ or ‘B’) corresponded to the actual (test) device and harness and which label corresponded to the placebo device and harness.

The placebo equipment was designed to have the same external physical appearance as the actual equipment. The software operating the placebo devices is designed to look exactly as the one operating the real device, with no distinguishing differences. The difference is that the placebo harness was not equipped internally with functional electrodes and the programming for the placebo device output only blank matrices. Neither the actual (test) nor the placebo harness produced any detectable noise, heat, light or other sensation output, so this also wasn’t a distinguishing factor for subjects or the investigator between the actual (test) and placebo devices.

To evaluate blinding, at the end of the session, both subject and investigator were asked to indicate which group they believed the subject to be assigned to (SCIO Test, placebo) and what led to that belief.

The study was completed in five testing sites on the following dates: Budapest, Hungary, 5-10 August 2009, Timisoara, Romania from 2-4 September 2009, Seattle, Washington, from 23 – 25 October 2009, Paris, France from 17-20 November 2009 and Speyer, Germany from 23 – 27 November 2009.

The investigation was initiated on the 2nd of September 2009, in Timisoara, Romania, and completed on the 27th of November 2009 in Speyer, Germany.

• Patients

A sample size of 40 patients per site was calculated. The sample size of 45 subjects per group (test and control, separately) has been determined using Table A.3. Sample sizes per group for a two-tailed test on proportions. P1=.20, on page 266 of the textbook, Statistical Methods for Rates and Proportions, Second Edition, Joseph L. Fleiss, Division of Biostatistics, School of Public Health, Columbia University, 1981, John Wiley & Sons, Inc. Publishers, New York, NY. To apply the values in this table to a one-tailed test, the alpha value of 2*alpha (0.05) was used.

From here, it was anticipated that about one-twelfth of subjects overall may withdraw from the study prior to completion for various reasons, including the length of the treatment period.

Final sample size = sample size X 1/(1-d); where d = # expected dropouts/# subjects enrolled.
Final sample size = \(45 \times \frac{1}{1 - 0.089}\); where \(d = \frac{4}{45}\)

Final sample size = \(45 \times \frac{1}{0.911} = 45 \times 1.098 = 49\) subjects per group.

Therefore, a minimum starting sample size of 49 subjects in each treatment group was needed to ensure that a sufficient number remains at the end of the trial (40 subjects per group) for any significant differences found between groups to be considered statistically valid and representative of the general population being sampled. For ease of division between the test sites, the number has been rounded up to 50 subjects per treatment group.

Patients were locally recruited in Speyer, Seattle, Paris, and Timisoara, from the pool of potentially suitable patients who normally attend the test sites for various services or form nearby consenting and suitable medical offices and other such suitable locations. Respondents were invited to the session. After giving written Informed Consent forms, patients were screened by investigators.

Inclusion criteria were perceived levels of high stress, injury and/or pain (based on a Quality of Lifestyle Questionnaire), age between 18 and 65. Exclusion criteria included extremely sick patients on more than 5 prescribed drugs, crippled and handicapped patients, diagnosed heart conditions, prior head traumas, pregnancy, breastfeeding or planning pregnancy, pacemaker use, serious mental illnesses, prior cardiac surgeries, seizure disorders, developmental disability or cognitive impairment, participation in other medical research in the past 30 days. At some sites,

Inclusion/Exclusion Criteria were evaluation via email or phone in advance.

Participants deemed eligible based on baseline assessment were randomly assigned to either the test (SCIO) group or the control (placebo) group. Subjects in the test group received the actual study treatment with an active, operational harness, and those in the control group received a ‘fake’ study treatment with a placebo harness that does not contain any active electrodes. The investigators follows the same protocol for all subjects. None of the participants could discern if they were in one group or the other as the devices and equipment looked and felt the same.

Levels of stress were assessed by self-report questionnaires. Body Wellness Indicators were evaluated through physiological measurements and electrical device measurements.

The potential for adverse reactions were monitored at each test site according to the Freiburger Ethics Committee International, Germany, which approved the study protocol. However, no adverse reactions were reported during the study or after the study.

• Outcome measures

The study was designed to determine the device’s efficacy by recording baseline measures in the pre-treatment phase, against which post-treatment changes were assessed. There were 8 types of tests performed pre- and post – treatment, for both test group and placebo group.

First, the levels of stress and pain/injury were assessed using a self-report Quality of Life Questionnaire. The questionnaire had 10 questions to be answered on a scale from 1 to 10 (the higher the stress, the higher the value).

Then physiological measurements were taken for the assessment of Body Wellness Indicators, defined as follows: Energy Index Factor, Strength Test, Oxidation Test, Flexibility Test, Memory Test, pH Test, VARHOPE test.

Energy Index Factor was calculated using the formula:
Systolic BP left arm sitting + the diastolic BP left arm sitting x pulse = energy index factor.

The Energy Index Factor indicates parasympathetic control below 9,000, balance at 14,000 and sympathetic neural control at 18,000+.

The strength test was performed using a hand-held Dynamometer. Subject held the Dynamometer in one hand, gave one big squeeze with one hand on the Dynamometer while the investigator documented how many kilograms of strength the patient was able to exert.

The same procedure was repeated for the other hand.

For the anaerobic oxidation test, the patients started in a seated position, relaxed and breathed at a normal breath rate for 1 minute. The subject took a deep breath at the same time that the investigator started a stop-watch counting minutes, seconds and tenths-of-a-second. Subject stood up at a normal speed and sat down again at a normal speed while still holding their breath as long as possible. As soon as the subject stopped holding breath and took a new breath then investigator stopped the stop-watch. Investigator documented the length of time that the subject held breath during Anaerobic Oxygenation Test.

To evaluate flexibility, there were three types of flexibility tests performed. Low back flexibility was measured with the subject sitting down on the floor with legs stretched out in front, heels approximately 20.32 cm (8 inches) apart. The subject extended both hands, outstretched fingers towards their heels keeping legs straight. Subjects were asked to do a maximum stretch and touch the floor as far as they can, even going past the heels, if they could. The distance in centimeters (or inches in The USA) from where their fingers touched the floor to the heels, with Zero at the heels, positive if they can extend past the heels, minus if they are before the heels. Normal scores are anywhere from -7.62 centimeters (-3 inches) to 0 centimeters (0 inches), scores below -7.62 centimeters (-3 inches) indicate low back difficulty. Ideal score should be 17.78 centimeters (7 inches) past the heels.

Side to Side Flexibility was measured with subject standing on their knees without bending forward or backward at the waste, and leaning to the left side, trying to touch their left palm to the floor.

Normal readings are touching fingers or knuckles to floor. An advanced subject would be able to touch their palm. If they cannot touch their fingers it indicates a lack of flexibility. A protractor was used to determine the angle of flexibility.

Neck flexibility is measured with the subject trying to touch their ear to the shoulder without raising their shoulder to their ear. A protractor was used to determine the angle of flexibility.

The suggested method for the memory test was to choose a first set of 5 random numbers. If the subject remembered these numbers in the proper sequence (either forward or backward depending on the memory test), then the investigator added 2 to the existing digit and one more digit to the end to increase the sequence by one digit. This method was continued until the end.

The investigators recorded the number of memorized digits (forward and backward).

An Over-The-Counter pH (acidity-alkalinity) Test Kit was purchased and used for pre-test and post-test measurements of pH.

VARHOPE is an acronym coined by the manufacturer in which V = Voltage, A = Amperage, R = Resistance, H = Hydration, O = Oxygenation, P = Protons and E = electrons.
The VARHOPE numbers are measured by the biofeedback device, during a 1.5-minute Calibration process and they are measured again at the end of the session.

The VARHOPE numbers are separated into two (2) categories where VARHO is one category and PE and the second category. The VARHO readings are set on a scale, determined by the manufacturer, from 0-110. Changes in the readings are shown on the scale where an increase in number shows in improvement.

The PE readings are set on a scale, determined by the manufacturer, where the closer the P reading to 75 the more the client’s state of wellness has stabilized, and the closer the E reading to 65 the more the client’s state of wellness has stabilized. This scale was chosen to make it easier for the layperson to accept readings on a simpler scale rather than electrical readings.

Voltage is derived directly from the skin electro-potential amplitude. Amperage is the amount of charged particles flowing and Voltage is the pressure behind the flow. All biofeedback instruments measure voltage which is the electro-potential of the skin underneath the electrode. Amplitude disorders in EEG or ECG, refer to the voltage vector.

Amperage is calculated from the volume of current over a short period of time coming off of the body. Knowing the skin’s electro-potential and impedance using Ohm law of V = A x R (Voltage equals Amperage times Resistance), the current or amperage at a transcutaneously measured point can be calculated. This is referred to in the industry as a virtual measured variable. Using simple collective mathematics global measures of our patient’s Voltage, Amperage and Resistance can now be established. Amperage is the amount of charged particles flowing and Voltage is the pressure behind the flow.

Resistance is a measure of how a substance resists current flow. In a complex situation of multiple liquids acting as electrolytes and multiple membranes, there are induction and capacitance enhancements to the flow (magnetic and capacitance). Resistance is measured in Ohms.

Hydration

The active stability of water concerning the amount of free water and its ability to permeate osmotic membranes. The electrical pulse of the SCIO stimulates osmosis. It is measured as the range of voltage scores over a period with the aberrant signals and cardio-signals filtered out give us a Hydration index. Voltage changes observed during the Calibration process give us a Hydration index (based on the free proton effect).

Oxygenation is given through the range of amperage scores over a period of time, with the aberrant signals and cardio-signals, which, when filtered out give an Oxygenation index. The range of scores, between maximum and minimum, reflects oxygenation giving an Oxygenation index. As the changing Voltage and Amperage is measured we get inductance and capacitance virtual scores and this allows us to find a Hydration and Oxygenation index. As Amperage changes slightly with each breath, we get an Oxygenation index from comparing maximum and minimum values. We need to observe several normal breaths to establish an Oxygenation index during the Calibration procedure.

Proton balance (in relation to Electron balance) The polarity of the signal and the collective global readings give us a proton-electron balance. Thus the system has an index which can be compared to established norms of patients based on age, stress, metal implants in teeth, smoking and behavior. Proton and Electron balance (or the charge stability of the client) affects the polarity and the resting potential. The slight changes in these electrical profiles can be measured. This is measured globally.
Improvements in the VARHOPE Scores show improvements in the body’s natural electrical functions.

For the purpose of this clinical study, the VARHOPE measurements were taken at the beginning of the test session with the SCIO and then again towards the end of the test session with the SCIO.

- **Intervention**

The SCIO biofeedback session was 45 minute in length. The SCIO device utilizes transcutaneous voltammetric evoked potential biofeedback technology, which consists of both hardware and software. The hardware consists of a digital interface box attached to the computer with electrodes attached to the wrists, ankles, and 8 on the forehead of the person (making up 12 transcutaneous carbon impregnated rubber electrode contact points). The software is a PC-based platform consisting of mathematical calculations and high-end graphics. The placebo equipment was designed to have the same external physical appearance as the actual equipment. The difference is that the placebo harness was not equipped internally with functional electrodes and the programming for the placebo device outputted only blank matrices. Neither the actual (test) nor the placebo harness produced any detectable noise, heat, light or other sensation output, so this also wasn’t a distinguishing factor for subjects or the investigator between the actual (test) and placebo devices.

After the pre measurements were taken, patients were invited to a quiet room, seated comfortably in a chair, connected to the device via harnesses and advised to be as relaxed as they can. The session protocol included relaxation treatments, Neuro Linguistic Programming treatments, electro-acupuncture, and biofeedback treatments.

After the 45 minute session, the subjects were ready for the post-treatment phase, were the measurements were repeated and recorded.

- **Statistical analysis**

Independent statisticians were hired to analyze the data and determine statistical significance. Success of the study was determined by simple calculation of the percentage of subjects in each treatment group who met the individual subject success criteria. If these percentages showed that the overall study success criteria are met, the study will be considered to have had a successful outcome.

In addition, the primary efficacy outcome measure was evaluated. A one-tailed z-test of proportions was conducted to assess for a statistically significant difference in the average post-treatment Body Wellness scores for test versus control group subjects. Using the paired t-test, the percentage differences between the mean differences in final post-treatment Body Wellness score minus pre-treatment Body Wellness score for subjects in the test group versus subjects in the control group was evaluated.

For the primary outcome measure, an intent-to-treat analysis (including all randomized patients), and a per-protocol analysis (subjects without major protocol deviations, incompletes excluded) were performed.

Handling of missing data in the per-protocol analysis was according to the multiple imputation method.

For the evaluation of the secondary efficacy outcome measures an ANOVA was used to evaluate the change in total Body Wellness score across the measurement time points of pre-treatment and post-treatment, comparing test and control group subjects. It was expected that more test group subjects will demonstrate an improvement in Body Wellness in one or more tests from pre-treatment to
treatment to post-treatment than will control group subjects. There were made correlations between scores on the various inventories and a z-test was used to evaluate differences in outcome satisfaction ratings between test and placebo group subjects. Changes recorded on the VARHOPE readings during treatment and comments provided by subjects were also evaluated. A safety outcome evaluation of any reported adverse events and reactions was performed.

The study flow chart, based on the Consolidated Standards of Reporting Trials recommendations, is shown in Figure 1. Subjects were recruited through local contacts on a voluntary basis. Some were screened beforehand via phone or e-mail and others were screened upon arrival. Potential candidates were excluded or deemed ineligible for the following reasons: a) did not meet inclusion criteria (n=3), b) scheduling problems (n=2), c) missed appointments (n=7). 151 subjects were enrolled and randomly assigned to either the SCIO Test Group (n=86) or Placebo Group (n=65). The subjects not included in analysis were those from whom no pre and/or post measurements were recorded at the end of the session. Reasons for not obtaining data were scheduling problems.

![Figure 1. Flowchart of the study of the application of the SCIO Universal Electrophysiological Biofeedback System for statistical evaluation of the SCIO’s ability to increase Body Wellness after one 45-minute session](image-url)
Results are shown in Figure 2. There was no statistical difference between groups in the following outcomes: Quality of Life Questionnaire, Energy Index Factor, Strength Test, Flexibility, Memory and pH.

