

**Title**

Psychobehavioral therapy for epilepsy

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**Abstract**

Growing evidence suggests a bidirectional interaction between epileptic seizures and psychological states, fuelling the interest in the development and application of psychobehavioral therapy for people with epilepsy (PWE). The objective of this article is to review the various psychobehavioral therapies in regard to their application, hypothesized mechanism and effectiveness. Most psychobehavioral therapy aims at improving psychological well-being and seizure control. Behavioral approaches, cognitive-behavioral therapy (CBT), and mind-body interventions are the most widely applied approaches for PWE. CBT, mind-body approaches and multi-model educative interventions have consistently demonstrated positive effects on enhancing well-being. Nevertheless, the effects on seizure control remain inconsistent partly attributable to small clinical trials and inadequate control groups. Assessor-blinded randomized controlled trials with sufficient power and carefully defined therapeutic components corresponding with objective and subjective outcome measures are recommended for future trial design.

**Keywords**

Epilepsy; stress; psychotherapy; mindfulness; mind-body; cognitive-behavioral therapy

## Questions

- 1 What psychobehavioral therapies are/may be useful in epilepsy?
- 2 What are the hypothesized mechanisms of action of psychobehavioral therapy for epilepsy?
- 3 Is psychobehavioral therapy effective in epilepsy?
- 4 What are the methodological challenges associated with the scientific study of psychobehavioral therapies and the proposed strategies for meeting those challenges?

### 1 Introduction

Accumulating evidence from animal models and clinical studies suggests a bidirectional relation between seizures and psychological states [1-3]. This understanding has been coupled by increasing research effort in incorporating psychobehavioral therapy in epilepsy care to improve psychological well-being and seizure control. This article aims to review the application and efficacy of psychobehavioral therapy for people with epilepsy (PWE), and to draw conclusions regarding the implications for future trial designs and clinical practice. The hypothesized mechanisms of these therapies are also discussed, which complements existing reviews [4, 5] For the purpose of this review, psychobehavioral interventions are defined as those based on theory of psychotherapy. Alternative remedies for epilepsy such as spirituality, energy healing, aromatherapy, although they appear to be psychological in nature, will not be considered.<sup>1</sup>

### 2 What psychobehavioral therapies are/may be useful in epilepsy?

Therapeutic elements in psychobehavioral therapies that are most extensively applied include behavioral approaches, cognitive behavioral therapy and mind-body approaches. The majority of therapies are based on a primary therapeutic approach with a correspondent conceptualization of the psychological mechanism that underlies seizure activity [6-8]. However, some interventions employed a composition of therapeutic elements from different approaches; which are delivered as multi-model interventions [9-11].

These therapies may be delivered through individual-based therapy and/or group-based sessions, although most prospective clinical trials employed a multiple-session group therapy design. The number of sessions varied from brief interventions consisting of 2 sessions to comprehensive therapy with up to more than 40 sessions [9, 12, 13]. Some therapies involved also caregivers of PWE[14, 15]. Most interventions include an evaluation of the individual's psychological stressors which could potentially increase seizure activity. An assessment of environmental and behavioral obstacles that prevent the individual from regulating lifestyle-related physiologic seizure risk factors, including drug noncompliance and sleep deprivation, are emphasized.

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<sup>1</sup> All literature published in English before August week 4, 2013 is included. Inclusion criteria regarding sample size and study design were not defined. Relevant studies are identified by search in MEDLINE (Ovid) and PsycINFO database with key words: epilepsy or seizure or convulsion, and one of the following key word: alternative therapy, aura interruption, behav therapy, biofeedback, breathing, conditioning, cognitive behav therapy, counseling, countermeasure, desensitization, education, mind-body, mindful, psychological, psychotherapy, relaxation, self control, self efficacy.

This section describes the application of the various techniques. Discussion on their hypothesized mechanisms and critical evaluations of their efficacy will follow.

