# A new approach on stress-related depression & anxiety: Neuro-Psycho- Physical-Optimization with Radio Electric Asymmetric-Conveyer

Salvatore Rinaldi<sup>1,2,3</sup>, Vania Fontani<sup>1,3</sup>, Elena Moretti<sup>4</sup>, Barbara Rosettani<sup>1</sup>, Lucia Aravagli<sup>1,3</sup>, Giorgio Saragò<sup>1,3</sup> & Giulia Collodel<sup>4</sup>

<sup>1</sup>Rinaldi Fontani Institute, Florence, <sup>2</sup>University of Florence, Medical School of Occupational Medicine, Postural Optimization, <sup>3</sup>University of Florence, Master of Neuro Psycho Physical Optimization & <sup>4</sup>University of Siena, Department of Medical Sciences, Applied Biology Section, Italy

Received January 23, 2009

*Background & objectives*: Chronic social stress is an important factor responsible for the worsening of depressive disorders in humans. In this study we present the relational Neuro-Psycho-Physical Optimization (NPPO) with Radio Electric Asymmetric Conveyer (REAC-CRM) as the treatment to tackle the unconscious dysfunction adjustments carried out by the central nervous system as a response to environmental stresses.

*Methods*: Psychological stress was measured in a group of 888 patients using the Psychological Stress Measure (PSM) test, a self-administered questionnaire. Data were collected immediately before and after the 4-wk therapy cycle. The detection of anxiety and depression clusters by PSM test has been based on the Diagnostic and Statistical Manual of Mental Disorders (DSM-IV-TR, APA, 2000) criteria. Six hundred and eighty eight patients (212 males, 476 females, average PSM test total scores 107.9  $\pm$  23.13) were treated with REAC-CRM therapy; 200 (64 males, 136 females, average PSM test total scores 107.86  $\pm$  25.80) were treated with "placebo REAC-CRM therapy" and used as control.

*Results*: This study showed a significant reduction in scores measuring subjective perceptions of stress in the patients treated with a cycle of REAC-CRM therapy. At the end-point the number of patients reporting symptoms of stress-related anxiety and depression on the PSM test was significantly reduced (P<0.001); in the placebo group no significant difference was highlighted.

*Interpretation & conclusions*: NPPO therapy with a cycle of REAC-CRM was shown to reduce subjective perceptions of stress measured by the PSM test and in particular, symptoms of stress-related anxiety and depression.

Key words Allostatic load - anxiety - depression - REAC-CRM - stress

Chronic social stress is one of the most important factors responsible in the triggering and/or worsening of anxiety and depressive disorders in humans<sup>1-3</sup>. In recent years, the impact of social stress on the development of psychopathologies has been thoroughly investigated in

preclinical animal studies<sup>4,5</sup>. It has been hypothesized that life stress alters the dynamic regulation of the autonomic, neuroendocrine, and immune systems<sup>6,7</sup>.

The central nervous system (CNS) is constantly and unconsciously adapting to accommodate changes in the environment. Unfortunately this unconscious process hides from our perception the adaptations that are detrimental to our health and to our quality of life. Allostasis is the ability to maintain the dynamic stability of the physiological systems facing a constantly changing environment (environmental stress or allostatic load). The allostatic load<sup>8-10</sup> is constituted by the combined environmental pressures that, when they can no longer be managed through the best physiological response, determine the allostatic state that is the comprehensive result of an altered physiological response (adaptive dysfunction) to the allostatic load. The allostatic state is unable to guarantee the good management of the physiological systems and therefore the health status and the wellbeing of the individual. The therapy to optimize the response to the allostatic load and the allostatic state is now available with the Neuro-Psycho-Physical-Optimization (NPPO) with a medical apparatus named the Radio Electric Asymmetric Conveyer (REAC)<sup>11,12</sup>.