<table>
<thead>
<tr>
<th>SCIO Treatment group (Test group)</th>
<th>Mean</th>
<th>N</th>
<th>Std. Deviation</th>
<th>Std. Error of Mean</th>
<th>Range</th>
</tr>
</thead>
<tbody>
<tr>
<td>x Quality of life Questionnaire 1</td>
<td>0,5412</td>
<td>85</td>
<td>0,92005</td>
<td>0,09979</td>
<td>5,00</td>
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<tr>
<td>x Quality of life Questionnaire 2</td>
<td>0,2353</td>
<td>85</td>
<td>0,66632</td>
<td>0,07227</td>
<td>4,00</td>
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<tr>
<td>x Energy Index Factor</td>
<td>871,3294</td>
<td>85</td>
<td>2 309,45944</td>
<td>250,49605</td>
<td>12 770,00</td>
</tr>
<tr>
<td>x Left Hand Strength (Kg)</td>
<td>0,4118</td>
<td>85</td>
<td>3,89839</td>
<td>0,42284</td>
<td>29,00</td>
</tr>
<tr>
<td>x Right Hand Strength (Kg)</td>
<td>-0,8235</td>
<td>85</td>
<td>5,06664</td>
<td>0,54955</td>
<td>48,00</td>
</tr>
<tr>
<td>x Oxyegnation Low Back</td>
<td>-1,1786</td>
<td>84</td>
<td>3,87193</td>
<td>0,42246</td>
<td>32,00</td>
</tr>
<tr>
<td>x Flexibility Side to Side</td>
<td>-0,6588</td>
<td>85</td>
<td>7,97549</td>
<td>0,86506</td>
<td>55,00</td>
</tr>
<tr>
<td>x Flexibility Neck</td>
<td>0,8706</td>
<td>85</td>
<td>11,22687</td>
<td>1,21772</td>
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<tr>
<td>x Memory Test (Forward)</td>
<td>-0,2706</td>
<td>85</td>
<td>1,31283</td>
<td>0,14240</td>
<td>6,00</td>
</tr>
<tr>
<td>x Memory Test (Backward)</td>
<td>-0,1624</td>
<td>85</td>
<td>1,31927</td>
<td>0,14309</td>
<td>8,00</td>
</tr>
<tr>
<td>x pH level</td>
<td>0,6012</td>
<td>84</td>
<td>6,63838</td>
<td>0,72431</td>
<td>62,20</td>
</tr>
<tr>
<td>x Voltage</td>
<td>-14,4286</td>
<td>84</td>
<td>8,98632</td>
<td>0,98049</td>
<td>40,00</td>
</tr>
<tr>
<td>x Amperage</td>
<td>-15,8333</td>
<td>84</td>
<td>9,78791</td>
<td>1,06795</td>
<td>58,00</td>
</tr>
<tr>
<td>x Resistance</td>
<td>-16,9167</td>
<td>84</td>
<td>11,80383</td>
<td>1,28790</td>
<td>62,00</td>
</tr>
<tr>
<td>x Hydration</td>
<td>-16,6071</td>
<td>84</td>
<td>9,84949</td>
<td>1,07467</td>
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<tr>
<td>x Oxygenation</td>
<td>-18,7381</td>
<td>84</td>
<td>12,49879</td>
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<tr>
<td>x Proton pressure</td>
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<td>84</td>
<td>4,09572</td>
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<td>27,00</td>
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<tr>
<td>x Electron pressure</td>
<td>3,6548</td>
<td>84</td>
<td>3,89159</td>
<td>0,42461</td>
<td>28,00</td>
</tr>
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</table>
Table 1. Summary results for the SCIO Treatment (Test) group

<table>
<thead>
<tr>
<th></th>
<th>Placebo group (Control group)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Mean</td>
</tr>
<tr>
<td>x Quality of life Questionnaire 1</td>
<td>0,4462</td>
</tr>
<tr>
<td>x Quality of life Questionnaire 2</td>
<td>0,2000</td>
</tr>
<tr>
<td>x Energy Index Factor</td>
<td>885,7077</td>
</tr>
<tr>
<td>x Left Hand Strength (Kg)</td>
<td>-0,0615</td>
</tr>
<tr>
<td>x Right Hand Strength (Kg)</td>
<td>0,0769</td>
</tr>
<tr>
<td>x Oxygenation</td>
<td>-1,2500</td>
</tr>
<tr>
<td>x Flexibility Low Back</td>
<td>-2,0580</td>
</tr>
<tr>
<td>x Flexibility Side to Side</td>
<td>1,5846</td>
</tr>
<tr>
<td>x Flexibility Neck</td>
<td>2,0615</td>
</tr>
<tr>
<td>x Memory Test (Forward)</td>
<td>-0,1846</td>
</tr>
<tr>
<td>x Memory Test (Backward)</td>
<td>-0,4923</td>
</tr>
<tr>
<td>x pH level</td>
<td>-0,1246</td>
</tr>
<tr>
<td>x Voltage</td>
<td>0,6923</td>
</tr>
<tr>
<td>x Amperage</td>
<td>-0,9077</td>
</tr>
<tr>
<td>x Resistance</td>
<td>-0,3538</td>
</tr>
<tr>
<td>x Hydration</td>
<td>-1,4769</td>
</tr>
<tr>
<td>x Oxygenation</td>
<td>-0,8462</td>
</tr>
<tr>
<td>x Proton pressure</td>
<td>0,5231</td>
</tr>
<tr>
<td>x Electron pressure</td>
<td>-0,2154</td>
</tr>
</tbody>
</table>
Table 2. Summary results for the Placebo (Control) group

[Diagram showing summary results with significant and not significant indicators]

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There was significant difference between groups in the VARHOPE measures ($p < 0.005$). The results were as follows:

- 86.90% of subjects had a V improvement of more than 5%,
- 90.47% of subjects had an A improvement of more than 5%,
- 88.09% of subjects had an R improvement of more than 5%,
- 88.09% of subjects had an H improvement of more than 5%,
- 89.28% of subjects had an O improvement of more than 5%,
- 47.61% of subjects had a P improvement of more than 5%,
- 48.80% of subjects had an E improvement of more than 5%.

There were no adverse events reported during or following the study.

**Discussion**

We conducted a double blind, randomized study with a placebo controlled design to evaluate an intervention comprising a 45 minute SCIO Biofeedback session. We looked at the impact of this session on electrical scores (VARHOPE), perceived levels of stress and pain/injury, blood pressure, strength, memory, flexibility and pH in subjects with high levels of stress. Significant effects associated with the intervention were observed in VARHOPE scores. The biofeedback session had no risks associated; compliance to study protocol was maintained during the study.

The body is an electrical process requiring volts and amps which conduct through resistance circuits to operate. All muscles are turned on by electrical impulse. Muscles themselves are magnetic using volts and amps for their operation. The brain is a massive collection of cells that have electrical activity that can be measured via the EEG. The heart itself is the largest electro-magnetic engine and uses electrical impulses to influence the muscles of the heart to circulate blood. Most EEG, ECG and EMG measures are only concerned with oscillatory properties and not so concerned with the volume or basic amperage of the circuit. The collective baseline of the circuit of EEG, ECG, EMG and GSR measurements can give us a rating of the global body voltage, body amperage and skin resistance (V, A, and R respectively). There are norms of the V, A and R and certain people, due to stress or other factors, can have low V, A and/or R.

The collective inductance and capacitance changes in the body are a reflection of redox potential and can reflect hydration (H) and oxygenation (O) indexes.

Irregularities in EEG, EMG, ECG and GSR can be corrected through guided electro-stimulation. The V and A are also greatly affected by the charge stability of the free protons and electrons, (negative charges and positive charges in the body), which collectively make up the proton pressure (P) and electron pressure (E). The acidity-alkalinity balance is an electrical measure of the amount of positive...
versus negative charged particles. This can be measured by the carbon based electrodes of the SCIO through electro-stimulation biofeedback.

This Clinical Study was developed in the context of substantiating the Indication for Use for the SCIO biofeedback device of rectifying charge stability imbalance and rectifying redox potential, two of the factors that influence Body Wellness. One study objectives was to determine if one 45-minute treatment with the SCIO would show a change on a person’s Body Wellness indicators as defined by the study hypothesis, which is now proven valid.

The other indicators of Body Wellness (Quality of Life Questionnaire, Energy Index Factor, Strength, Anaerobic Oxygenation Test, Flexibility, Memory and pH) were not statistically significant. This does not mean that the results are not valuable. A closer analysis shows that there are trends in the improvement levels between the Test and Control (Placebo) group. As per the hypothesis defined in the Clinical Study protocol it was expected that any positive change in post-treatment measures for control subjects is expected to occur to a significantly lesser degree than for subjects in the test group.

The subjects were not equally assigned to the placebo versus test group, but as follows:

Timisoara, Romania
Control Group n=17
Treatment Group n=21

Seattle, Washington, U.S.A.
Control Group n=20
Treatment Group n=20

Paris, France
Control Group n=20
Treatment Group n=20

Speyer, Germany
Control Group n=8
Treatment Group n=25

Because the test/placebo assigned ratio was not equal, in order to determine trends of improvement per the study hypothesis, we have analyzed subjects from Seattle, USA and Paris, France (40 subjects in test group, 40 subjects in placebo group).
Strength test results analysis showed that 60% of the subjects in the test group had a 5% or more improvement in left hand strength, as opposed to only 35% of the placebo group which had a 5% or more improvement. In the case of the right hand strength, difference is smaller, however, 47% of the test group subjects improved more than 5%, and 37.5% of the placebo group improved more than 5%. The results show that there is definitely a possibility of improving strength with the SCIO biofeedback device, the question that remains is how many SCIO sessions would have a statistically significant result. Also, another thing to consider while analyzing the results is that the protocol followed included mostly general stress reduction therapies, as opposed to using specific muscle strengthening treatments, which could have a greater impact on strength.

It is interesting how the anaerobic oxygenation test had a greater improvement for the placebo group (68.42%). Test group had an 55.26% of subjects that had a 5% or more improvement. The question that rises is whether the improvement occurred because the subjects had a chance to relax during the 45 minute protocol or because of the biofeedback session. The anaerobic oxygenation test requires a clear mind and a rested body. So whether the results are due to the relaxing 45 minute session or not is unclear.

Analysis of flexibility back shows that 60% of the subjects in test group had a 5% or more improvement versus 55% of the placebo group that showed a 5% or more improvement. 10% more of the subjects in the test group had a 5% or more improvement of flexibility side versus subjects in placebo group. In Seattle, side to side flexibility showed statistically significant improvement. Even though the overall analysis does not support these findings, it is a strong basis for future studies. Also, evaluating pathological versus non-pathological data revealed that, for one site (Paris) subjects with pathological reactions to the low back flexibility test had non-pathological reactions after the biofeedback session.

Flexibility of the neck has again quite similar results. 25% of the test group and 27.5% of the placebo group had a 5% or more improvement. It would be interesting to find out whether flexibility would be significantly improved by specific muscle therapies with the SCIO device. Considering the relaxation therapies that were applied in this study, it is definitely worth finding out how many specific SCIO treatments would have a considerable effect of flexibility measurements.

Memory test did not suggest any trends, improvement of more than 5% being observed at an exactly the same percent of the test group subjects as the placebo group subjects.

Energy Index Factor, a variable based on blood pressure measurements did not show statistical significance. However, a study that evaluated the effects of GSR Biofeedback and Progressive Muscle Relaxation showed that PMR induced a significant decrease in blood pressure whereas GSR biofeedback training showed a decrease in respiratory rate. The GSR treatment was administered for 20 minutes daily, for 10 consecutive days. This provides a strong basis for a further analysis of the SCIO device sessions and the effects they might have on blood pressure.

This study has limitations, primary among them the lack of follow-up. Nevertheless, the results obtained after one session provided valuable data on feasibility and plausibility. Another limitation was the short duration of the study. An alternative design would provide data obtained over a

significantly longer period of time, which may also influence the results of the study and offer better understanding of the efficacy of a sequence of session with the SCIO device.

- Conclusions

The SCIO device appears to be a valuable tool in improving Body Wellness. One 45 minute session had significant results in improving the natural electrical parameters of the body. The results also showed trends of improvement in other body variables, therefore providing a basis for future studies. The fact that there were no adverse events reported shows device safety.

- Acknowledgements

The study was funded by Maitreya Ltd., manufacturer of the SCIO Universal Electrophysiological Biofeedback System. Maitreya Ltd. was involved in the design and conduct of the study and provided logistical support during the trial. Employees of the sponsor worked with the investigators, but the analyses were performed by two independent statisticians. The manuscript was prepared by Dr Mutschler. Maitreya Ltd. was permitted to review the manuscript and suggest changes, but the final decision on content was exclusively retained by the authors.

- References


IJMSSHNEM journal #089 2013 United States of America Research/
IJMSSHNEM journal #090 2013 Romanian Study of the Stimulation of Sports Ability/

2013 USA new studies Validating the SCIO auto focusing Cybernetic loop

STUDY INFORMATION:

SUPERVISING RESEARCHERS: Dr. Danis György, MD,
Dr. Hilf Klara MD, Jozsef Mezei MD

MEDICAL CONSULTANT:
Dr. Pauline Willis, USA, Dr. Gebhard Gehring MD Bavaria, Germany

DATE and PLACE: 2008 – 2013 Arizona, USA

SPONSORS:
SCIO International / Maitreya Kft.

INSTITUTIONAL MONITOR:
IMUNE / University of Timisoara (Victor Babes University of Medicine) Dr. Bacean Aurel MD
USA IRB-Freiburger Ethik-Kommission International (FEKI)
Registered at Amtsgericht Freiburg i.Br. HRB 5010,
Registered according to § 20 Abs. 7 MPG at Bundesinstitut für Arzneimittel und Medizinprodukte (BfArM) under Reg. No.: GS 4.1-A 1871 2375/95

Double Blind Study of Sport Performance with the SCIO device versus Placebo control 2013 USA

Written by Darwin Davidson Doctor of Quantum Biofeedback

This study took 46 healthy athletic subjects over a period from 2007 to 2012 and measured their strength power performance before and after a SCIOtherapy and compared to Placebo control group.

This study showed an increase in strength performance in the treatment SCIO group versus the control group in most patients.
Trauma Sport Pain Electro Healing With SCIO-2013 USA

Written by Darwin Davidson Doctor of Quantum Biofeedback

In this study 27 fit healthy subjects in Arizona USA were hit with a sport injury of the same strength on each leg one at a time. The one leg would get real SCIO therapy the other leg would get Placebo. After the SCIO or control treatment the athletes rated the pain in 10 min intervals till pain recovery was stable. The SCIO showed ability to lower pain after a slight sport injury and promote flexibility recovery quicker than placebo treatment. It is proposed that the increase in osmosis and the autofocused injury treatment pulse increases the body’s natural ability to deal with pain and heal. Transcutaneous Electro-Nerval Stimulation for pain and Electro Wound Healing for injury have been well documented in the literature. This study has shown conclusively that the SCIO technology is significantly safe and effective in treating sport pain and minor injuries.

MCES and Addiction Control a DbI Blind Clinical Study -2013 USA

Written by Darwin Davidson Doctor of Quantum Biofeedback

This study was done in a medical doctor supervised clinical setting in Arizona USA from 2008 through 2013. 37 patients with tobacco addiction and 21 patients with alcohol addiction diagnosis were given SCIO MCES treatments or Placebo treatments for three to ten sessions. There was three report of headache logged and no report of any significant risks. Patients were asked to rate their desire or cravings as scalar numbers from one to ten rating. There was significant evidence of the SCIO MCES reducing craving versus the Placebo control group. The literature discussion proves that there is a wealth of evidence for MCES ability to reduce addiction cravings. The Autofocused Cybernetic Stimulation of the SCIO technology has an improved ability to help stabilize emotional and reduce aberrant addictive impulses. The literature shows MCES has positive results to lower addiction craving and to stabilize emotional depression. The significant evidence of the SCIO technology’s ability to make this claim is now firmly established. Discussion will show a statistically significant positive effect on addiction and emotions using MCES and trans-cranial-cutaneous electrodes.

SCIO’s Effect on Body Osmosis2013 -USA

Written by Darwin Davidson Doctor of Quantum Biofeedback

In this study 41 subjects were hooked to the SCIO with the SCIO off and a line drawn firmly on their forearm with a finger nail. Then we count the seconds it takes before the line turns red. This is an indication of osmosis as that the traumatized tissue will have histamine rush in the traumatized tissue of the forearm. The time it takes indicates the osmosis ability. Norms are 8 to 10 sec. Then the SCIO is turned on and the line drawn on the forearm again of the subject. In the control group there was an average of 12 seconds and 4 did not have the line appear after the 20 sec deadline. The SCIO group had an average of 9 seconds and all were under the deadline. This improvement of treatment group over control demonstrates the SCIO’s ability to increase osmosis thru its auto-focused electrical pulsation. This verifies the SCIO ability to enhance osmosis.

Stimulating Eye Hand Coordination With SCIOVARHOPE Update 2013

Written by Darwin Davidson Doctor of Quantum Biofeedback

In this 53 subject study we review the history of the SCIO sport medicine use with an eye on eye hand coordination 21 athletic males from 13 to 43 were asked to shoot basketball free throws in a double blind fashion after being on the SCIO or after a placebo treatment in 2009 -2013.
In July 2009 15 athletes were given the same double blind test with very similar results. In Arizona, USA 18 subjects were tested with darts accuracy to determine the ability of the SCIO to increase eye hand coordination. The SCIO treatment was proven effective in safely and reliably increasing eye hand coordination. So in total 53 people participated in a double blind study with reversal patterns that prove the ability of the SCIO autofocused stimulation to improve eye hand coordination. Significant results will show that increase in VARHOPE from the treatment group correlate to increased performance of eye hand coordination needed for free throws, darts and other coordination challenges.

SCIO Effects on Oxidation/Oxygenation 2013
Written by Darwin Davidson Doctor of Quantum Biofeedback

In our test of the VARHOP index in Arizona 2009 we saw significant ability of the SCIO to improve the VARHOP Profile (Voltage, Amperage, Resistance, Hydration, Oxidation and Proton pressure). Here we saw an increase in the Oxidation index. There was also a strong trend of increase in our double blind stand-ups sit-downs while holding the breath test. This was a test of the anaerobic strength of the body or the ability of the subjects to do a physical task while in a deprived oxygen condition. Our hypothesis is that the increase of Oxidation and osmosis proven before would be able to increase oxygenation and endurance. Much experiential study has shown the benefits of SCIO on athletes. We thought that a double blind direct measure of time of holding the breath would be easier to measure and make less opportunity for interference. So in Arizona USA 32 volunteers had the stimulation of the SCIO with pre and post measures of the time they can hold their breath. On certain random sampled volunteers Placebo control was performed. The results show a significant increase in breath retention and thus endurance. The SCIO autofocused cybernetic pulse increase osmosis as well as the VARHOP index. So an increase in body voltage and amperage coupled with an increase with oxidation produces a distinct improvement in oxygenation and endurance.