## **2.1 Behavioral approaches**

The behavioral approach for epilepsy is based on one of the earliest paradigms of psychotherapy and remains an essential element in many protocols. It operates by applying countermeasures upon seizure triggers and early stages of seizure activity to abort or reduce the likelihood of developing a seizure from the outset. Techniques such as conditioning, aura interruption, systematic desensitization and EEG biofeedback are the major therapeutic components [16-18].

Aura interruption and systematic desensitization operate similarly based on the recognition of pre-ictal and early ictal phenomena, with specific emphasis on the correlating psychosocial and emotional contexts. Countermeasures to interrupt the behavioral chain are applied when these pre-ictal or early ictal phenomena arise. The most common techniques are relaxation, abdominal breathing, attention shifting and inward monologue [6, 19, 20].

Operant conditioning by pairing an epileptic seizure with punishment and a seizure-free period with reward was mainly found in the early literature[21]. This method is used in conjunction with EEG biofeedback later on. Individuals are trained to increase 11- to 15-Hz EEG activity localized to the sensorimotor cortex to resemble a state of mental relaxation [22-25]. EEG activities are explicitly displayed on a computer screen. Rewards (games, token) appear on the screen when the desirable EEG patterns occur, and disappear when the EEG patterns fall beyond the range.

## **2.2 Cognitive behavioral therapy (CBT)**

CBT emphasizes the examination of the relationship between thoughts and emotions in the context of a specific event. Individuals learn to identify maladaptive patterns of thought and to establish alternative improved habitual cognitive and behavioral response patterns in order to enhance subjective emotional well-being. It is applied on a wide range of psychiatric disorders in systematized approaches. Therapeutic elements for PWE include psycho-education on disease knowledge, mood regulation techniques, behavioral and cognitive countermeasures to auras, the resolution of cognitive dissonance that may result in intense emotional distress, stress management as well as elements that facilitate lifestyle modification.

Most CBT protocols emphasize the individuals' perception of health locus of control and aim at empowering individuals to establish subjective control over their seizures. Further, their role and participation in treatment are promoted with motivational techniques [10, 15, 26, 27]. Journaling as a tool to record contextual seizure characteristics, relaxation logs and self-affirmation statements are commonly used. Some include structured workbooks that guide and involve individuals through the exploration of their seizure experiences. Andrews and Reiter pioneered with the development of a systematic, semi-directive, step-by-step counseling technique that is based on the workbook "Taking Control of Your Epilepsy" as an on-going guideline for individual sessions. It comprises 12 sessions

and includes detailed behavioral and cognitive restructuring techniques, lifestyle recommendations, assertive communication skills and goal setting exercises[28]. Since an increased awareness of the capacity for taking control of seizure precipitants may lead to a tendency to react with self-blame and guilt in some individuals, alternative behavioral responses and cognitive reframing strategies are offered in case such feelings should arise.

Some interventions for PWE are specifically designed to treat or prevent psychiatric comorbidities and to decrease the build-up of seizure activities [7, 29-32]. For example, Goldstein and colleagues reported using a 12-session standardized CBT individually for adults PWE with psychiatric comorbidity[7]. Similarly, Macrodimitis et al. developed group-based CBT for PWE with comorbid depression and anxiety[31].

### **2.3 Mind–Body approaches**

Mindfulness and acceptance-based meditation have become one of the most common mind-body approaches used in PWE in recent years. Although meditation styles vary, all practices involve self-observation of mental and bodily activity, attention training and the cultivation of process-oriented awareness. The therapeutic components of mindfulness are to acquire attention control by focusing on internal processes (breath, bodily sensations, thoughts, and emotions) and external stimuli (sights, sounds, smells, and texture) at the present moment (“here-and-now”), with non-elaborative attitude and nonjudgmental acceptance. These techniques have been formalized into standardized psychotherapy such as Mindfulness Based Stress Reduction (MBSR), Mindfulness Based Cognitive Therapy (MBCT) and Acceptance and Commitment Therapy (ACT). MBCT and ACT have been applied among individuals with seizures. Various skills are involved; basic techniques include mindful breathing, mindful eating, mindful awareness on sensations, thought and emotion labeling [8, 33-36].