The REAC-CRM emits a radio frequency (RF) in the field of very low power microwaves. The REAC-CRM, due to the nature of its construction, can concentrate the signal emitted in the environment on specific points of the body of the patient to be treated. The interaction between the field of microwaves emitted by the instrument and the body of the patient is different to that produced by exogenous cerebral stimulation with transcranial magnetic stimulation<sup>13,14</sup> and with vagal nerve stimulation which was recently approved by the Food and Drug Administration for treatment-resistant major depression<sup>15,16</sup>. Indeed, REAC-CRM produces an autogenous signal which, when suitably transported and concentrated in specific points of the auricular pavilion, can produce biological responses. These biological responses have been the subject of repeated clinical observations over the years, always showing a general improvement in the physical-psychological characteristics of the subjects treated with NPPO<sup>17-21</sup>

In conditions of acuteness of stress, at a level of the CNS, the secretion of adrenaline and cortisol hormones stimulates memory and attention, to let the organism deal with the environmental stressors. In the case of chronic stress there is progressive damage to some structures of the nervous system and in particular there is an atrophy of the apical dendrite of the neurons of the hippocampus, of the amygdale and damages to the dendrite of the neurons of the prefrontal cortex. The most serious and frequently observed symptoms were those of the clusters for mood depression and anxiety. Anatomic pathological damage to the cerebral cortex has also been shown in mood disturbances with a reduction of the glial tissue of the prefrontal cortex<sup>22</sup>.

The purpose of the present study was to verify whether the use of REAC-CRM was effective in reducing subjective perceptions of stress measured by a validated questionnaire, the PSM test<sup>23-25</sup> that allows the precise classification of the subject studied in a stress well-being scale to accurately assess the effectiveness of the treatment. In particular the influence on the symptoms of anxiety and depression, according to the definition of mixed anxiety and depressive disorder given by Diagnostic and Statistical Manual of mental disorders (DSM-IV-TR)<sup>26</sup> and correlated to stress conditions, were considered.

### **Material & Methods**

*Patient selection*: Between January 1999 and December 2007, a total of 888 selected subjects, who attended at Rinaldi-Fontani Institute showing different types of stress-related symptoms (such as tension headaches, high blood pressure, migraine headaches, anxious tremors, colitis, irritable bowel syndrome, bruxism, neck and back pain, chronic pain syndrome, bronchial asthma, peptic ulcer disease, skin disorders, insomnia) gave their consent and were included in the study. No patients were taking psychotropic medication.

The subjects (randomly selected for each group) were divided into 2 groups, Group A patients were given active treatment and patients in Group B were given placebo treatment. Group A comprised 688 subjects (476 females, average age  $42.3 \pm 13.7$  yr, and 212 males, average age  $42.9 \pm 15.4$  yr) treated with a cycle of NPPO of "active" REAC-CRM, and Group B comprised 200 subjects (control group) (136 females, average age  $42.5 \pm 16$  yr, and 64 males, average age  $43.8 \pm 16.7$  yr), treated with a cycle of NPPO of "matter and for the second seco

The clinical trial reported here was registered in the Primary Register of the Australian New Zealand Clinical Trial Registry (ANZCTR) (No. ACTRN12607000429459) and reported in the International Clinical Trials Registry Platform Search Portal - WHO.

*Psychological test and psychiatric assessment:* The Psychological Stress Measure (PSM)<sup>23-25</sup> was specifically developed to detect the stress levels in nonclinical population. The PSM is usually a 49-item selfreport paper and pencil questionnaire but in this study we used an electronic version to collect and process the data, and analyze the results. Each item is based on clusters of stress conditions: loss of self-control, irritability, psychological sensations, confusion, anxiety, depression, physical pain, hyperactivity and acceleration. The patient was asked to answer questions about his/her psychological stress using a 4-point scale to describe the intensity of his/her psychological stress condition (very much=4, much=3, little=2, none=1). The final score is expressed in Total Points (TP, T=Z\*10+50) according to the summary of the results of each item, and also as a percentile. The Total Points report normative data in the tables in percentiles and in T Points (T=Z\*10+50). In our study, we used the Total Points. As the focu s of this study was on anxiety and depression, the scores obtained from items 6-13-15-29 of the PSM test were specifically used.