TVEP reactivity scores to Allersode compounds measured 2013 USA
Written by Darwin Davidson Doctor of Quantum Biofeedback

In this continuing study started in Arizona, USA 2007, we have tested 53 males and 78 females with known allergies using the Transcutaneous Voltammetric Evoked Potential (TVEP) electrical reactivity in the SCIO. The SCIO readings to the allersodes of the know allergies of the subjects was compared to TVEP xroid scores of the non-allergic trivector readings. The reactivity scores of the known allergies were significantly higher than the non-allergic items.

This research adds to the continuing steam of evidence that proves the TVEP reactivity reaction of the SCIO. The SCIO technology is able to test and display allergy reactions.

TVEP reactivity scores to compounds measured update 2013 USA
Written by Darwin Davidson Doctor of Quantum Biofeedback

In this study we tested 65 subjects Transcutaneous Voltammetric Evoked Potential (TVEP) electrical reactivity to five compounds given internally. One was diluted orange juice to act as a placebo and the next four were safe weak dilution of common herbal poisons. Atropine, Convallariana, Aconite and Podophyllum were used because of their toxicity but safety in a 4 x dose. The subjects had a very significant reaction to homeopathic compounds containing the herbals detox and other detox compounds. They also had a no measurable reaction to placebo orange juice after
testing. The placebo test showed no reaction to the sensitive compounds were as the treatment group had significant reactions. This points to the efficacy of the TVEP method.

**Voltammetric Sarcode Hormone Streaming of Testosterone Update 2013 USA**

Written by Darwin Davidson Doctor of Quantum Biofeedback

In our study 28 men (ages 13 to 60) were told to lie down and use their mind to turn themselves on and get an erect penis. They are not allowed to touch or move to do this but only in the mind. The men were connected to the SCIO device and told it would help. The SCIO device was set on placebo for the first round and the SCIO was then operative on visit 2. The time it takes to get an erect penis is an indication of available testosterone. Testosterone is richer in young men and in the morning hours when you get an early morning erection. All tests were done after 12 AM to minimize circadian effects. Thus there was a single blind test of testosterone streaming. In the control measure there was an average of 13.5 minutes and several could not do it within the 15 minute allowed time. The second time with the SCIO on testosterone streaming the time was nearly half with an average of 7.8 minutes and all achieved erections within the 15 allowed period.

Thus it appears that hormone streaming works and the body builders success is real from hormone streaming.

**VARHOPE and EPR Validation Of the SCIO technology -2013 USA**

Written by Darwin Davidson Doctor of Quantum Biofeedback

In 1989 the American FDA registered the EPFX (Electro-Physiological-Feedback-Xrroid) as medical equipment based on research done by the AAQBT and on an equivalency 510k application.

Massive research has been done to further validate the EPFX device. The basic design is still the same since 1989 even though the device has had other names like QXCI, SCIO, Indigo, Eductor, Indigo Pro.

Most recently a series of research projects were done in Europe and America over the last five years. It is well documented that slight oscillating electro stimulation will increase osmosis. We suppose that a harmonious stimulation from an autofocusing cybernetic loop will help to perfect this process, and thus all physiological processes will be improved.

We did simultaneously studies on Sport Performance on performance, breath retention, osmosis, eye hand coordination, addiction urge control, VARHOPE changes, injury repair and EPR (Electro-Physiological-Reactivity which is now called TVEP—Transcutaneous-Voltammetric-Evoked-Potential). We used the same exact protocols in our America study and thus we shared wiring format as well in our presentation.

Working with the American approved IRB of the sponsor and under strict medical supervision we did our studies from 2008 till 2013 in similar fashion to the studies done in Europe. We got similar results showing the SCIO technology valid in these areas.

**Romanian Sports Studies**

Ethical supervision for the following studies has been done by Ethics International Romania, University of Timisoara and the Psychology Faculty of the University of Bucuresti.
The following sports studies have been conducted under the ethical supervision of the aforementioned ethics committees, and have been published in peer reviewed medical journals.

- SKIN SCRATCH
- BASKETBALL FREETHROWS - EYE HAND COORDINATION
- LOW BACK FLEXIBILITY TEST
- HOLD BREATH TEST
- FOOTBALL KICK ACCURACY TEST – EYE FOOT COORDINATION
- HANDBALL THROW ACCURACY - EYE HAND BODY COORDINATION
- DARTS THROW ACCURACY - EYE HAND COORDINATION
- VARHOP MEASURE

2013 Romanian Study of the Indigo Plus Stimulation of Sports Ability

Co Authored by Professor of Medicine Desire’ Dubounet and Hilf Klara MD

STUDY INFORMATION:
SUPERVISING RESEARCHERS: Dr. Danis György MD, Dr. Hilf Klara MD
Licensed Hungarian Medical Doctors
DATE and PLACE: May, 2013, Saut Marie, Romania
SPONSOR:
Sterling Srl / QX World
INSTITUTIONAL MONITOR:
IMUNE / University of Timisoara (Victor Babes University of Medicine) Dr. Bacean Aurel MD

ABSTRACT:
Our previous European and American research has fully shown how the Indigo Plus/Indigo Pro device can increase osmosis with an autofocused micro-current stimulation. Then using a trickle charge system that measures VARHOP (Voltage, Amperage, Resistance, Oxidation and Ph) and can slightly correct aberrations of the body electric. We have shown in previous studies how this has helped a normal population to improve skills in holding breath, eye hand coordination, strength and addiction reduction. This study is designed to test a professional / semi-professional group of athletes for direct sport improvement among base wellness measures. 81 professional / semi-professional athletes age 12 to 45 were tested. 33 basketball players, 28 handball players, and 20 footballers. These athletes were measured for skin osmosis, holding breath, low back flexibility, ability to throw darts, shoot baskets, handball and football skills as well as VARHOP improvement.

And 83 tests results showed a significant improvement versus placebo control measures. Our discussion has shown that these studies have more than proven the claim that the autofocused VARHOP test and intervention is real and enhances sport performance.

INTRODUCTION:

When we apply a micro charge electro-pulse through a biological membrane process, Osmosis increases. Everything in the body depends on osmosis. When Osmosis increases enzymes work better, hormones work better, detox works better, nutrition works better, all cellular functions works better.
Osmosis is the movement of solvent molecules through a selectively permeable membrane into a region of higher solute concentration, aiming to equalize the solute concentrations on the two sides.\textsuperscript{[1]}[2] It may also be used to describe a physical process in which any solvent moves, without input of energy,\textsuperscript{[3]} across a semipermeable membrane (permeable to the solvent, but not the solute) separating two solutions of different concentrations.\textsuperscript{[3]} Although osmosis does not create energy, it does release kinetic energy\textsuperscript{[4]} and can be made to do work,\textsuperscript{[5]} but is a passive process, like diffusion. Everything is made of atoms that never touch each other because of the charge of the outer electrons. The charge of the outer electrons allows for osmosis to occur. When we apply a micro charge electro-pulse through a process, Osmosis increases.

This helps explain the tremendous results the INDIGO patients get on all types of diseases. There is a universal stimulation of osmosis. The INDIGO measures the body level of Voltage, Amperage, Resistance, Hydration, Oxidation and Ph (VARHOP). By stimulating an autofocusing cybernetic harmonic frequency to the body the INDIGO can maximize the osmosis increasing effect without doing any damage. Since it is through Osmosis that the cells bring nutrition and remove toxins, all of life’s processes are improved. Injury improves from the Electrical field stimulation of the INDIGO.

This trickle charge can have maximum benefits in a simple 45 min session. The total change is limited to the body factors of free ions, free minerals, free fatty acids pools, and specifically the membrane potentials of the body.

**SUBJECT AND INVESTIGATOR PROFILE:**

The study took place in Satu Mare, Romania, at the Satu Mare Sports Highschool. Subjects were 60% male and 40% female, aged between 12 and 45 years old semi-professional and professional athletes.

The Romanian Competent Authority granted permission to do studies in Romania, and the Hungarian Ethics Committee has allowed Hilf Klara to do studies in Hungary. (See Appendix)

Our Ethic committee, And written informed consent was approved by all participants.

Medical supervisor:

Dr. Hilf Klára, MD

Placebo officer: Calin Pap

Taflan Andreea, Regulatory Site Manager.

Dates: 20-22 May 2013

There were 4 therapists performing the tests:

Dr. Hilf Klára

Tavman Gabriella- IMUNE Certified Biofeedback Therapist

Pop Gheorghe- IMUNE Certified Biofeedback Therapist
Ruff Krisztian- IMUNE Certified Biofeedback Therapist

METHOD:

SKIN SCRATCH

In this study subjects were hooked to the INDIGO with the INDIGO off and a line drawn firmly on their forearm with a finger nail. Then we count the seconds it takes before the line turns red. This is an indication of osmosis as that the traumatized tissue will have histamine rush in the traumatized tissue of the forearm.

The time it takes indicates the osmosis ability. Norms are 8 to 10 sec. Then the INDIGO is turned on and the line drawn on the forearm again of the subject. In the control group there was an average of 12 seconds and 2 did not have the line appear after the 20 sec deadline. The INDIGO group had an average of 9 seconds and all were under the deadline. This improvement of treatment group over control demonstrates the INDIGO’s ability to increase osmosis thru its auto-focused electrical pulsation. Pre and post tests indicate a possible effect versus placebo group results.

BASKETBALL FREETHROWS - EYE HAND COORDINATION

to test this effect in professional patient blind fashion we got 23 athletic players to do 10 freethrows as a baseline and then 10 more after a 20 min INDIGO treatment and 10 after a placebo treatment. The subjects were blinded as to when the placebo versus therapy happened. VARHOPE measures were also calculated and compared for improvements.

The subject was told to shoot freethrows. The measure was of success of freethrows out of ten.
Pre and post tests indicate a possible effect versus placebo group results.

LOW BACK FLEXIBILITY TEST

Subjects sat on the ground with feet outstretched and knees straight. They reach for their heels and see if they can go past the heels with both hands touching the ground at the longest extension. If they reach their heels they get a zero. If they cannot reach to their heels we use the number of cm shy of the heels as a negative number. If they reach past the heels they get a positive number reflecting the cm they can extend past their heels. Pre and post tests indicate a possible effect versus placebo group results.

HOLD BREATH TEST

Simple measure of how long the subjects can hold their breath in minutes and seconds, pre and post. Pre and post tests indicate a possible effect versus placebo group results.

FOOTBALL KICK ACCURACY TEST – EYE FOOT COORDINATION
The kicker was told to aim at the vertical post from the penalty kick. The measure was of success of hitting the post how many times out of ten.

Pre and post tests indicate a possible effect versus placebo group results.

**HANDBALL THROW ACCURACY - EYE HAND BODY COORDINATION**

The thrower was told to aim at the vertical post from the penalty kick. The measure was of success of hitting the post how many times out of ten.

Pre and post tests indicate a possible effect versus placebo group results.

**DARTS THROW ACCURACY - EYE HAND COORDINATION**

The subject was told to aim at the bullseye and counted the points from the board. The measure was of success of hitting the bullseye and or accumulated points.

Pre and post tests indicate a possible effect versus placebo group results.

**VARHOP MEASURE**

The INDIGO/Indigo Pro electrically measures the body voltage, amperage, hydration, oxidation index, and Ph. The pre and post scores are made as part of the test.
The VARHOPE Scale is made up of the following terms:

V = Voltage where normal is 80-100, below 50 is chronic and from 100-110 is above normal. Voltage is derived directly from the skin electro-potential amplitude.

A = Amperage where normal is 80-100, below 50 is chronic and from 100-110 is above normal. Amperage is calculated from the volume of current over a short period of time coming off of the body.

R = Resistance where normal is 80-100, below 50 is chronic and from 100-110 is above normal. Resistance shows the body’s reaction to the electrical input from the INDIGO.

H = Hydration where normal is 80-100, below 50 is chronic and from 100-110 is above normal. Voltage changes observed during the Calibration process give us a Hydration index.

O = Oxidation where normal is 80-100, below 50 is chronic and from 100-110 is above normal. We get an Oxidation index from comparing maximum and minimum values of changing Amperage.

P = Proton Pressure (commonly interchanged with Proton Balance and refers to pH) where 75 is normal for humans (much like neutral pH is 7.0 and above is more alkaline and below is more acidic.) Changes toward our established norm show an improvement. Proton and Electron pressure (or the charge stability of the system) affects the polarity and the resting potential.

RESULTS:

Treatment Group 58

Subject 1

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<th>Test type</th>
<th>Pre treatment</th>
<th>Post treatment</th>
<th>Improvement Percentage</th>
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<tr>
<td>Skin scratch (seconds)</td>
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<td>Low back extension (cm)</td>
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### Subject 3

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Subject 5

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Subject 6

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Subject 7

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**Subject 8**

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**Subject 9**

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Subject 12

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Subject 13

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Subject 15

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Subject 18

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Subject 19

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Football (how many out of 10)

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**Subject 26**

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Subject 28

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<td>Hold Breath (seconds)</td>
<td>Pre treatment</td>
<td>Post treatment</td>
<td>Percentage of improvement</td>
</tr>
<tr>
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<tr>
<td>36</td>
<td>73</td>
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<tr>
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<td>14</td>
<td>36</td>
<td>157%</td>
</tr>
<tr>
<td>Basketball (how many out of 10)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Handball (how many out of 10)</td>
<td></td>
<td></td>
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<td>Football (how many out of 10)</td>
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Subject 29

<table>
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<th>Percentage of improvement</th>
</tr>
</thead>
<tbody>
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<td>Skin scratch (seconds):</td>
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<td>34</td>
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<td>6</td>
<td>-33%</td>
</tr>
<tr>
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<td>37</td>
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<td>62%</td>
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<td>45</td>
<td>30</td>
<td>-33%</td>
</tr>
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<td></td>
</tr>
<tr>
<td>Handball (how many out of 10)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Football (how many out of 10)</td>
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<td>8</td>
<td>14%</td>
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Subject 30

<table>
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<th>Percentage of improvement</th>
</tr>
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<tbody>
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<td>Skin scratch (seconds):</td>
<td>19</td>
<td>14</td>
<td>26%</td>
</tr>
<tr>
<td>Low back extension (cm):</td>
<td>10</td>
<td>10</td>
<td>0%</td>
</tr>
<tr>
<td>Hold Breath (seconds)</td>
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<td>67</td>
<td>-30%</td>
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160
<table>
<thead>
<tr>
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<th>9</th>
<th>33</th>
<th>267%</th>
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<tr>
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</tr>
<tr>
<td>Handball (how many out of 10)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Football (how many out of 10)</td>
<td>8</td>
<td>8</td>
<td>0%</td>
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**Subject 31**

<table>
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</tr>
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<td>-60%</td>
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<td>-65%</td>
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<td>98</td>
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<td>27</td>
<td>-21%</td>
</tr>
<tr>
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</tr>
<tr>
<td>Handball (how many out of 10)</td>
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<td></td>
</tr>
<tr>
<td>Football (how many out of 10)</td>
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**Subject 32**

<table>
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</tr>
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<tbody>
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<td>Skin scratch (seconds):</td>
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<td>22%</td>
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<tr>
<td>Low back extension (cm):</td>
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<td>56%</td>
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<tr>
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<td>39</td>
<td>30%</td>
</tr>
<tr>
<td>Darts (how many points out of 3)</td>
<td>30</td>
<td>28</td>
<td>-7%</td>
</tr>
</tbody>
</table>
Basketball (how many out of 10) |  |  |  |
| Handball (how many out of 10) |  |  |  |
| Football (how many out of 10) | 8 | 10 | 25% |

**Subject 33**

<table>
<thead>
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<th>Pre treatment</th>
<th>Post treatment</th>
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</tr>
</thead>
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<td>16</td>
<td>33%</td>
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<tr>
<td>Low back extension (cm):</td>
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<td>100%</td>
</tr>
<tr>
<td>Hold Breath (seconds)</td>
<td>56</td>
<td>57</td>
<td>2%</td>
</tr>
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<td>Darts (how many points out of 3)</td>
<td>17</td>
<td>80</td>
<td>371%</td>
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<td></td>
<td></td>
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<tr>
<td>Handball (how many out of 10)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Football (how many out of 10)</td>
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**Subject 34**