Generic stress reduction techniques, such as relaxation exercise and breathing, are widely employed in conjunction with other psychobehavioral approaches for epilepsy. Most breathing techniques utilize awareness of breathing rate, rhythm and volume to achieve a state of calmness and to minimize physiological reactions to stress [37-39]. Moreover, they are often used jointly with muscle relaxation exercise, such as Jacobson’s progressive muscular relaxation (PMR) and yoga exercises.

### **3 What are the hypothesized mechanisms of action of psychobehavioral therapy for epilepsy?**

The application of psychobehavioral therapy on epilepsy as a chronic neurological disease does not necessarily aim at resolving psychopathology as its role assumed in the traditional psychiatric setting. Instead, therapies aim at assisting individuals to increase their ability of coping with their disorder on a psychological level with possible underlying neurobiological alterations that might contribute to improved seizure control and psychological well-being.

Despite their diverse approaches, all psychobehavioral therapies for epilepsy operate with the

aim to increase the individual's ability to prevent the build-up of seizure activity. In general, therapies aim to reduce both non-specific seizure precipitants (e.g. sleep deprivation, stress, physiologic risks), and specific internal (e.g. emotional distress, cognitive distortion) and/or external (e.g. sensory and environmental) contextual seizure precipitants, as illustrated by Figure 1a-1c and Figure 2. Several hypothesized mechanisms of action are discussed below.

### **3.1 Stress hypothesis**

Stress is widely accepted as a risk factor of seizures and can exacerbate seizure manifestation in humans and animal models [1, 40-42]. Since stress may alter autonomic nervous system functioning by increasing sympathetic activity, stress reduction techniques are possibly effective by increasing the parasympathetic response. The restoration of balance within the autonomic nervous system can hence import beneficial effects for epilepsy patients by reducing the physiological correlates of psychological stress, thereby preventing the build-up of seizure activity. Besides, hyperventilation could be an acute stress reaction towards the emergence of a seizure and consequentially a trigger of a seizure in itself. Stress reduction breathing such as deep abdominal breathing might reverse this effect by increasing partial pressure of carbon dioxide (PCO<sub>2</sub>) and thus lowering seizure activity [43]. Evidence is found in approaches that utilize a stress reduction model e.g. relaxation training, mindfulness meditation, and yoga exercises. Most of them demonstrated a reduction of seizure occurrence [8, 33, 37, 44-47].

### **3.2 Psychiatric comorbidity hypothesis**

Multiple studies have revealed a high prevalence of psychiatric disorders in PWE. The prevalence is fairly consistent across studies suggesting around 1 in 4 PWE have a psychiatric comorbidity [48, 49]. The high prevalence of comorbidity is attributed to a variety of factors including reactive processes, iatrogenic causes, genetic predisposition and seizure related endogenous changes impacting neurochemical and neurophysiological processes.

Furthermore, the relation between epilepsy and psychiatric disorder has been demonstrated to be bidirectional, whereby the latter can be a risk factor of the former [2, 3, 50]. There is evidence that individuals who have a history of a formal psychiatric diagnosis have a higher risk of developing seizures at a later time [51]. Patients with a history of a psychiatric disorder were also less responsive to antiepileptic drug treatment and had worse seizure outcome following epilepsy surgery [52, 53]. It has been hypothesized that both disorders share common pathogenic and psychogenic mechanisms that, in turn, explain their high comorbidity [54-56].

According to this hypothesis, treating psychiatric disorders and psychological symptoms in PWE has the potential to improve not only psychological well-being but also seizure manifestation. This has become the rationale that underpins several CBT protocols that aim at resolving psychological complications and improving well-being.

### **3.3 Self-efficacy hypothesis**

Health locus of control is the degree of an individual's subjective sense of control over personal health. Lacking or externalization of control is associated with health compromising behaviors such as drug noncompliance and psychiatric comorbidity [57, 58]. Patients with epilepsy suffer from persistent worries about seizure recurrence, probably arising from a sense of unpredictability of the course of the disorder as well as subjective helplessness. Evidence suggests that the level of depression and anxiety at one year following the diagnosis of epilepsy correlates with the degree to which a patient senses a lack of self-control, but not with the actual number of seizure[58]. This indicates that the assumed uncertainty towards having a seizure could be more disabling than the seizure itself.