The PSM demonstrates sound psychometric properties of validity, reliability, and convergent/ divergent validity. Internal consistency of 0.97 was demonstrated, as well as test-retest reliabilities of 0.63 over a two-week interval, 0.67 over a one month interval, and 0.55 over a 6 month interval. Assessment data were collected at two points in time: immediately before (t0) and immediately after (t1) the therapy/ placebo cycle of 18 sessions, which took place within a period of about 4 wk. The PSM scores are correlated by a factor of 0.68 with behavioural anxiety, and by a factor of 0.69 with the state of anxiety (P < 0.05). On the basis of this close correlation, we analyzed in particular the cluster relating to anxiety and depression. Both group of patients were clinically evaluated at t0 and t1 by a psychiatrist.

The detection of anxiety and depression clusters by PSM test was based on the Diagnostic and Statistical Manual of Mental Disorders (DSM-IV-TR, APA, 2000) criteria for mixed anxiety and depressive disorder (entry F41.9)<sup>26</sup>.

Description of the Radio Electric Asymmetric Conveyer (REAC-CRM) and of Neurologicalpsycho-physical optimization: The REAC-CRM is an innovative medical device<sup>11,12</sup> aimed at promoting the Neuro-Psycho-Physical Optimization (wellbeing and a reduction in the adaptive dysfunctional modifications in the nervous system induced by stress). It is a new medical instrument that uses the effects produced by the interaction between the electromagnetic field of the human body (~30-300 GHz, of about 3 mW/m2)<sup>27</sup> and that produced by the instrument (2.4 or 5.8 or 10.5 GHz, measurable from the emitter about 0.1 mW/m2) which lasts approximately a few milliseconds. This emission is dispersed into the environment and it is then received by a probe (conveyor) placed on the body of the patient. During the receiving process this probe (conveyor) allows an interaction between the emitted electromagnetic field and that of the treated patient. The target of the process is to allow the CNS to selfperceive the dysfunctional adjustments occurred, without the conditioning of the dysfunctional cognitive filters that maintain the-neuro-psycho physical-pathological alterations (dysfunctional adjustments). This allows optimization of the allostatic responses, progressively reducing the various dysfunctions.

The instrument that we used is registered under the trademark of CRM (ASMED, Florence, Italy). The result is an activation of the central nervous system that can optimize neuropsycomotorial function and reduce the adaptive dysfunctional modification of the nervous system induced by stress. The NPPO auricular therapy protocol was used to manage and optimize these modifications<sup>17-21</sup>. The REAC-CRM probe was applied to seven specific points of the auricular pavilion, the same points that are also used in auricular therapy to treat neurovegetative symptoms and diseases (Shenmen, kidney, stomach, heart, occiput, ipotalamus, prefrontal cortex). Only 18 sessions of NPPO with the REAC-CRM active (group A) and inactive (group B) were administered to each patient on alternate days, during a 4 wk therapy cycle after the first PSM test. The aim of REAC-CRM therapy was to optimize the responses of the CNS and of the whole organism against unknown alterations due to stress from continuous interaction with the environment.

Each therapeutic session lasted for approximately three seconds. The protocol was painless, non invasive, did not require the collaboration of the patient and was completely without side effects.

After 18 sessions of NPPO with the active REAC-CRM (group A) and the inactive REAC-CRM (group B), the PSM test was repeated (t1).

Statistical analysis: Statistical analysis of Total points and Point T scored at t0 and t1 was performed using the Wilcoxon Signed Ranks test and P<0.05 was considered significant. The number of patients showing anxiety/ depression before and after therapy in both groups was compared using the McNemar test of Symmetry for statistical significance. The mean percentage change of Total points and T points before and after therapy between Groups A and B was calculated and compared by the Wilcoxon-Mann Whitney Rank Sum Test.

## Results

The mean of Total Points and T Points obtained by the PSM test before and after "active" and "placebo" REAC-CRM therapy in both groups of patients are shown in the Table.

In Group A, before the treatment 312 patients (45%) reported stress related symptoms of anxiety/ depression, whereas only 84 (12%) reported the persistence of these clinical targets after a cycle of "active" REAC-CRM therapy. In particular, the mean of Total Points and T Points decreased from 107.9 to 87.3 and from 57.1 to 48.2, respectively and was statistically significant (Wilcoxon Signed Rank Test P<0.001 for Total Points).