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<td>100%</td>
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<td>Hold Breath (seconds)</td>
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<td>42</td>
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</tr>
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<td>66</td>
<td>230%</td>
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<td></td>
<td></td>
</tr>
<tr>
<td>Test type</td>
<td>Pre treatment</td>
<td>Post treatment</td>
<td>Percentage of improvement</td>
</tr>
<tr>
<td>---------------------------------</td>
<td>---------------</td>
<td>----------------</td>
<td>--------------------------</td>
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<td>100%</td>
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</tr>
<tr>
<td>Handball (how many out of 10)</td>
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<td></td>
<td></td>
</tr>
<tr>
<td>Football (how many out of 10)</td>
<td>9</td>
<td>9</td>
<td>0%</td>
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<table>
<thead>
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<th>Pre treatment</th>
<th>Post treatment</th>
<th>Percentage of improvement</th>
</tr>
</thead>
<tbody>
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<td>0%</td>
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<tr>
<td>Low back extension (cm):</td>
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<td>100%</td>
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<td>Hold Breath (seconds)</td>
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<td>4%</td>
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<td>-7%</td>
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<tr>
<td>Handball (how many out of 10)</td>
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<td></td>
<td></td>
</tr>
<tr>
<td>Football (how many out of 10)</td>
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<td>9</td>
<td>13%</td>
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**Subject 37**

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</tr>
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<tr>
<td>Low back extension (cm):</td>
<td>2</td>
<td>4</td>
<td>100%</td>
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<td>-37%</td>
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</tr>
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<td>Handball (how many out of 10)</td>
<td>5</td>
<td>9</td>
<td>80%</td>
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**Subject 38**

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</tr>
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<td>0%</td>
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<td>Hold Breath (seconds)</td>
<td>34</td>
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<td>6</td>
<td>-83%</td>
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<td>Handball (how many out of 10)</td>
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<td>10</td>
<td>11%</td>
</tr>
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Subject 39

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<td>10</td>
<td>11%</td>
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Subject 40

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<td>-17%</td>
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<tr>
<td>Handball (how many out of 10)</td>
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<td>9</td>
<td>29%</td>
</tr>
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Subject 41

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<td>Post treatment</td>
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</tr>
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<td>Low back extension (cm):</td>
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**Subject 42**

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<td>Low back extension (cm):</td>
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</tr>
<tr>
<td>Hold Breath (seconds)</td>
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<td>35</td>
<td>13%</td>
</tr>
<tr>
<td>Darts (how many points out of 3)</td>
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<td>9</td>
<td>13%</td>
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**Subject 43**

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</tr>
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<tr>
<td>Test type</td>
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<td>Post treatment</td>
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<td>17%</td>
</tr>
<tr>
<td>Low back extension (cm):</td>
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<td>10</td>
<td>0%</td>
</tr>
<tr>
<td>Hold Breath (seconds)</td>
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<td>37</td>
<td>6%</td>
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<td>Handball (how many out of 10)</td>
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<td>10</td>
<td>0%</td>
</tr>
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**Subject 44**

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<tr>
<td>Low back extension (cm):</td>
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<td>14%</td>
</tr>
<tr>
<td>Hold Breath (seconds)</td>
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<td>14%</td>
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<tr>
<td>Darts (how many points out of 3)</td>
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<td>12</td>
<td>14%</td>
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<tr>
<td>Basketball (how many out of 10)</td>
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<td>12</td>
<td>14%</td>
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<td>14%</td>
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<td>14%</td>
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<tr>
<td>Subject 46</td>
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<td>Post treatment</td>
<td>Percentage of improvement</td>
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<td>20%</td>
</tr>
<tr>
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<td>-3%</td>
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<td>30</td>
<td>100%</td>
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<td>0%</td>
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<td>Handball (how many out of 10)</td>
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</tr>
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Subject 47

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Subject 48

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Subject 53

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### Subject 54

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### Subject 55

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### Subject 57

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### Subject 58

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### PLACEBO Group 15

**Subject 1**

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**Subject 2**

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<td>Handball (how many out of 10)</td>
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**Subject 8**

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<td>6</td>
<td>14%</td>
</tr>
<tr>
<td>Low back extension (cm):</td>
<td>12</td>
<td>11</td>
<td>-8%</td>
</tr>
<tr>
<td>Hold Breath (seconds)</td>
<td>43</td>
<td>35</td>
<td>-19%</td>
</tr>
<tr>
<td>Darts (how many points out of 3)</td>
<td>51</td>
<td>49</td>
<td>-4%</td>
</tr>
<tr>
<td>Basketball (how many out of 10)</td>
<td>7</td>
<td>6</td>
<td>-14%</td>
</tr>
<tr>
<td>Handball (how many out of 10)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Football (how many out of 10)</td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Subject 12</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Test type</strong></td>
</tr>
<tr>
<td>Skin scratch (seconds):</td>
</tr>
<tr>
<td>Low back extension (cm):</td>
</tr>
<tr>
<td>Hold Breath (seconds)</td>
</tr>
<tr>
<td>Darts (how many points out of 3)</td>
</tr>
<tr>
<td></td>
</tr>
<tr>
<td>----------------------</td>
</tr>
<tr>
<td><strong>Skin scratch (seconds):</strong></td>
</tr>
<tr>
<td><strong>Low back extension (cm):</strong></td>
</tr>
<tr>
<td><strong>Hold Breath (seconds)</strong></td>
</tr>
<tr>
<td><strong>Darts (how many points out of 3)</strong></td>
</tr>
<tr>
<td><strong>Basketball (how many out of 10)</strong></td>
</tr>
<tr>
<td><strong>Handball (how many out of 10)</strong></td>
</tr>
<tr>
<td><strong>Football (how many out of 10)</strong></td>
</tr>
</tbody>
</table>

**Subject 14**

<table>
<thead>
<tr>
<th></th>
<th>Pre treatment</th>
<th>Post treatment</th>
<th>Percentage of improvement</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Skin scratch (seconds):</strong></td>
<td>9</td>
<td>14</td>
<td>-56%</td>
</tr>
<tr>
<td><strong>Low back extension (cm):</strong></td>
<td>10</td>
<td>7</td>
<td>-30%</td>
</tr>
<tr>
<td><strong>Hold Breath (seconds)</strong></td>
<td>38</td>
<td>33</td>
<td>-13%</td>
</tr>
<tr>
<td><strong>Darts (how many points out of 3)</strong></td>
<td>29</td>
<td>28</td>
<td>-3%</td>
</tr>
<tr>
<td><strong>Basketball (how many out of 10)</strong></td>
<td>9</td>
<td>6</td>
<td>-33%</td>
</tr>
</tbody>
</table>
### Subject 15

<table>
<thead>
<tr>
<th>Test type</th>
<th>Pre treatment</th>
<th>Post treatment</th>
<th>Percentage of improvement</th>
</tr>
</thead>
<tbody>
<tr>
<td>Skin scratch (seconds):</td>
<td>9</td>
<td>8</td>
<td>-11%</td>
</tr>
<tr>
<td>Low back extension (cm):</td>
<td>6</td>
<td>7</td>
<td>-17%</td>
</tr>
<tr>
<td>Hold Breath (seconds)</td>
<td>25</td>
<td>23</td>
<td>-8%</td>
</tr>
<tr>
<td>Darts (how many points out of 3)</td>
<td>15</td>
<td>1</td>
<td>-93%</td>
</tr>
<tr>
<td>Basketball (how many out of 10)</td>
<td>6</td>
<td>6</td>
<td>0%</td>
</tr>
<tr>
<td>Handball (how many out of 10)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Football (how many out of 10)</td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

Treatment 58
Placebo 15

Treatment group all tests improvement: +35.23%
Placebo Group all test improvement: -19.88%
Skin scratch (real): +13.85%
Skin scratch (placebo): -15.88%
Low back extension (real): +41.32%
Low back extension (placebo): -9.12%
Hold Breath (real): +18.39%
Hold Breath (placebo): -14.43%
Darts (real) +45.23%
<table>
<thead>
<tr>
<th>Activity</th>
<th>Treatment Improvement</th>
</tr>
</thead>
<tbody>
<tr>
<td>Darts (placebo)</td>
<td>-28.06%</td>
</tr>
<tr>
<td>Basketball (real)</td>
<td>+34.19</td>
</tr>
<tr>
<td>Basketball (placebo)</td>
<td>-14.25%</td>
</tr>
</tbody>
</table>

**Treatment 58**

**Placebo 15**

- Treatment group all tests improvement: +35.23%
- Placebo Group all test improvement: -19.88%
- Skin scratch (real): +13.85%
- Skin scratch (placebo): -15.88%
- Low back extension (real): +41.32%
- Low back extension (placebo): -9.12%
- Hold Breath (real): +18.39%
- Hold Breath (placebo): -14.43%
- Darts (real): +45.23%
- Darts (placebo): -28.06%
- Basketball (real) 17 people: +55.05%
- Basketball (placebo) 16 people: -17.32%
- Handball (real) 24 people: 30.91%
- Handball (placebo) 4 people: 0%
- Football (real) 16 people: 25.25%
- Football (placebo) 4 people: 0%

**Treatment 58**

**Placebo 15**

- Treatment group all tests improvement: +35.23%
- Placebo Group all test improvement: -19.88%
- Skin scratch (real): +13.85%
- Skin scratch (placebo): -15.88%
Low back extension (real): +41.32%
Low back extension (placebo): -9.12%
Hold Breath (real): +18.39%
Hold Breath (placebo): -14.43%
Darts (real): +45.23%
Darts (placebo): -28.06%
Basketball (real) 17 people: +55.05%
Basketball (placebo) 15 people: -17.32%
Handball (real) 24 people: 30.91%
Handball (placebo) 4 people: 0%
Football (real) 16 people: 25.25%
Football (placebo) 4 people: 0%

VARHOP PRE POST MEASURES

VARHOPE TREATMENT GROUP IMPROVEMENT:
V 14.07%
A 19.23%
R 15.36%
H 21.44%
O 18.24%
P 4.56%

VARHOPE PLACEBO IMPROVEMENT:
V 0.01%
A 0.01%
R 0.10%
H -0.03%
O -0.10%
P -0.20%

DISCUSSION:
After selling over 35,000 equivalent such devices without any cases of any significant risks, we can unequivocally say that this device is safe. Our safety risk analysis and the ISO safety tests show there is insignificant risk at best from the INDIGO/Indigo Pro.

This study along with our 2012 European and 2013 American research has fully shown how the Indigo Plus/Indigo Pro device can increase osmosis with an autofocused micro-current stimulation. Then by using a trickle charge system that measures VARHOP (Voltage, Amperage, Resistance, Oxidation and Ph) and can slightly correct aberrations of the body electric. We have shown in this study how this has helped a normal population to improve skills in holding breath, eye hand coordination, strength and addiction reduction versus a placebo group. The professional / semi-professional group of athletes demonstrated direct sport improvement among base wellness measures.

Results showed a significant improvement versus placebo control measures. Our discussion has shown that these studies have more than proven the claim that the autofocused VARHOP test and intervention is real and enhances sport performance.

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28. 978-615-5169-13-7 Injury and Sport Medicine, IMUNE Press
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4. VARHOPE (Voltage, Amperage, Resistance, Oxidation, Hydration, Proton and Electron pressure, the body electric's vital signs), 978-615-5169-17-5, IMUNE Press 2011
5. VARHOPE and STRESS, Vol.XXV.1. 2010 VARHOPE and Stress ISSN 2041-4293
6. INDIGO's ability to increase Body Wellness, Vol.XXXV.1. 2010 VARHOPE and Stress ISSN 2041-4293
7. VARHOPE Large scale study, Vol.XXXV.1. 2010 VARHOPE and Stress ISSN 2041-4293
8. Varhope Improvements in a Clinical Setting, Vol.XXXV.1. 2010 VARHOPE and Stress ISSN 2041-4293
9. Large Scale Studies of the INDIGO, Vol.XXIII.1. - 2008 The Large Scale Study of the INDIGO ISSN 2041-4293
APPENDIX:

MINISTERUL SĂNĂTĂȚII
DIRECȚIA PENTRU INFRASTRUCTURĂ SANITARĂ,
LOGISTICĂ, ACHIZIȚII ŞI DISPOZITIVE MEDICALE
COMPARTIMENT DISPOZITIVE MEDICALE
Str. Cristian Popişteanu Nr. 1-3, Sector 1, 010024, BUCUREȘTI
Telefon: 021 307 25 82; Fax: 021 307 25 85; E-mail: apmed@ms.ro

Nr. 20474/31.03.2010
Referitor la solicitarea dumneavoastră transmisă prin e-mail și înregistrată la Ministerul Sănătății cu nr. 20474/23.03.2010 vă comunicăm următoarele:
- investigația clinică pentru dispozitive medicale este reglementată de Hotărârea Guvernului nr.54/2009 privind condițiile introducerii pe piață a dispozitivelor medicale, publicată în Monitorul Oficial nr. 94 din 17 februarie 2009 și Ordinul ministrului sănătății nr. 792/2006 privind desfășurarea procedurii de investigație clinică și a procedurii de evaluare a performanței pentru dispozitive medicale, publicat în Monitorul Oficial Nr. 595 din 10 iulie 2006;
- cererea pentru emiterea autorizației privind desfășurarea procedurii de investigație clinică trebuie însoțită de documentele prevăzute în anexa nr. 2 din OMS nr 792/2006 și aceste documente se depun la registratura Ministerului Sănătății din str. Cristian Popişteanu nr.1-3, sector 1, 010024, București;
- Ministerul Sănătății nu percepe taxe pentru emiterea autorizației menționată mai sus;
- potrivit HG 54/2009 se va prezenta punctul de vedere al comitetului de etică implicat.

Cu stimă,
DIRECTOR,
Ing. Alexandru STERIU
Ref.Specialitate Margareta Mihalache

ORDIN Nr. 792 din 29 iunie 2006
privind desfășurarea procedurii de investigație clinică și a procedurii de evaluare a performanței pentru dispozitive medicale

Text în vigoare începând cu data de 22 aprilie 2009
REALIZATOR: COMPANIA DE INFORMATICĂ NEAMȚ

Text actualizat prin produsul informatic legislativ LEX EXPERT în baza actelor normative modificatoare, publicate în Monitorul Oficial al României, Partea I, până la 22 aprilie 2009.

Act de bază

#B: Ordinul ministrului sănătății publice nr. 792/2006

Acte modificatoare

#M1: Ordinul ministrului sănătății nr. 465/2009

Modificările și completările efectuate prin actul modificator sunt scrise cu font italic. În fața fiecărei modificări sau completări este indicat actul normativ care a efectuat modificarea sau completarea respectivă, în forma #M1.

#B

Având în vedere prevederile art. 9 și 11 din Legea nr. 176/2000 privind dispozitivele medicale, republicată, ale art. 35 din Hotărârea Guvernului nr. 911/2005 privind stabilirea condițiilor de introducere pe piață și de punere în funcțiune a dispozitivelor medicale, ale art. 19 din Hotărârea Guvernului nr. 798/2003 privind stabilirea condițiilor de introducere pe piață și de utilizare a dispozitivelor medicale pentru diagnostic in vitro și ale art. 23 din Hotărârea Guvernului nr.
ART. 1
(1) Prezentul ordin stabileşte condiţiile şi modul de desfăşurare a procedurii de investigaţie clinică şi a procedurii de evaluare a performanţei pentru dispozitive medicale.
(2) Prevederile prezentului ordin se aplică dispozitivelor medicale, dispozitivelor medicale implantabile active şi dispozitivelor medicale pentru diagnostic in vitro, denumite în continuare dispozitive medicale.

ART. 2
În prezentul ordin sunt aplicabile definiţiile şi procedurile stipulate în Hotărârea Guvernului nr. 911/2005 privind stabilirea condiţiilor de introducere pe piaţă şi de punere în funcţiune a dispozitivelor medicale, în Hotărârea Guvernului nr. 798/2003 privind stabilirea condiţiilor de introducere pe piaţă şi de utilizare a dispozitivelor medicale pentru diagnostic in vitro şi în Hotărârea Guvernului nr. 344/2004 privind stabilirea condiţiilor de introducere pe piaţă şi/sau de punere în funcţiune a dispozitivelor medicale implantabile active, cu modificările ulterioare.