Hence, the individuals' role and participation in their own epilepsy condition are emphasized in many psychobehavioral protocols [10, 26-28]. The understanding that seizure occurrence might not be entirely random and unpredictable, but could be triggered by multiple risk factors that are highly controllable, is a motivating insight for many patients since it increases their sense of control over their health condition. This possibly increases the likelihood and openness for them to engage in psychobehavioral therapy, in which their cooperation and practice are central to treatment efficacy.

### **3.4 Learning hypothesis**

Learning to stop the progression of seizure activity is the central idea of most behavioral approaches for PWE. In behaviorism, it is believed that desirable behaviors can be learned and unlearned, strengthened and weakened by applying structural behavioral modification, whereby the process of long-term potentiation (LTP) underlies the synaptic mechanism that correlates with and eventually facilitates the behavioral change[16, 59]. Behaviorists regard seizures as a learned action resulting from a habitual chain of seizure-related behaviors. According to their idea, damaged neurons create only a predisposition for the occurrence of seizures and it is the individuals' response to the dysfunction that is more critical to trigger seizure activity[19, 60]. The learning hypothesis is supported by the observation that the detection of seizure triggers can be facilitated by eliciting the biographical context of the first seizure events[61]. The underlying psychological theme will often resurface in the individual's life and trigger seizures even though the psychological stimulus might be less intense than prior to the first seizure events.

Behavioral approaches for PWE are applied based on this mechanism of action. Aura interruption and systematic desensitization allow individuals to learn new sets of reactions to pre-ictal and early ictal phenomena, which probably lead to neurobiological changes. Presumably, newly learned actions could be able to engage neurons adjacent or contralateral to the area of hyper-excitabile neurons in order to prevent further recruitment and hence the spread of seizure activity. Another hypothesis is that these new behaviors might be able to engage neurons of complex networks in order to prevent loss of consciousness [62-65]. In operant conditioning and EEG biofeedback, desirable EEG patterns were learned by positive and negative reinforcement. These

learning processes are presumably established through LTP[23].

### **3.5 Generic adjustment hypothesis**

Adjustment issues may arise when an individual starts to have seizures, or becomes seizure free after treatment. Research even suggests that the de-novo development of psychogenic non-epileptic seizures might be an underestimated entity in adults after epilepsy surgery, particularly in female patients with preoperative psychiatric comorbidity[66]. Problems with psychosocial adjustment were prominent in PWE in various areas including education, employment, interpersonal relationships, treatment options and even pregnancy. Some psychobehavioral therapies enhance adjustment by providing educational materials, introducing adaptive coping, communication skills, general problem solving skills and non-judgmental awareness [8, 10, 33-36, 67]. These processes are supposed to allow a psychologically healthy transition of an individual's sense of self-identity from being "chronically ill" to being "normal". These forms of interventions might also reduce one's sense of helplessness, as well as secondary gain issues and could promote psychological well-being, which might in turn prevent the increase of seizure activity.

## **4 Is psychobehavioral therapy effective in epilepsy?**

Although many individual trials have reported beneficial effects of psychobehavioral therapy in terms of psychological well-being and seizure control, robust evidence is scarce. This multi-method review section assesses the published evidence from prospective trials in regard to its effectiveness. Table 1 presents a summary of evidence from randomized controlled trials (RCT).

### **4.1 Behavioral approaches**

Behavioral therapies for PWE, which aimed at the seizure reduction, were mainly found in the early literature and many of them had significant methodological deficiencies. Although clinically significant seizure reduction was reported in case studies; the findings were often based on observations or unsophisticated counting without reliable statistical bases[6, 21, 68]. There were also biases in patient selection; therapies were applied on institutionalized patients with mental retardation[21, 68], the positive changes were difficult to generalize. The only RCT was conducted by Dahl et al. in 18 PWE. This un-blinded study reported significant reduction on seizure frequency only in those who received behavioral therapy[69].