In Group B, before the treatment 76 patients (38%) reported stress related symptoms of anxiety/depression, and after the cycle of "placebo" REAC-CRM therapy 72 patients (36%) reported the presence of these clinical targets. The difference was not significant.

In Group A, the PSM scores (Total Points and T Points) after the cycle of REAC-CRM treatment were lower and less variable (Fig. 1). In contrast, the distribution of scores for the "placebo" group (Group B) was similar in both the first and second tests (Fig. 2).

Table.	Total	points a	and T p	points	obtain	ed ir	n group	A and ii	1 group	E
by the	PSM	test bef	fore (t0	) and	after (1	t1) tł	herapy/j	olacebo		

Patients	Total points PSM test	T points PSM test	Anxiety/ depression No. of patients (%)						
Group A (n=688) (t0)	107.9 ± 23.13	57.06 ± 9.70	312 (45)						
Group A (t1)	87.3 ± 16.21	$48.25\pm6.90$	84*(12)						
Group B (n=200) (t0)	$107.86 \pm 25.80$	57.05 ± 10.62	76 (38)						
Group B (t1)	$106.32 \pm 25.88$	56.09 ± 11.02	72 (36)						
Values are mean $\pm$ SD * <i>P</i> <0.05 compare to before therapy									



**Fig. 1.** Distribution of the PSM scores in treated patients (Group A,  $n^{\circ}$  688). Scores obtained in the first test were significantly different (*P*<0.05) to those of the second test performed after the therapy. The scores after therapy were lower.



Fig. 2. Distribution of the PSM scores in untreated patients (Group B,  $n^{\circ}200$ ). The distribution of scores was similar in both the first and second tests.

The McNemar test was used to analyze the difference in number of patients showing anxiety/ depression before and after therapy. Only 8 patients of Group B, who reported stress related symptoms of anxiety/depression before the cycle of "placebo" REAC-CRM therapy, did not show the same symptoms after the cycle of placebo. On the contrary, 240 patients belonging to Group A found relief from anxiety/depression as they did not report these symptoms after the cycle of REAC-CRM therapy (P<0.001).

The mean percentage change of Total Points and Points T between Group A and B was also analysed. The difference between the two groups was significant (P<0.001) for both Total points and Point T, confirming the positive effect of the REAC-CRM therapy on anxiety/depression.

### Discussion

The use of electricity and magnetic fields in biomedical studies, and particularly in the treatment of disturbances of the nervous system, is not a new idea<sup>28</sup>. The weak environmental emission of a radiofrequency field and its dispersion due to the interaction with the body was able to induce weak currents in the human body that activated the nervous system. Preliminary studies have suggested a reduction in general stress levels and especially in correlated stress disorders when REAC-CRM therapy is applied<sup>18-21</sup>.

This study was aimed at assessing the significant possible improvement of stress conditions in examined patients treated with REAC-CRM therapy, in particular on behavioural anxiety/depression. After a cycle of REAC-CRM therapy, a significant reduction in the scores measuring subjective perceptions of stress in the studied patients was observed. In particular, a significant reduction of symptoms was reported by the group showing signs of behavioural anxiety-depression. On the other hand, in the untreated control group, no significant difference was found. Also, the cluster measuring behavioural anxiety/depression remained unchanged.

Growing literature indicates that the symptoms of stress-related anxiety and depression are the results of allostatic processes on specific cerebral areas<sup>29</sup> and our data also showed that REAC-CRM was an effective instrument to optimize the responses of the CNS. Recently Collodel *et al* <sup>17</sup> suggested that NPPO therapy with REAC-CRM treatment may realize a general improvement in spermatogenetic condition, as demonstrated with sophisticated tools such as TEM and FISH, in males with idiopathic infertility.

The results of this study showed that NPPO therapy with REAC-CRM could reduce subjective perceptions of stress measured with the PSM psychometric test and in particular the symptoms of stress-related anxiety and depression. Hendriks *et al* <sup>30</sup> reported that cognitive behavioural therapy is efficacious for the treatment of late-life anxiety disorders.