ART. 3
Autorizaţia pentru desfăşurarea procedurii de investigaţie clinică şi a procedurii de evaluare a performanţei pentru dispozitive medicale este eliberată de structura de specialitate din cadrul Ministerului Sănătăţii Publice, cu avizul comisiilor de specialitate ale Ministerului Sănătăţii Publice.

ART. 4
Producătorii sau reprezentanţii lor autorizaţi care vor să desfăşoare procedura de investigaţie clinică sau procedura de evaluare a performanţei pentru dispozitive medicale trebuie să notifice în scris structura de specialitate din cadrul Ministerului Sănătăţii Publice.

ART. 5
(1) Pentru aprobarea desfăşurării procedurii de investigaţie clinică şi a procedurii de evaluare a performanţei pentru dispozitive medicale, producătorul sau reprezentantul său autorizat înaintează o cerere la structura de specialitate din cadrul Ministerului Sănătăţii Publice, al cărei model este prevăzut în anexa nr. 1.
(2) Cererea prevăzută la alin. (1 va fi însoţită de documentele prevăzute în anexa nr. 2, după caz.

ART. 6
(1) În baza cererii şi a documentelor prevăzute la art. 5 alin. (2, structura de specialitate din cadrul Ministerului Sănătăţii Publice autorizează:
a) începerea investigaţiei clinice, potrivit prevederilor art. 36 şi 37 din Hotărârea Guvernului nr. 911/2005 sau ale art. 24 din Hotărârea Guvernului nr. 344/2004;
b) începerea evaluării performanţei, potrivit art. 19 din Hotărârea Guvernului nr. 798/2003. 
(2) Pentru începerea procedurii de investigaţie clinică şi a procedurii de evaluare a performanţei pentru dispozitive medicale, structura de specialitate din cadrul Ministerului Sănătăţii Publice eliberează autorizaţia al cărei model este prevăzut în anexa nr. 3, respectiv în anexa nr. 4, după caz.

ART. 7
Investigaţiile clinice trebuie să se desfăşoare în concordanţă cu prevederile anexei nr. 10 la Hotărârea Guvernului nr. 911/2005 sau ale anexei nr. 7 la Hotărârea Guvernului nr. 344/2004, iar evaluarea performanţei, potrivit prevederilor anexei nr. 8 la Hotărârea Guvernului nr. 798/2003.
ART. 8
Producătorul sau reprezentantul său autorizat trebuie să păstreze la dispoziția Ministerului Sănătății Publice raportul privind investigația clinică prevăzut la pct. 2 subpct. 2.3.7 din anexa nr. 10 la Hotărârea Guvernului nr. 911/2005 sau raportul prevăzut la subpct. 2.3.7 din anexa nr. 7 la Hotărârea Guvernului nr. 344/2004, respectiv raportul privind evaluarea performanței prevăzut la pct. 3 din anexa nr. 8 la Hotărârea Guvernului nr. 798/2003.

ART. 9
Potrivit art. 56 lit. b) și d) din Hotărârea Guvernului nr. 911/2005, nerespectarea prevederilor art. 4, 5, 7 și 8 din prezentul ordin constituie contravenție și se sancționează cu amendă de la 2.500 lei (RON) la 5.000 lei (RON).

#M1

ART. 10 *** Abrogat

#B

ART. 11
Anexele nr. 1 - 4 fac parte integrantă din prezentul ordin.

ART. 12
Structura de specialitate din cadrul Ministerului Sănătăţii Publice, direcţiile cu atribuţii în activitatea de asistenţă medicală din Ministerul Sănătăţii Publice, comisiile de specialitate ale Ministerului Sănătăţii Publice, precum şi toate unităţile sanitare vor aduce la îndeplinire prevederile prezentului ordin.  

ART. 13

Prezentul ordin va fi publicat în Monitorul Oficial al României, Partea I.

ANEXA 1

CERERE pentru eliberarea autorizaţiei de desfăşurare a procedurii de investigaţie clinică sau a procedurii de evaluare a performanţei pentru dispozitivele medicale

Nr. ........... din .................

Câtre Ministerul Sănătăţii Publice

Structura de specialitate în domeniul dispozitivelor medicale

Producătorul/Reprezentantul autorizat de dispozitive medicale ........................................ , cu sediul în ...................., telefon .........., fax ..........., e-mail................., reprezentat de ........................................, solicit autorizarea desfăşurării procedurii de investigaţie clinică sau a procedurii de evaluare a performanţei pentru dispozitivele medicale:

..............................................................................................................................................

Anexez documentele prevăzute în lista verificărilor din anexa nr. 2.

Data .......... Semnătura ..............

ANEXA 2

INVESTIGAŢIA CLINICĂ/EVALUAREA PERFORMANŢEI

LISTA VERIFICĂRILOR

1. INFORMAŢII GENERALE

1.1. Numele producătorului/reprezentantului autorizat, adresa, telefonul, date pentru contactare în vederea comunicării

1.2. Dacă este prima cerere pentru investigaţie/evaluare sau resolicitare

1.3. Dacă este resolicitare legată de acelaşi dispozitiv, numărul referinţei sau referinţelor şi datele anterioare rezultate din cele mai recente investigaţii

1.4. Alte țări membre participante la investigaţia clinică/evaluarea performanţei ca parte a studiului multinaţional/in multicentre

1.5. Declaraţie semnată din care să rezulte că dispozitivul în cauză este conform cu cerinţele esenţiale, cu excepţia acelor aspecte care fac obiectul investigaţiei şi în conformitate cu care au fost luate toate măsurile pentru protejarea siguranţei şi sănătăţii pacientului.

2. DATE CARE PERMIT IDENTIFICAREA DISPOZITIVULUI

2.1. Numele comercial al dispozitivului

2.2. Numele generic al dispozitivului

2.3. Numele de model al dispozitivului

2.4. Numărul modelului, dacă există.

3. ALTE DETALII PRIVIND DISPOZITIVUL

3.1. Clasificare
3.2. Descrierea completă a dispozitivului, inclusiv o listă a accesoriilor, principiile de operare și desenele de ansamblu și ale componentelor de bază, împreună cu o scurtă descriere a dispozitivelor destinate să fie folosite în combinație, în scopul investigației/evaluării

3.3. Identificarea oricăror caracteristici de proiectare care sunt diferite față de cele ale produsului similar introdus anterior pe piață

3.4. Detaliul privind caracteristicile dispozitivelor noi sau netestate anterior, care să prevadă, unde este posibil, funcția și principiile de operare

3.5. Rezumat al experiențelor cu orice dispozitiv asemănător, făcut de același producător, care să conțină data când a fost introdus pe piață și o prezentare a problemelor legate de performanțe, incidente și măsurile luate pentru rezolvarea acestora

3.6. Analizele beneficu-risc, care să cuprindă identificarea hazardului și estimarea riscurilor legate de fabricație (inclusiv cele referitoare la alegerea dispozitivului, a materialelor și a softului) și de utilizare a dispozitivului, precum și descrierea măsurilor care trebuie să fie luate pentru minimalizarea sau eliminarea riscurilor identificate

3.7. Rezumat și analize ale testelor preclinice și ale datelor experimentale, care să cuprindă rezultatele calculurilor de proiectare, testelor mecanice, testelor electrice, testelor de validare a softului, verificarea siguranței în funcționare și orice performanță sau teste de siguranță efectuate pe animale

3.8. Descrierea materialelor care vin în contact cu organismul uman, motivul pentru care au fost alese astfel de materiale și standardul aplicabil, dacă este relevant

3.9. Descrierea biocompatibilității și siguranței biologice și modul în care a fost abordată astfel încât să cuprindă și identificarea riscurilor și hazardului legat de utilizarea dispozitivului

3.10. Identificarea oricăror componente farmacologice ale dispozitivului, cu descrierea scopului propus și experiența anterioară în utilizarea acestor substanțe

3.11. Principiul de proiectare și diagramele de funcționare, inclusiv materiale și biomateriale, însoțite de descrierea și explicațiile necesare pentru a înțelege proiectul

3.12. Descrierea softului, logica și condițiile de utilizare, dacă este cazul

3.13. Metoda de sterilizare și validare (metode, justificare)

3.14. Identificarea oricăror țesuturi de origine animală încorporate în dispozitiv și informații privind sursa țesuturilor ținute de fabricație; detaliul privind validarea procedurilor de fabricație utilizate pentru reducerea sau inactivarea agenților neconvenționali

3.15. Identificarea oricăror condiții speciale de fabricație ca cerințe speciale și modul în care trebuie să fie îndeplinite aceste cerințe

3.16. Lista standardelor armonizate aplicabile în întregime sau parțial ori descrierea soluțiilor adoptate pentru îndeplinirea cerințelor esențiale ale directivei, dacă standardele de referință nu sunt aplicabile în întregime

3.17. Instrucțiuni de utilizare

3.18. Ce măsuri au fost luate de producător - dacă există - pentru reconstruirea (reproiectarea) dispozitivului (aplicabilă în cazul dispozitivelor implantabile, dispozitivelor cu utilizări multiple) și prevenirile ulterioară a unei utilizări neautorizate.

4. PLANUL INVESTIGAȚIEI CLINICE/EVALUĂRII PERFORMANȚEI

Informații generale
4.1. Numele, calificările, funcția profesională, adresele investigatorilor clinici, ale investigatorului coordonator, dacă este cazul, din multicentrele de investigație clinică

4.2. Precizări privind experiența și calificarea necesare pentru utilizarea dispozitivului investigat

4.3. Numele, adresele și instituțiile în care se vor desfășura investigațiile
4.4. Copie de pe opinia Comitetului de etică, cuprinzând informații cu privire la faptul dacă documentele de studiu au fost aprobate parțial sau total sau aprobate cu unele condiții, dacă este cazul
4.5. Copie de pe consimțământul avizat al pacientului
4.6. Copie de pe documentul privind modul de despăgubire a pacientului în cazul deteriorării stării lui de sănătate în urma investigației clinice
4.7. Sumarul literaturii științifice de referință care a stat la baza studiului, cu analiză și bibliografie, dacă este cazul.
5. PLANUL INVESTITIIEI CLINICE/EVALUĂRII PERFORMANȚEI
Planul și parametrii investigației
5.1. Scopul și obiectivele investigației
5.2. Planul investigației, de exemplu dacă este prevăzută utilizarea unui grup controlat de pacienți - cu motivația corespunzătoare; dacă s-a luat în considerare concomitent tendința datorată evoluției naturale a bolii față de efectele tratamentului
5.3. Numărul de pacienți - justificare
5.4. Durata studiului, cu precizarea datelor de început și sfârșit și perioada de urmărire a realizării finale a investigației - justificare
5.5. Populația studiată
5.6. Criterii de selecție a pacienților
5.7. Criterii de includere și excludere
5.8. Criterii pentru retragerea din studiu
5.9. Descrierea și justificarea incidentelor cauzate de procedurile invazive care nu sunt de natură medicală
5.10. Descrierea metodelor generale de diagnostic sau a condiției medicale de tratament pentru care a fost propusă investigația.
6. PLANUL INVESTITIIEI CLINICE/EVALUĂRII PERFORMANȚEI
Colectarea datelor/Analize/Statistici
6.1. Descrierea rezultatelor finale pentru a demonstra performanța și siguranța utilizării dispozitivului și datele înregistrate pentru realizarea scopului final, metoda de urmărire a pacienților, evaluarea și monitorizarea în timpul investigației
6.2. Descrierea procedurilor și detalii privind înregistrarea și raportarea incidentelor, inclusiv detalii ale incidentelor deosebite care trebuie să fie raportate autorității competente
6.3. Descrierea și justificarea datelor statistice, metodei și procedurilor analitice.
7. ALTE PREVEDERI
Compensație în cazul agravării stării de sănătate a pacienților
ANEXA 3
ROMÂNIA
MINISTERUL SĂNĂTĂȚII PUBLICE
Structura de specialitate în domeniul dispozitivelor medicale
AUTORIZAȚIE PENTRU INVESTITIIEA CLINICĂ
a dispozitivelor medicale
Nr. ........ din ............ În conformitate cu prevederile Hotărârii Guvernului nr. 168/2005 privind organizarea și funcționarea Ministerului Sănătății, cu modificările și completările ulterioare, ale Hotărârii Guvernului nr. 911/2005 privind stabilirea condițiilor de introducere pe piață și de punere în funcțiune a dispozitivelor medicale, ale Hotărârii Guvernului nr. 344/2004 privind stabilirea condițiilor de introducere pe piață și/sau de punere în funcțiune a dispozitivelor medicale implantabile active, cu modificările ulterioare, și în baza documentației înaintate, Ministerul Sănătății Publice autorizează desfășurarea investigației clinice pentru dispozitivul medical:

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(denumirea, tipul)
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Orice modificare a condițiilor stabilite prin reglementările Ministerului Sănătății Publice, care au stat la baza autorizării, atrage anularea prezentului document.

Director,

.................................

ANEXA 4
ROMÂNIA
MINISTERUL SĂNĂTĂŢII PUBLICE
Structura de specialitate în domeniul dispozitivelor medicale
AUTORIZAŢIE PENTRU EVALUAREA PERFORMANŢEI dispozitivelor medicale
Nr. ..... din ............
În conformitate cu prevederile Hotărârii Guvernului nr. 168/2005 privind organizarea și funcționarea Ministerului Sănătății, cu modificările și completările ulterioare, ale Hotărârii Guvernului nr. 798/2003 privind stabilirea condițiilor de introducere pe piață și de punere în funcțiune a dispozitivelor medicale pentru diagnostic in vitro și în baza documentației înaintate, Ministerul Sănătății Publice autorizează evaluarea performanței pentru dispozitivul medical:

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Producător: .....................................
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Director,

.................................

1932 őszén letesztelte a paprika C-vitamin tartalmát. Bebizonyosodott, hogy a paprika egy nagyon gazdag forrása a C-vitaminnak, és a kínálat nem volt gond - Szeged volt a Magyarország paprika fővárosa. Szent-Györgyi azonnal mozsósította a munkatársait, és megkezdték a paprika nagyszabású C-vitaminkitermelését. Egy héten belül több mint három font tiszta kristályos anyagot, C-vitamint készítettek.

Szent-Györgyöt kutatásai az izomszövet légzésével kapcsolatban, arra a kérdésre vezette, hogy hogyan mozog az izom. 1939-ben a kutatók arról számoltak be, hogy az izom fehérjéi kölcsönhatásba léphetnek és megoszthatják az ATP-t. Bár 1929-ben fedezték fel ATP-t, még nem azonosították fő energiaforrásként a sejtekben (hatalmas energia szabadul fel, amikor a foszfát, kötésekre van felosztva). Szent-Györgyi indoklása szerint a miozin-ATP kölcsönhatás megmagyarázhatja az izom mozgását.

Szent-Györgyi izomműködésről írt munkáját rövid, elegánsan írt sorozatú könyvekben összegzi.

Szent-Györgyi András, Albert fiatalabb unokatestvére, és felesége Éva, felfedezték a miozin alegységeit ("meromyosin-okat"), és elkezdték az izomfehérjék működésének elemzését több elektronikus szinten. Szent-Györgyi és más kollégák úttörő munkát végeztek az izomszövet elektronikus szinten való elemzésével kapcsolatban.

Legtöbb termés a paprikából Magyarországon volt, javasoljuk, hogy tanulmányozza a paprika hatását, és az elektromágneses stim (szabványos TENS készülékkel) elemzését. 1. Izom növelése diéta állat
2. Paprika a diétában és az Elektromágneses stim (szabványos TENS készülékkel)
3. TENS készülék önmagában
4. Kontroll csoport

5 és 10 témára számítunk minden csoportban. És, egy egy hónapos időszakot vennénk figyelembe a fejlődés során.

Előtte és utána mérések: vér, hormon szint, tesztoszteron, humán növekedési hormon, kortizol, lélegzet visszatartás, szem és kéz koordináció, fájdalom tolerancia, az érzelmi állapot összpontosítása az ingerlékenységre, fogyás, az erő és izom tömege.

Ezzel a tanulmányval, szeretnénk felkelteni az érdeklődést C-vitaminnal kapcsolatban, és javítani a magyar paprika exporton.
Szent-Györgyi kiadványok

Az oxidáció, erjesztés, vitaminok, egészség, és a betegség (1940)
Bioenergetika (1957)
Bevezetés egy szubmolekuláris biológiába (1960)
Az izommal kapcsolatos tanulmányok az Orvostudományi Vegyészeti Intézettől

Referenciák.