This new therapy has the advantage of being painless, non-invasive and totally free of side-effects. Moreover, REAC-CRM therapy is not pharmacological and can represent an efficient support in many medical fields, as it does not interfere with the simultaneous use of other therapeutic approaches. This research highlights the efficacy of NPPO with REAC-CRM therapy on anxious-depressive scale only in non psychiatric subjects; these conditions represent the most comprehensive mental suffering in the general population.

The main limitation of this research was that the efficacy of NPPO with REAC-CRM therapy was demonstrated on subjects who although identified as psychometric anxious-depressive, did not need any clinical intervention. In current psychiatric terminology, the observed affective phenomenology is classified as "below threshold" (presence of certain symptoms of anxiety or depression, which are however insufficient to set up a "primary diagnostic entity") or "sub-clinical" (complete clinical picture, but without serious symptoms). Moreover, the patients who have responded positively to REAC-CRM therapy, showed clearly stress-related symptoms and therefore "reactive" psychic symptoms. They did not show any "endogenetic" features such as "melancholic" forms of depression, for which, the best choice in terms of speed of response and global therapeutic efficacy was a psychopharmacological treatment.

Further studies are needed to verify the stability over time when using more than one cycle, although it could be very difficult to obtain and mantain a selected group, especially after therapy.

In conclusion, our results showed that REAC-CRM therapy will help to speed up the physiological capability of recovery of the person, optimising the adaptive response to environmental stressors and contributing to the elimination of dysfunctional adaptive responses.

#### Acknowledgment

The authors thank Matteo Lotti Margotti, for helpful discussion in statistical analysis, Alessandro Castagna, for helpful discussion, and to Piero Mannu, psychiatrist, for the clinical evaluation of the patients and fruitful discussions.

#### References

- 1. St-Jean-Trudel E, Guay S, Marchand A. The relationship between social support, psychological stress and the risk of developing anxiety disorders in men and women: results of a national study. *Can J Public Health* 2009; *100* : 148-52.
- Fagring AJ, Kjellgren KI, Rosengren A, Lissner L, Manhem K, Welin C. Depression, anxiety, stress, social interaction and health-related quality of life in men and women with unexplained chest pain. *BMC Public Health* 2008; 8 : 165.
- Clow A, Hamer M. The iceberg of social disadvantage and chronic stress: Implications for public health. *Neurosci Biobehav Rev* 2010; Epub Mar 17.
- 4. Barsy B, Leveleki C, Zelena D, Haller J. The context specificity of anxiety responses induced by chronic psychosocial stress in

rats: a shift from anxiety to social phobia? *Stress* 2010; *13* : 230-7.

- Schmidt MV, Scharf SH, Liebl C, Harbich D, Mayer B, Holsboer F, *et al.* A novel chronic social stress paradigm in female mice. *Horm Behav* 2010; 57 : 415-20.
- 6. Lucini D, Norbiato G, Clerici M, Pagani M. Hemodynamic and autonomic adjustments to real life stress conditions in humans. *Hypertension* 2002; *39* : 184-8.
- Bellinger DL, Lubahn C, Lorton D. Maternal and early life stress effects on immune function: relevance to immunotoxicology. *J Immunotoxicol* 2008; 5: 419-44.
- 8. Kuchel GA. Frailty, allostatic load, and the future of predictive gerontology. *J Am Geriatr Soc* 2009; *57* : 1704-6.
- Juster RP, McEwen BS, Lupien SJ. Allostatic load biomarkers of chronic stress and impact on health and cognition. *Neurosci Biobehav Rev* 2009; Epub Oct 12.
- Dowd JB, Simanek AM, Aiello AE. Socio-economic status, cortisol and allostatic load: a review of the literature. *Int J Epidemiol* 2009; 38 : 1297-309.
- Rinaldi S, Fontani, V. Inventor Rinaldi S, Fontani V, assignee. Radioelectric Asymmetric Conveyer for therapeutic use patent EP1301241 (B1). 2000 October 11, 2006.
- Rinaldi S, Fontani V. Inventor Rinaldi S, Fontani V, assignee. Radioelectric Asymmetric Conveyer for therapeutic use. USA patent 7,333,859 2001.
- George MS, Lisanby SH, Avery D, McDonald WM, Durkalski V, Pavlicova M, *et al.* Daily left prefrontal transcranial magnetic stimulation therapy for major depressive disorder: a sham-controlled randomized trial. *Arch Gen Psychiatry* 2010; *67*: 507-16.
- Yip AG, Carpenter LL. Transcranial magnetic stimulation for medication-resistant depression. *J Clin Psychiatry* 2010; 71 : 502-3.
- 15. Henderson JM. Vagal nerve stimulation versus deep brain stimulation for treatment-resistant depression: show me the data. *Clin Neurosurg* 2007; *54* : 88-90.
- 16. Fitzgerald PB, Daskalakis ZJ. The use of repetitive transcranial magnetic stimulation and vagal nerve stimulation in the treatment of depression. *Curr Opin Psychiatry* 2008; *21* : 25-9.
- Collodel G, Moretti E, Fontani V, Rinaldi S, Aravagli L, Sarago G, *et al.* Effect of emotional stress on sperm quality. *Indian J Med Res* 2008; *128* : 254-61.
- Mannu P, Rinaldi S, Fontani V, Castagna A, Lotti Margotti M. Radio Electric Treatment vs. Es-Citalopram in the treatment of panic disorders associated with major depression: an open-

label, naturalistic study. *Acupunct Electrother Res* 2009; *34* : 135-49.

- Rinaldi S, Fontani V, Aravagli L, Lotti Margotti M. Psychological and symptomatic stress-related disorders with radio-electric treatment: psychometric evaluation. *Stress and Health* 2010 Epub Jan 20.
- 20. Castagna A, Rinaldi S, Fontani V, Aravagli L, Mannu P, Lotti Margotti M. Does osteoarthritis of The knee also have a psychogenic component? Psycho-emotional treatment with a radio-electric device vs. intra-articular injection of sodium hyaluronate: An open-label, naturalistic study. *Acupunct Electrother Res* 2010 (in press).
- Rinaldi S, Fontani V, Aravagli L, Mannu P. Psychometric evaluation of a radio electric auricular treatment for stress related disorders: a double-blinded, placebo-controlled controlled pilot study. *Health Qual Life Outcomes* 2010; 8 : 31.
- 22. Ongur D, Drevets WC, Price JL. Glial reduction in the subgenual prefrontal cortex in mood disorders. *Proc Natl Acad Sci USA* 1998; 95 : 13290-5.
- Trovato GM, Catalano D, Martines GF, Spadaro D, Di Corrado D, Crispi V, *et al.* Psychological stress measure in type 2 diabetes. *Eur Rev Med Pharmacol Sci* 2006; *10*: 69-74.
- 24. Lemyre LTR. Mesure du stress psychologique. Se sentir stressé-e. *Rev Canad Sci Comport* 1988; 20: 302-21.
- 25. APA. *Diagnostic and statistic manual of mental disorders*. Washington DC: American Psychiatric Press; 2000.
- Lemyre L, Tessier R. Measuring psychological stress. Concept, model, and measurement instrument in primary care research. *Can Fam Physician* 2003; 49 : 1159-60, 66-8.
- Valberg PA, van Deventer TE, Repacholi MH. Workgroup report: base stations and wireless networks-radiofrequency (RF) exposures and health consequences. *Environ Health Perspect* 2007; *115*: 416-24.
- Daban C, Martinez-Aran A, Cruz N, Vieta E. Safety and efficacy of vagus nerve stimulation in treatment-resistant depression. A systematic review. *J Affect Disord* 2008; *110*: 1-15.
- 29. Magarinos AM, McEwen BS. Stress-induced atrophy of apical dendrites of hippocampal CA3c neurons: comparison of stressors. *Neuroscience*1995; *69* : 83-8.
- Hendriks GJ, Oude Voshaar RC, Keijsers GP, Hoogduin CA, van Balkom AJ. Cognitive-behavioural therapy for late-life anxiety disorders: a systematic review and meta-analysis. *Acta Psychiatr Scand* 2008; *117*: 403-11.

Reprint requests: Dr Giulia Collodel, Department of Medical Sciences, Applied Biology Section, University of Siena Policlinico Le Scotte, Viale Bracci, 14, 53100 Siena, Italy e-mail: collodel@unisi.it

194