3. ^ Dr. Czeizel E. : Az érték még mindig bennünk van, 172 oldal, Akadémiai kiadó, Budapest
IMUNE's medical text books are a part of the curriculum of the 'Victor Babes' Medical University in Timisoara, Romania, and have been for the last 5 years, since 2009.

As a general rule, a medical device study cannot be n a university approved medical text book until it has been in a peer reviewed medical journal for at least 5 years. Our studies have been in peer reviewed medical journals for more than 7 years thus our studies are included in the following medical text books of the University of Timisoara:

- **Energetic Medicine - Science over Convention**
- **VARHOPE (Voltage, Amperage, Resistance, Oxidation, Hydration, Proton and Electron pressure, the body electric's vital signs)**
- **TVEP and Medication Testing (the research)**
- **Stress as THE Medical Concern**
- **Electro-Physiology-Feedback - Measures of Interstitial Fluids**
- **Medical Research Validation of the SCIO**
- **SOC Index and the Evidence for Lifestyle Medicine**
- **The Body Electric Simplified**
- **To Be a Professional Biofeedback and Energetic Medicine Therapist**
- **Today’s Modern Research in Electro Stim and the Eductor**
- **VASO-VAGAL Reaction what you need to know to operate the SCIO**

Since 2009, University of Timisoara has been the host of a 4 module Postgraduate Study on Neuroanatomy, Neuroelectrophysiology and Biofeedback, on the SCIO device. The professors involved with the postgraduate study, together with SCIO International Romania, are:

Dr. Aurel I. Bacean (Romania)

Dr. Igor Cetojevic (Cyprus)

Dr. Matthias Heiliger (Switzerland)

Stephanie Heiliger (Switzerland)

Dr. Codruta Bacean

Dr. Onut Bacean

This is the first international postgraduate study on biofeedback credited by the Romanian College of Physicians with CME (Continuous Medical Education) credits with an internationally recognized certificate.

This is a picture of Dr. Igor’s (Novak Djokovic’s doctor) Romanian license professional qualification a Neuro-Physiologist-Bio-Feedback-Bio-Resonance SCIO therapist.
DICHTIFICAT DE ABSOLVIRE

Igor Cetojevic

S-a absolvit cursurile postuniversitare de perfecționare cu durata de 16.06.2010 - 16.06.2011, în speciﬁcarea Neuroanatomie, neuroelectrofiziologie și biofeedback, cu apărarea: SCIO - placa numărată și biocoordonate.

Creat pentru acestui certificat i se acordă toate drepturile legale.

Rege
d

Secție Sf. Cătălin

A. M. 13.05.2011

Seminatul titularului

"The SCIO turned around my career"

Novak Djokovic

Neuro-Anatomy, Neuro-Electro-Physiologist in Biofeedback
SCIO Therapist with Bio-Resonance
Medical regulators do not decide what is or is not medicine. Medical Universities decide what is or is not medicine. When the University of Timisoara allowed medical text books to include our publications, which have been in peer reviewed journals for more than 5 years, this became undeniably a medical art. And beyond that, when the Government of Romania has issued a professional work qualification for this device, we have achieved complete compliance, validation and verification.
Click to read the Swedish Clinical Research

Click to see the German case study

Ovarian torsion of a pedicile, left, with bleeding on 18.04.2006 in the morning, same ultrasound BUT with no complaints at all after 12-hour SCIO treatment
The SCIO is 85% to 95% accurate

Abstract: We have used the SCIO device for 4 years now and have seen over 5000 patients. The device shows reaction profiles of the patient to vitamins, minerals, enzymes, homeopathics, and various other items. We have seen the accuracy of the reactions be uncanny accurate. This study will reflect the patient's profiles in our practice for over the last 3 years. As indicated in the title the device is very accurate.

Key Words:
CBR = Computerized Body Resonance scan = Scola
EPF = Electro-Physiological Reactivity
Input = the known diagnosis eg Shingles, Chickenpox etc. (undisputable inputs have been used)
Output = What the Scola / CBR scan finds.
Purpose of study: To check the accuracy of the Scola INPUT measurement.
The H1N1 measurement had a long, long, negatives. Using the older software. The newer one is basically 90% to 100% accurate.

Author:
Dr. Francois Theor

B. Med. Sc. (Medical Science) University of Pretoria, 1979
M.B.Ch.B. Bachelor in Medicine and Bachelor in Surgery Unv Preto's 1984

Research degrees:
Ph.D. (Biochemistry) 2010
D. Med. Sci. (Medical Science and Technology) 2010

What is Biofeedback Technology?

Biofeedback technology is an astonishing response tool that has changed the way we look at normal therapeutic processes. The QXCI/SCIO bio-feedback machine scans through your body completely, searching for viruses, fungi and other harms. A total of ninety two thousand areas are checked in a span of minutes. By means of quantum physics, the root of the QXCI/SCIO expertise is the diffusion of sixty five million minuscule electron-magnetic pointers into the body, several times for every second.

Legal England Research Publication + Presentation at the Royal Society of Medicine in London, England

The Butterfly Device

It has been said that something as small as the flutter of a butterfly’s wing can ultimately cause a typhoon halfway around the world.

Small effects can have large results and this device has made a large effect on Medicine

- Chaos Theory

The Royal Society of Medicine
Bioenergetische Diagnose und Therapie mit umweltmedizinischem Potenzial

Anwendungsmöglichkeiten des computergestützten Biorezonanssystems SCIO

Seit zwei Jahren macht auch in Deutschland ein bioenergetisches HighTech-System (SCIO) mehr und mehr von sich reden. Seine Attraktivität liegt in der Möglichkeit begründet, nicht nur die energetischen Störungen des Patienten zu diagnostizieren, sondern ihm gleichzeitig die meisten energetischen Therapien durchführen zu können. Dieser Beitrag befasst sich mit der Fragestellung, welche Optionen dieses System bietet, funktionale Störungen, die durch Substanzfehler der Umwelt hervorgerufen werden, zu analysieren und zu behandeln. Um zu einem besseren Verständnis zur nächsten in die Funktionsprinzipien dieses bioenergetischen Verfahrens eingeführt.

Abb. 1: Über das bioenergetische Diagnoseprogramm lassen sich auch gezielte umweltmedizinische Bündelungstherapien erstellen.

SCIO ist ein hochkomplexes bioenergetisches Verfahren, dass wie ein digital energetischer Muster eines Organismus erfasst und auf Grund einer tiefen registrierte Datenstandardisierte Therapiesätze auflegt, die dann auch direkt durch das System ausgeführt werden können. Es vereinigt in sich viele Prinzipien der energetischen Medizin, wie z. B. den Therapieplan eines individuellen Medizin- und Behandlungsprogramms.

Biophysikalische Steuerung des Organismus

Das System macht sich das Grundprinzip der Quantenphysik zunutze, nach dem zu jedem Molekül ein elektromagnetisches Feld gehört, also auch zu jeder Zelle, jedem Organ und zu jedem Elektron.

Gestörte Schwingungsmuster deuten auf ein anziehendes Krankheitspotenzial hin, das sich zu einem akuten oder auch chronischen Leiden auswachsen kann.

Mit dem System werden die energetischen Rest- und Wiederherstellungen behandelt, wobei die Frequenz von außen ausgewählt wird, um mit welcher Resonanzfrequenz des Organismus auf die Störung der energetischen Netzwerke abgestimmt wird, und welches halloenergetische Resonanzfeld ermittelt wird. Dieser Wirkungsmechanismus lässt sich mit einem Halbmond vergleichen, in dem ein hochsensibles elektronisches Ortagewerke zur Verfügung steht, mit dem es sich auf mehrere Kilometer Entfernung auslöschen kann, obwohl er nicht wahrnehmbar ist. Obwohl es sich hier um eine weitreichende Technologie handelt, ist die Wirkung auf den Organismus sehr gering. Intensiv in der Forschung ist der Effekt, dass Elektronen in der Umgebung des SCIO-Stroms aufgenommen werden und die energetische Struktur des Organismus verändert wird.
Click to see Spanish Research
Universidad Internacional de Cataluña

TRATAMIENTO CON EL SISTEMA DE BIOFEEDBACK-SCIO MEDIANTE BIORRESONANCIA EN PACIENTES DIAGNÓSTICADOS DE FIBROMIALGIA

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Universidad Internacional de Cataluña

IMUNE
International Medical University for Natural Education
Evidence Based Natural Energetic Medicine Education

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Cranial electrotherapy stimulation (CES) is a well-documented neuroelectrical modality that has been proven effective in some good studies of fibromyalgia (FM) patients. CES is no panacea but, for some FM patients, the modality can be valuable. This article discusses aspects of both CES and FM and how they relate to the individual with the condition. FM frequently has many comorbidities such as anxiety, depression, insomnia and a great variety of different rheumatologic and neurological symptoms that often resemble multiple sclerosis, dysautonomia, chronic fatigue syndrome and others. However, despite long-standing criteria from the American College of Rheumatology for FM, some physicians believe there is probably no single homogeneous condition that can be labeled as FM. Whether it is a disease, a syndrome or something else, sufferers feel like they are living one disaster after another. Active self-involvement in care usually enhances the therapeutic results of various treatments and also improves the patient's sense of being in control of the condition. D-ribose supplementation may prove to significantly enhance energy, sleep, mental clarity, pain control and well-being in FM patients. A form of evoked potential biofeedback, the EFx, is a powerful stress reduction technique which assesses the chief stressors and risk factors for illness that can impede the FM patient's built-in healing abilities. Future healthcare will likely expand the diagnostic criteria of FM and/or illuminate a group of related conditions and the ways in which the condition relate to each other. Future medicine for FM and related conditions may increasingly involve multimodality treatment that features CES as one significant part of the therapeutic regimen. Future medicine may also include CES as an invaluable, cost-effective add-on to many facets of clinical pharmacology and medical therapeutics.

Cranial electrotherapy stimulation (CES) with Alpha-Stim® is a well-documented neuroelectrical modality that has been proven effective in some good studies of fibromyalgia (FM) patients (REFERENCES 1 & 2). This article discusses aspects of both CES and FM and how they relate to the individual with the condition. CES is an FDA- and EU-recognized generic category for medical devices using microcurrent levels of electrical stimulation applied across the forehead via transcutaneous electrodes for the treatment of anxiety, insomnia and depression. Microcurrent (<1000 μA) stimulation usually means 1 mA or less, whereas transcutaneous electrical nerve stimulation (TENS) involves higher currents in the 60-100 mA range and with very different waveforms. CES treatment of anxiety and depression began in the USA in the early 1960s, and is still being prescribed routinely by several hundreds of physicians today, but has yet to achieve ubiquitous acceptance in medical practice. That is possibly because sufficient information has not been made available to practitioners regarding the safety and efficacy of CES as a treatment for the approved indications of anxiety, insomnia and depression. Using an electromedical device requires much of an additional learning curve for both practitioners and patients who are accustomed to the pharmacological model of intervention. Ingesting a capsule or a tablet does not always require the attention to detail that correct application of ear clip electrodes, for example, demands. We have been conditioned
Click to read Canadian Research on TVEP

Trivector Analysis of Bioterrain

ORP Map

ORP SCALE
Oxidized Ozone Rich +1300 mv
H2O=0mv
Hydrogen Rich – 600 mv Reduced

Y= VOLTS
H= Hydride Electron Donors
Z= Minerals

OCEAN

X=AMPS H+ Protons

22 Atm H2
0

Sacred Birthing

What is Sacred Birthing

Sacred Birthing is truly for one purpose: to birth a baby in a way that can best preserve the essence of divinity that accompanies each new being.

Extensive research of the EPFX-QXCI was done to show it effective in helping the Natural Birth
A Study to Detect the Efficacy of Micro-Current Electrical Therapy on Decubitus Wound

M.O. Ullah

This study tried to point out the effectiveness of Micro-current electrical therapy on decubitus wound of patients in different hospitals. The analysis shows that Micro-current Electrical Therapy (MET) has significant effect on healing the wound. The analysis also show that wound of female patients are healing significantly better than that of male patients and age is significantly influence for slightly decreasing the healing of wound.

Keywords: Decubitus wound, Micro-current Electrical Therapy (MET), multiple regression.
Dr. Hilf Klara’s research

In 2012 Dr. Klara Hilf has made an application to the Hungarian Ethic Committee for a study based on Albert Szent Gyorgy’s work. She has received one request from the EEKH for additional information, and after submission, after the passing of the 60 days, the study has been started.

On the 31st of January 2013, Andreea Taflan and Edit Barota have attended a course organized by SAASCO Kft. on Medical Device technical File. As a part of the presentation, was a section on Clinical Evaluations and Clinical Investigations with Fodor Eszter from Pharma-Hungary. Mrs. Fodor has stated that after the passing of 60 days without a response from the Ethics Committee, the study is considered approved and can be initiated.

Thus Dr. Hilf Klara has received approval from the Hungarian Ethics Committee to conduct research when they have not refused her after the passing of 60 days from her submission. In addition, the Ethics Committee of the Psychology Faculty of the University of Bucuresti has further approved Dr. Hilf Klara’s work and research.

In her office, with researcher’s help, Dr. Hilf Klara has duplicated the study conducted in 1973 initially by Prof. Desire’ Dubounet. This has brought us back to the start. After 40 years of research, validation, verification, we have proven that this device is safe and effective to the indications for use.

**Quantum Entwinement as a Principal of Human Communication 2014**

**Abstract:**

A research study first done in 1973 is being redone forty years later in 2013-2014. Volunteer subject teams were chose for their intimacy relationships. Mothers and child, marital partners, and close friends each in a pair are chosen. One member is isolated in a dark room with a stroboscope in front of their face. The other is hooked to an EEG device (the Eductor). The Eductor measures their Brain wave, heart electrical pattern, skin resistance and the VARHOPE of their body electric. At random intervals the strobe light in the room with the remote subject will flash for 30 sec. this will make a subtle shock to the system of the person and induce an ocular evoked potential. In the original experiment in 1973 the system shock to the one member of the pair provoked a similar evoked potential to the other.

In our 2013-2014 study we had 33 teams. In twenty-eight of the subjects there was an evoked potential at a distance detected in the VARHOPE of the receiver on the first stimulation. On the second strobe stimulation there was a reduced but measurable evoked potential. Verbal mind guesses were inaccurate but a bioelectric response was demonstrated.

This can be explained thru a type on quantum entwinement/entanglement principle of the body electric. But since this process is beneath the reticular activating system and thus not connected to the word area. Over emphasis of the word area in science has prevented us from truly accepting the ability of telepathy. Over 20 different research scientists have validated this incredible result in independent medical supervised studies.

There is a distinct dichotomy of the brain in a word area (left hemispheric logic) versus a Intuitive body electric a global Gestalt systemically wired Holistic nonverbal mind. The word area’s logic has dominated over the last few centuries. New developments in Insight Creativity have shown it occurs elsewhere in a nonverbal mind.

As we can see, there are basic barriers to acceptance of ESP. most of these barriers are intellectual and come from the verbal mind. First is a lack of a logical plausible explanation for the word area of the
minds of limited thinking scientists to comfort. The basic idea of Quantum Entwinement/Entanglement has now offered us such a plausible explanation. But the word area of limited verbal minded scientist struggle with this. Einstein struggled with this and called it “Spooky Action at a Distance”. And indeed most all things beneath verbal minded analysis are spooky indeed. But most of our societies and their inhabitants have held fast to their beliefs in spooky action at a distance. Who does not have a story of a thought or communication from someone at a distance? More than 75% of the people believe in ESP communication. Studies have shown its effect but not its reliability or at least its verbal reliability. We did this study in 1973 at Youngstown State University, and hence printed it in the graduate department publication. Forty years of further research into the components in the face of incredible resistance has led to a complete analysis of the concerning factors. Now with the advent of a new technology we have repeated the experiment. And now that modern science of Quantum Electro Dynamics has caught up with us a scientific explanation is apparent. And the belief of the masses has been verified. In fact with quantum entwinement/entanglement the ability of close people to communicate nonverbally at a distance is expected.

**Strobe Stimulation:**
One member of each subject team is isolated in a dark room with a stroboscope in front of their face. The strobe stimulation room is placed in a different building, to remove the electrical interference effect. At random intervals over an hour the strobe light in the room with the remote subject will flash for 30 sec. Hypothetically this will make a subtle shock to the system of the person and induce an ocular evoked potential. Verbal guesses had no results.

The other member of the study team is hooked to an EEG device (the Eductor). The Eductor measures their Brain wave, heart electrical pattern, skin resistance and the VARHOPE of their body electric. In the original experiment in 1973 the system shock to the one member of the pair provoked a similar evoked potential to the other.

**Result processes:** measures of attention were the moments of stimulus beginning and discontinuation, 1-2 seconds, in the measured subjects’ EEGs. An affirmative relationship was hypothesized to appear between the collective modifications of the stimulated subjects’ EEGs versus a strobe stimulated subjects. Control data using the same equipment and test conditions, but normal subjects tested, was collected to see if there was equipment and systematic artifacts.

**Results:** The placebo test resulted in a correlation of \( r = 0.05, p = 0.61 \); the experimental test resulted in \( r = 0.25, p = 0.0005 \).

Twenty eight (28) of the 33 pairs of participants showed independently significant correlations. Five (5) of the 33 pairs of participants showed no significant reactions.

Inspection of the stimulated subjects’ event-related evoked potentials showed that the stronger their responses in the theta band.

The analytical procedure was as followed:

1. Determine for each S period \( j \) the maximum value from the onset or offset of each stimulus up to one second post partner stimulus; call these maximum values \( \text{max} \).

2. Identify those S period where \( \text{max} \) values were larger than a maximum threshold value selected to identify at least 50 such periods; call this subset of periods \( \{ \text{max} \} \).

3. Find the peak value of the R ensemble variance array \( v \), that is, variance across all Rs, from each stimulus up to 1 second afterwards; call the time where this peak occurred \( p \) and the associated variance value \( v_p \).
4. Determine the R ensemble variance array $v$ for the subset of $\{\text{max}\}$ periods identified.

Brainwave frequencies drifting towards the theta band were found. And a slight change in volt potential max reading was displayed.

**Conclusion:** Under certain conditions, the EEG of a sensorial isolated human subject can influence event-related evoked potentials of their meaningful other at a distant. Showing a quantum entwinement of people at a distance.

**Discussion:**
In this study we reproduced the 1973 results showing that there was indeed spooky action at a distance between intimate pairs. Things once so joined maintain a level of communication regardless of distance.

**Eductor research**

The Eductor device is substantially equivalent to the SCIO, EPFX, QXCI devices. The Eductor device is today’s technology, today’s design, but operating within the same specs, only with a higher range of efficacy. We now present research done on the Eductor, echoing research done on the SCIO, EPFX and QXCI to show completely that our device is valid, verified and compliant with all the regulatory requirements and standards.
GSRtDCs Biofeedback Stimulation Increases Math, Insight and Language Memory Eductor 2014

**Supervising Researchers:** Dr Klara Hilf, Dr Danish
**Biofeedback Research Technician:** Neményi Rita

Permission of the Hungarian Ethics Committee and the Ethic Committee of the University of Bucharest Faculty of Psychology

**Institution:** International Medical University
**Sponsor:** QX World

**Dates:** September 2014  **Place:** Budapest, Hungary

**Abstract:**

93 subjects male and female were measured for basic Math skills, Insight and Language Memory.

Three GSR Cybernetic systems were compared to a placebo group. The Indigo, SCIO, Eductor 2014 with single signal generator and double signal generator setting were compared to placebo control testing. Cybernetic autofocusing of micro-current stimulation and biofeedback correction is used to maximize the effect.

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Two numbers added together make _A_ and Multiplied by each other make _B_

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<tr>
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<td>2-2</td>
</tr>
<tr>
<td>4</td>
<td>3</td>
<td>3-1</td>
</tr>
</tbody>
</table>

Start control Pre Test

| 7  | 12 | 3-3   |
| 8  | 12 | 2-6   |
| 12 | 36 | 6-6   |
| 16 | 48 | 4-12  |
Start stimulation tell them to relax with eyes closed wait one minute while getting one channel of CES

| 9  | 20 | 4-5 |
| 11 | 30 | 6-5 |
| 10 | 21 | 3-7 |
| 18 | 81 | 9-9 |
| 12 | 35 | 5-7 |
| 11 | 10 | 10-1 |
| 22 | 121 | 11-11 |
| 9 | 18 | 3-6 |

Next we tell them to relax with eyes closed wait one minute while getting two channels of CES

| 7  | 10 | 2-5 |
| 20 | 99 | 11-9 |
| 10 | 16 | 2-8 |
| 13 | 42 | 6-7 |
| 12 | 27 | 3-9 |
| 8  | 15 | 3-5 |
| 10 | 25 | 5-5 |
| 14 | 45 | 9-5 |

**Part two word memory retention**

| one | ichi |
| two | ni |
| three | san |
| four | yon |
| five | go |
| six | roku |
| seven | nana |
| eight | hachi |
| nine | kyuuu |
| ten | juu |

どうして？ (doushito?) = Why?
なに？ (nani) = What?
時間 (jikan) = Time
だれ (dare) = Who.
いつ (itsu) = When.
人 (hito) = Person.
どこ (doko) = Where.
日本 (nihon) = Japan.
Results of the math studies:

**Pre - Test**

- Man
- Woman
- Boy
- Shemale
- Woman (Born with…)
- Man (With Alzheimer)
- Man (Slower brain)
- Transexual
- Man
- Woman
- Man
- Woman
- Man
- Woman
- Woman
- Woman
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- Woman
### Results of the "Words"

<table>
<thead>
<tr>
<th>Age</th>
<th>Gender</th>
<th>Condition</th>
<th>Pre-Test</th>
<th>With Machine</th>
</tr>
</thead>
<tbody>
<tr>
<td>24</td>
<td>Woman</td>
<td>40 - Woman (Doctors...)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>46</td>
<td>Woman</td>
<td>49 - Man</td>
<td></td>
<td></td>
</tr>
<tr>
<td>21</td>
<td>Shemale</td>
<td>25 - Woman (Born with brain...)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>29</td>
<td>Woman</td>
<td>31 - Man (With Alzheimer)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>34</td>
<td>Woman</td>
<td>34 - Man</td>
<td></td>
<td></td>
</tr>
<tr>
<td>28</td>
<td>Woman</td>
<td>40 - Shemale</td>
<td></td>
<td></td>
</tr>
<tr>
<td>25</td>
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<td></td>
<td></td>
</tr>
<tr>
<td>17</td>
<td>Woman</td>
<td>21 - Woman (Slower brain...)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>50</td>
<td>Man (With Alzheimer)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>148</td>
<td>Woman</td>
<td>40 - Woman (Doctors...)</td>
<td></td>
<td></td>
</tr>
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</table>
Much better result with machine:
Better result:
Same result:
Worse result:
Pre - Test
With Machine
Second Wave

Math Test Results After Second Wave
Word Test Results After Second Wave

- Better result: 55%
- Same result: 38%
- Worse result: 6%
- Much better result with machine: 6%
In the test there were no reported significant risks. Two small headaches were reported on treatment that passed after several minutes. Of the 93 test subjects 90% had improvement in the word memory performance and the same in mathematical performance. The comparison to our placebo control group shows the effect of stimulation of the mental cognition similar effect to recently quoted research in the literature. Our hypothesis has been confirmed in this research. This has resulted in a p-value of 0.038 proving significance in our study. The GSRtDCs is safe + effective.
Discussion:

There were no reported risks during the study. The study showed clearly that the CES can stimulate math ability and memory retention. The history of micro-current CES positive effects on learning dates back decades. There have been no safety issues in the literature. There has been subtle but positive effects demonstrated on thousands of research documentation. This research shows the extra boost of positive effects of the second wave form generator only in the Eductor.
References:

1. ^ab 21CFR882.5800, Part 882 ("Neurological Devices")

2. ^ab Smith RB, Cranial Electrotherapy Stimulation: Its First Fifty Years


5. ^a Stillings D. A Survey Of The History Of Electrical Stimulation For Pain To 1900 Med.Instrum 9: 255-259 1975


13. ^a doi:10.1300/J184v09n02_02


15. ^a DOI: 10.1007/s11940-008-0040-y


19. ^a doi:10.1300/J184v09n02_02


30. "FDA medical device classifications


GSRtDCs Biofeedback Stimulation Increases Math, Insight and Language Memory Eductor 2014

Supervising Researchers: Dr Klara Hilf, Dr Danish

Therapist: Andrea Fantan

Permission of the Ethic Committee of the Univ of Bucharest Faculty of Psych Institution: International Medical University

Sponsor: QX World

Dates: October 2014 Place: Saut Marie Romania

Abstract:
75 subjects male and female ages 17 to 61 and 15 extra subjects were measured for basic Math skills, Insight and Language Memory. They were asked to report any changes in focus and confidence after the therapy.
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Then the same researcher asked the questions to the subjects. The subjects were read an example, then asked to solve with no stimulation, then with a single generator and then with two signal generators.

Pre Questions:

Do you like Mathematics???

Do you have confidence while doing Math???

Can you Focus while doing Math???

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</tr>
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Examples - give answers

Start control Pre Test - Now do not give answers

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</tr>
<tr>
<td>15</td>
<td>56</td>
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Start stimulation tell them to relax with eyes closed wait one minute while getting one channel of CES

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Post Questions after single wave generator:
Do you now have more confidence while doing Math???
Can you now Focus better while doing Math???
Does your ability to think seem clearer????
Anything else you feel.

Next we tell them to relax with eyes closed wait one minute while getting two channels of CES

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Post Questions after single wave generator:
Do you now have any more confidence while doing Math???
Can you now Focus any better while doing Math???
Does your ability to think seem any clearer????
Anything else you feel.

Results:
In the placebo group 4 of the 15 increased performance which was 27%. 5 out of the 15 reported noticeable increases in focus and confidence.

In the Eductor treatment group first wave form generator there was a noticeable increase in performance in %, confidence %, and focus %.

In the Eductor treatment group 2nd wave form generator there was a noticeable increase in performance in %, confidence %, and focus %.

<table>
<thead>
<tr>
<th></th>
<th>Performance</th>
<th>Confidence</th>
<th>Focus</th>
<th>all as increase over baseline</th>
</tr>
</thead>
<tbody>
<tr>
<td>Placebo</td>
<td>27%</td>
<td>33%</td>
<td>27%</td>
<td></td>
</tr>
<tr>
<td>1st WFG</td>
<td>67%</td>
<td>65%</td>
<td>64%</td>
<td></td>
</tr>
<tr>
<td>2nd WFG</td>
<td>76%</td>
<td>77%</td>
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This shows a dramatic increase in performance, confidence and focus over placebo control.

The GSRtDCs part of the program works to stimulate the brain for math performance and confidence as well.
GSRtDCs Biofeedback Stimulation Increases Math, Insight, Confidence, Focus and Language Memory in Students - Eductor 2015

**Supervising Researchers and Medical Review:** Dr Klara Hilf, Dr. Marco, Antonio, Rodriguez Infante, Dr. Hobian Veronica and Dr. Maria Baicu

**Therapist:** Andreea Fanatan, IMUNE Qualified GSRtDCs Research Technician

**Permission of the Ethic Committee of the National Institute of Recovery Physical Medicine and Balneo-Climatology, Bucharest, Romania**

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The technology has used a single wave form generator for CES since first registered with the US FDA in 1989. After over 35,000 such devices with not one reported significant risk, safety is obvious. Hundreds of studies have shown this technology to be effective, and now a second wave form generator will be tested.
Abstract:

22 subjects male and female ages 17 to 61 were asked to compete in 15 different chess games with another subject. Each player was hooked to a harness, one harness was active to the Eductor, and the other was a placebo control. No one knew which was which, so we have a perfect double blind study. They were asked to report any changes in focus and confidence after the therapy.
6 of the subjects were asked to play in 2 games and 2 subjects played in 3 games. In our games the player receiving the Eductor stimulus won each time. Even after losing the first time the player receiving the stimulus in the second game won when they got the stimulus. 2 subjects played 3 games. They lost when they got the placebo treatment and won when they got the treatment. The stimulation group reported increasing insight, play expertise, board vision, ability to see moves ahead and chess skills.

The Eductor 2015 with single signal generator and double signal generator setting were compared to placebo control testing. Cybernetic autofocusing of micro-current stimulation and biofeedback correction is used to maximize the effect.

There was a measurable chess performance increase in the treatment group. We asked the subjects to report focus, perception, creativity and confidence after the treatment. There was a dramatic 80% increase in confidence and focus.

Many new studies have shown the safety and efficacy of GSR trans-cranial stimulation inducing improved performance in mental acuity. These devices showed superior effect largely due to the autofocused cybernetic loop technology first developed in the 1980’s by IMUNE and first clinically proven in 2002 and proven again in several studies over the last two decades.

The technology has used a single wave form generator for CES since first registered with the US FDA in 1989. After over 35,000 such devices with not one reported significant risk, safety is obvious. Hundreds of studies have shown this technology to be effective, and now a second wave form generator will be tested.

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**GSRtDCs Biofeedback Cortical Excitation Stimulation Increases Insight in Students – with Eductor 2015**

**Supervising Researchers and Medical Review Staff:** Dr Klara Hilf, Dr. Marco, Antonio, Rodriguez Infante, Dr. Hobian Veronica and Dr. Maria Baicu

**Therapist:** Rita Nemenyi, IMUNE Qualified GSRtDCs Research Technician

**Permission of the Ethic Committee of the National Institute of Recovery Physical Medicine and Balneo-Climatology, Bucharest, Romania**

**Institution:** International Medical University

**Sponsor:** Biofeedback Srl

**Dates:** Feb, 27, 2015 **Place:** Budapest, Hungary
Abstract:

28 subjects, male and female ages 16 to 63, were asked to use Insight to solve a mental puzzle. They were also asked to report any changes in focus, perception, creativity and confidence after the therapy.

The subjects were asked to solve the nine dot problem to cover each of the nine points with four connected lines. The points of the problem are shown here on the left and the solution on the right. To solve this problem one must develop the insight to go outside the box of the lines.

The 28 subjects were shown the problem and give 5 minutes to solve it. 3 solved it with no stimulation and they were thus removed from the study. 25 could not and were thus entered into the study. The subjects were then given 5 minutes of single channel and the 10 minutes of the 2nd wave form generator (WFG) making a total of 15 minutes of stimulation with the Eductor. 5 solved it.
in the first 5 min. and 13 did it with the 2nd WFG. 18 total of the subjects could solve the puzzle in the
15 minutes all subjects were asked to rate their focus, perception, creativity and confidence.

The Eductor 2015 with single signal generator and double signal generator setting were compared. The lack of signal stimulation at the start of the test was used as a control. Cybernetic autofocusing of micro-current stimulation and biofeedback correction is used to maximize the insight effect.

There was a measurable performance increase in the treatment group. There was a dramatic 77% increase in confidence and focus. Confidence and focus is key for children in school.

We analyzed speed, accuracy and stress during insight problem solving. Once a base-line was established, the trans-cranial GSR Biofeedback cybernetic operation was turned on. After stimulation there was a significant noticeable increase in accuracy and speed of the insight problem solving. The second wave form generator performed better in the test.

Many new studies have shown the safety and efficacy of GSR trans-cranial stimulation inducing improved performance in mental acuity. These devices showed superior effect largely due to the autofocused cybernetic loop technology first developed in the 1980’s and first clinically proven in 2002 and proven again in several studies over the last two decades.

The technology has used a single wave form generator for CES since first registered with the US FDA in 1989. After over 35,000 such devices with not one reported significant risk, safety is obvious. Hundreds of studies have shown this technology to be effective, and now a second wave form generator will be tested.

GSRtDCs Biofeedback Stimulation Increases Math, Insight and Language Memory Eductor 2015

Supervising Researchers and Medical Review: Dr Klara Hilf, Dr. Marco, Antonio, Rodriguez Infante and Dr. Hobian Veronica
GSRtDCs Biofeedback Research Technician: Neményi Rita, IMUNE Qualified
Permission of the Hungarian Ethics Committee and the Ethic Committee of the University of Bucharest Faculty of Psychology
Institution: International Medical University
Sponsor: QX World

Dates: January 2015  Place: Budapest, Hungary

Abstract:

96 subjects male and female were measured for basic Math skills, Insight and Language Memory.

Three GSR Cybernetic systems were compared to a placebo group. The Indigo, SCIO, Eductor 2014 with single signal generator and double signal generator setting were compared to placebo control testing. Cybernetic autofocusing of micro-current stimulation and biofeedback correction is used to maximize the effect.
We analyzed speed, accuracy and stress during math problem solving and learning new words in a new language. Once a base-line was established, the trans-cranial GSR Biofeedback cybernetic operation was turned on. After stimulation there was a significant noticeable increase in accuracy and speed of the math and word skills. The second wave form generator performed better in the test.

Many new studies have shown the safety and efficacy of GSR trans-cranial stimulation inducing improved performance in mental acuity. These devices showed superior effect largely due to the autofocused cybernetic loop technology first developed in the 1980’s by Desire’ and first clinically proven in 2002. And proven again in several studies over the last two decades.

The technology has used a single wave form generator for CES since first registered with the US FDA in 1989. After over 35,000 such devices with not one reported significant risk, safety is obvious. Hundreds of studies have shown this technology to be effective, and now a second wave form generator will be tested.

Quantum Strength and Muscle Enhancement with the Eductor

Written by Professor Desire’ Dubounet

STUDY INFORMATION:
SUPERVISING RESEARCHERS + MEDICAL REVIEW STAFF:
Dr. Marco Antonio Rodriguez Infante, MD, Dr. Hobian Veronica, and Dr. Hilf Klara MD
Licensed Hungarian, Mexican and Romanian Medical Doctors
Therapists: Rob Wright, Rita Nemenyi
Permission of the Hungarian Ethics Committee, and the Ethic Committee of the National Institute of Recovery Physical Medicine and Balneo-Climatology, Bucharest, Romania
Institutional Monitor: International Medical University
Sponsor: Biofeedback Srl
DATES: March 2015

Abstract:

Nobel Laureate Hungarian born Albert Szent-Györgyi was one of the first researchers to look into the Quantum Electro Physiological nature of Biology. Albert Szent-Györgyi’s 1957 book on ‘Bio-Energetics” was one of the first to look into the electrical nature, and quantum nature of the electro-chemical process of life. I worked with Albert Szent-Györgyi in America before his death. Albert was truly one of the great minds with vision in all of humanity. I have followed up on his work ever since being with him.
In a series of studies we wanted to test the hypothesis that a CE mark registered Bio-Energetic device could measure the body electric of patients and through energetic intervention we could affect the biology in a positive way. We have tested such things as Chess playing, Insight, Math skills, Memory, Blood, Telepathy, Strength, Stamina, eye-hand-foot coordination among others were tested and found positive.

In this study we have tested body builders before and after Bio-Energetic therapy to see effects on power, stamina, and blood hormone level. In a series of different test we found complete safety and a surprising efficacy from the Bio-Energetic therapy.

**GSRtDCs Biofeedback Cortical Excitation Stimulation Increases Insight in Students – with Eductor 2015**

Supervising Researchers and Medical Review Staff: Dr Klara Hilf, Dr. Marco, Antonio, Rodriguez Infante, Dr. Hobian Veronica and Dr. Maria Baicu

Therapist: Rita Nemenyi, IMUNE Qualified GSRtDCs Research Technician

Permission of the Ethic Committee of the National Institute of Recovery Physical Medicine and Balneo-Climatology, Bucharest, Romania

Institution: International Medical University

Sponsor: Biofeedback Srl

Dates: Feb, 27, 2015  Place: Budapest, Hungary
Abstract:

As we have shown everything is an energetic collection of fields that hold atoms in their places. These fields that make us up are reactive with the environment. We must decide what is appropriate to eat and what to avoid. This education starts at the earliest of ages. Most of our current biological electro detection of what is good or bad for us takes place in the nasal-pharynx between smell and taste. The shape receptors of the smell and taste buds are electronic voltammetric field detectors. They sense a proper voltammetric fields that says it is good for nutrition or what is bad. The taste receptors do not absorb or metabolize the nutrient they only test it for intake by measuring the shape of the fields with the shape receptors of the tongue. Voltammetry is the science of analysis of the electrical fields of a substance.

We have shown the patent for the process of the QQC voltammetric analyzer. This device has been designed to work like the human tongue and to recognize the voltammetric signatures of items. These signatures are maintained as a 22x22x22 matrix of 10,648 separate shape vectors that constitute one signature. Since these fields reflect shapes they have a 3 Dimensional component and are referred to as the trivector voltammetric signatures. These complex signatures can be amplified and inputted into the body as part of the Xroid process in the EPFX, SCIO, Indigo, or what is now known as the Eductor.

The Xroid analysis is where the SCIO device measures the reaction of the body to over 10,000 substances at the calibrated speed of the body reaction. During the calibration of the SCIO device to the subject the device will measure the voltammetric field of the patient and then send in the QQC. Then the voltammetric signature of what is generally known as the weakest reactive substance (distilled water) is sent in over 20 times and the highest known reactive substance combination is
sent in 4 times. The starting speed is 103rd of a second. If the subject does not react significantly to the reactive substance versus the non-reactive distilled water the speed is reset minus one to 102nd of a second. The speed drops in this increment till the subject has a significant reaction to the reactive substances and it is repeatable. This then gives us a measure of the speed the subject reacts to items. Research in the 1980s showed that patients on morphine reacted much slower to norm patients. Then a variety of reactive speeds was shown thus making a speed of reactivity calibration need for proper testing.

There are several factors that can interfere with the testing of reactivity. If we test and test an item over and over there is adaptation. An aberrant movement, electrical wave form, or a brain wave surge can affect a reading. So we have seen that the reading of reactivity to a single item is not as significant as we would like. Till we could put a subject into a Faraday cage and perfectly control mental aberrations it is not likely. But we have seen that if we measure family reactions we can get some good insight into the reactivity fields of a subject. Research has shown that these families that we use to develop risk profiles are worth medical attention. In this review over one hundred thousand subject studies have verified the TVEP reactive families and the risk profiles have resulted from this work.

In this study over one hundred subjects were measured for Xrroid analysis on either a normal setting or a placebo setting. This was done to validate the TVEP validity and show that on normal setting there would be much more replication of data.

Proper Ethics committee and IRB were used and informed consent from subjects. The study took place in Europe and in America. Subjects were asked to do several measures of their wellness and they were measured for their Xrroid reactivity profiles before and after the test. Repeated items were counted in placebo versus real testing.


**Complementary and alternative therapy use in adult survivors of childhood cancer: a report from the Childhood Cancer Survivor Study.**


University of Minnesota, Minneapolis, Minnesota, USA. mertens@epi.umn.edu

BACKGROUND: Little information is available on the use of complementary and alternative medicine (CAM) in long-term survivors of childhood and adolescent cancer. PROCEDURE: The Childhood Cancer Survivor Study (CCSS) is a resource evaluating the long-term effects of cancer and associated therapies in 5 years survivors of childhood and adolescent cancer diagnosed between 1970 and 1986 before the age of 21 years. A survey of CAM use during the previous year was distributed in 2000-2001 and completed by 9,984 survivors and 2,474 sibling controls. RESULTS: CAM use reporting was similar in cases (16.4%) and siblings (41.1%). Compared to female siblings, female survivors were more likely to use biofeedback (OR=3.8; 95% CI=1.6-9.4) and hypnosis guided imagery (OR=1.2; 95% CI=1.0-1.4). Male survivors were more likely than male siblings to use herbal remedies (OR=1.3; 95% CI=1.1-1.6). Factors associated with CAM use in survivors included elevated scores on the brief symptom inventory (BSI) (OR=1.6; 95% CI=1.3-1.9), prolonged pain (OR=1.5; 95% CI=1.1-1.8), and having seen a physician in the past 3 years (OR=1.6; 95% CI=1.4-1.8). Survivors reporting low alcohol intake and excellent or good general health reported lower levels of CAM use (OR=0.7; 95% CI=0.6-0.8 and OR=0.8; 95% CI=0.7-0.9, respectively). CONCLUSIONS: Survivors have a similar reported use of CAM compared to a sibling cohort. However, our data suggest that survivors turn to CAM for specific symptoms related to previous diagnosis and treatment. Future research is needed to determine whether CAM use reflects unmet healthcare in this population. (c) 2007 Wiley-Liss, Inc.

PMID: 1736533 [PubMed-indexed for MEDLINE]
TVEP Explanation on one page
1. We are all made of atoms with electrons on the outside. The electrons and the atoms never touch each other they repel. The atoms are held in place by energetic fields. We are 99.9999999% energetic fields.
2. Every molecules of a substance have a specific signature energetic field which can be measured with Voltammetry
3. Every living creature has a reactive field that is seeking nutrition and repelling toxins. The global field of the body maintains a changing field of Voltage, Amperage, Resistance, Hydration, Oxidation and Ph variables. This reactive field reacts to stimulation.
4. By measuring the Voltammetric field of a substance and amplifying the field we can send the field into the body and measure its reactance to the stimulation.
5. We use over the skin thus Transcutaneous electrodes. We send in a known Voltammetric
6. Research has shown that the TVEP reactions are interesting not diagnostic. The family trends of reactivity are of more interest but still not accurate enough to be diagnostic. Thus a disclaimer of these patterns needs to be displayed.
Eductor - Cybermagnetic™ Chair

Vibrational Treatments for Weight Loss

Supervising Researchers: Dr Klara Hilf and Dr. Maria Baicu

Therapist: Rita Nemenyi

Permission of the Ethic Committee of the National Institute of Recovery Physical Medicine and Balneo-Climatology, Bucharest, Romania

Institution: International Medical University

Sponsor: Biofeedback Srl

Dates: December, 7, 2015  Place: Budapest, Hungary

Abstract:

Much study has been done on the effects of vibrations on the membrane of adipose tissues. Weight loss has become a major industry with bad diet and less action lifestyle becoming more prevalent. With the Eductor generating an electro-magnetic oscillation pulse, the cybermagnetic chair technology producing a magnetic pulse vibration and the sonic vibrations have been combined in this pre-study to determine safety and efficacy of the treatment.

15 subjects with weight loss issues were measured for weight before and then after a 30min therapy. There was an unexpected immediate loss measured in 8 of the subjects. (Averaging at 0.1 kilogram.)

The therapy was deemed safe and further study and long term follow up is needed to know more about the effectiveness.
Eductor - Cybermagnetic™ Chair Vibrational Treatment of Alzheimer’s

Supervising Researchers: Dr Klara Hilf and Dr. Maria Baicu

Therapist: Rita Nemenyi

Permission of the Ethic Committee of the National Institute of Recovery Physical Medicine and Balneo-Climatology, Bucharest, Romania

Institution: International Medical University

Sponsor: Biofeedback Srl

Dates: September, 27, 2015   Place: Budapest, Hungary

Abstract:

12 subjects, 7 males and 5 females ages 40 to 78, were asked to use 7 different words in different sentences. This is a classic medical test for Alzheimer’s. After the use of the words in sentences the subjects were asked to then remember the words. 2 or less can mean the possibility of Alzheimer’s. 5 of the subjects demonstrated memory weakness of Alzheimer’s.

Then a memory test of number chains was given to each subject. And we found how many numbers from a chain they could remember. They were also asked to report any changes in focus, perception, creativity and confidence after the therapy.

The Eductor 2015 with double signal generator setting were used with the Cybermagnetic™ Chair. The lack of signal stimulation at the start of the test was used as a control. Cybernetic autofocusing of micro-current stimulation and biofeedback correction is used to maximize the insight effect.

There have been reports in the literature of a sonic pulse of 28khz to 30khz being able to dissolve Alzheimer’s plaque in rats. We proposed using a vibrational medicine similar pulse imbedded in an entertaining movie to see if we could get positive effects on Alzheimer’s patients.

Also here have been reports of memory improvement with blue light on the eyes. We used a blue light stimulating goggles on the subjects during the study as well.

There was a measurable performance increase in the treatment group. There was a dramatic 67% increase in confidence and focus. Confidence and focus is key for Alzheimer’s recovery.
Conclusion

After all this research conducted to the letter of the law, in the entire Europe (Spain, Germany, Romania, Hungary, Italy, France, England), Switzerland, United states of America, Mexico, China, Mozambique, South Africa, Canada, now we present to you complete evidence of validation, verification, safety and efficacy of our technology.

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- Environmental Medicine by Dr. Med. Rainer Mutschler and Dr. Andreas Kott, Umweltmedizin, 09/07


- A study to detect the efficacy of Micro-current Electrical Therapy on Decubitus Wound, M.O.Ullah, Med. Sci. 7(8) 1320-1324, 15 November 2007.

- SCIO technology - An inverted ovarian tumor on the left side, Stephanie Heliger and Matthias Heliger M.D.

- Sacred Birthing, Extensive Research of the EPFX-QXCI was done to show it effective in helping the Natural Birth, Kauai, Hawaii
EMOTIONS FROM EEG
towards real-time valence monitoring

Raw EEG data
F3/4 or F7/8

Frequency spectra:
alpha power
2 second windows

Real-time Valence
Ratio of alpha power
right/left

High ratio = positive emotions
Low ratio = negative emotions

http://www.downloads.imune.net/journals/2018%20Unconscious%20Human%20Emotion%20Displayed%20in%20Real%20Time/Unconscious%20Human%20Emotion%20Displayed%20in%20Real%20Time%20ijmshnem.pdf
Possible cures for:

- Anxiety
- Addiction
- Chronic pain
- Sleep disorders
- Attention Deficit Disorder
- Migraine
THE SPECTACULAR SCIENCE BEHIND SONOGENETICS: CONTROLLING THE BRAIN THROUGH SOUND

The Eductor is the Only
Cybernetic Feedback
System for Energetic
Medicine Therapy
By Measuring the
Body Electric then
Adjusting Therapy, We
Increase safety+Efficacy

Legal TUV Registration
of Claims to Address Stress
and Lifestyle, Reduce Pain,
Increase Performance,
Reduce Anxiety,
Heal Injuries,
Correct Ph,
Balance the Body
Electric

TREATS
MTENS, MCES,
GSR, TDCS, EWH

Computer Software
reads VARHOPE
(Volts, Amps, Resistance,
Hydration, Oxidation + Ph)
and Calculates the
best Stimulation to
send out in the next
signal then software
remeasures. Thus it
makes a cybernetic
auto-focusing loop

Volt Meter with Freq
Counter to Measure
VARHOPE Strength
EEG, ECG, EMG, GSR

Signal Generator to send Stimulation
for TENS, MCES,
EWH + Voltammetric
signatures
DECLARATION OF CONFORMITY

Maitreya Srl of Romania

We declare that the device: Cybernetic Biofield Adjuster

meets all essential requirements for:
Class 1 Safety and Efficacy Standards

- Protection of the health and the safety of the user:
  - EN 60950-1:2007

- Electromagnetic compatibility
  - ETSI EN 301 489-1 V1.6.1
  - ETSI EN 301 489-8 V1.2.1

- Effective use of radio frequency spectrum:
  - ETSI EN 301 502 V8.1.2
  - ETSI EN 300 609-4 V8.0.2

Maitreya Srl Quality Control Officer
Desire' Dubounet

Issued 2-2-2019
Here is a list of Eductor Medical Claims that have been OKed by the CE

Mark and other regulators

1. SOC- Stress, Behavior and Lifestyle Questionnaire and Risk Analysis
2. EEG- Dual channel for Bi-hemispheric comparison
3. GSR- Quadrant measures to guide Physiological Therapy
4. EMG- Quadrant measures to guide Physiological Therapy and Muscle Resonance
5. ECG- Heart Rate and Heart Pulse Amplitude Variability
6. TVEP (Transcutaneous Voltammetric Evoked Potential) or, EPR (Electro-Physiological Reactivity). This is the ability to measure a person's reactivity to the QQC Voltammetric electro signatures of isodes, alesodes, nosodes, and sarcodes
7. VARHOPE this measuring the basic body electrical factors of Voltage, Amperage, Resistance, Hydration, Oxidation and Ph of the body. all registered on the FDA510 k as well as you see from the second excerpt
8. VARHOPE correction, Thru trickle charge electro-stimulation we can fix the body electric factors, and enhance Sport performance
9. EWH this is Electro-Wound- Healing for stimulating repair of injured tissue
10. Electro-Acupuncture is OKed in our Clinical Evaluation
11. CES Cranial Electro-Stimulation is fully OKed for Migraine, Stress, Insomnia, Addiction, Anxiety, Depression
12. MTENS Micro-current transcutaneous-electro-nerval-stimulation for PAIN therapy
13. GSRtDCs Galvanic Skin Reactivity trans Direct Current Stimulation- for cybernetic loop auto focused therapy
14. GSRtDCs- Intellect Enhancement
15. GSRtDCs- Math Ability Enhancement
16. GSRtDCs- Insight Enhancement