The most promising behavioral approach appeared to be EEG biofeedback. Tan and colleagues conducted a meta-analysis on 10 studies using EEG biofeedback on PWE[25]. All studies reported an overall mean reduction of seizures after treatment and 64 out of 87 patients (74%) reported fewer weekly seizures following EEG biofeedback training.

### **4.2 CBT**

In general, the effects of CBT on enhancing psychological well-being are more established than

its role on seizure control as evident by a handful of RCTs and multiple single trials.

A RCT (n = 37) showed significant seizure reduction in CBT group (number of seizure per month at baseline = 6.33; post-treatment = 3.68; 3-month post-treatment = 1.39) compared to no change in control (4.95, 5.00 and 4.42, respectively)[70]. The achievement of complete seizure freedom was reported between 13% and 80% of patients by a few uncontrolled studies [13, 71-73]. The approach based on a 12-session multi-model CBT-like intervention protocol, developed by Andrews and Reiter, is one of the most effective and widely studied CBT approaches into improving seizure control[12, 13, 73]. Despite the positive findings, some studies suggested otherwise and did not demonstrate a significant effect of CBT on seizure control[7, 74].

The effects of CBT on improving psychological states has been consistently demonstrated by RCTs[27, 32, 75]. Patients reported reduction of anxiety and depression symptoms, improvement on adjustment, social functioning as well as quality of life (QOL).

### **4.3 Mind-body approaches**

Studies using mind-body approaches consistently revealed clinical improvement on both seizure control and psychological well-being in PWE [8, 33, 36, 76]. Among various approaches, mindfulness-based ACT has been studied in two RCTs and showed significant effects on improving QOL and seizure control in patients with drug-resistant epilepsy; these positive results were sustained at one year after intervention[8, 33]. Similar finding was reported by Thompson et al. using a mindfulness and cognitive-behavioral based intervention delivered via telephone conference<sup>33</sup>.

### **4.4 Multi-model educative interventions**

Some multi-model interventions include a composition of education, skill-based training with or without cognitive-behavioral elements. Many of them have been formalized into standardized program and are the most extensively and systematically studied psychobehavioral approach for PWE. Despite the heterogeneity of therapeutic content, length of intervention and mode of delivery, these approaches have showed consistent effects on enhancing knowledge and management of seizures. Significant positive effects include improvement on psychological states[9, 10, 15, 77], epilepsy knowledge[9, 14, 15, 77], self-management[9, 10], perceived competencies and social skills[15], contentedness with therapy[10], drug compliance[9] as well as seizure control[9, 10].

### **4.5 Summary of effects using psychobehavioral therapy for PWE**

CBT, mind-body approaches and multi-model educative interventions have consistently demonstrated significant effects on improving psychological well-being, enhancing epilepsy knowledge and adjustment as compared to controls. However, the effects on seizure control remain inconsistent across studies, thus no reliable conclusion can be drawn.

## **5 Challenges in research methodology and way forward**

## **5.1 Deficiencies in study design**

One of the most common shortcomings of these studies relates to the lack of control groups. There was precedence for “significant” findings in uncontrolled studies to fail reproducibility when the intervention was subjected to a controlled trial as reported by Mittan in a comprehensive review[5]. Most RCTs were, however, compromised by the inadequate statistical power, which critically affected the validity of the results.

Second, many psychotherapy protocols consist of a mixture of therapeutic components, each of which could be considered as an independent variable. For instance, a multi-model educative program, Modular Service Package Epilepsy (MOSES), was designed with multiple aims spread over nine separate modules of interventions; each module was featured by a specific topic spanned from imparting knowledge to the adjustment and emotional aspects[10]. Although this form of intervention often demonstrate beneficial effects, it is difficult to clearly delineate the therapeutic components. Moreover, because of the nonspecific therapeutic target, an adequate control group and corresponding outcome measures are difficult to define. In addition, the lack of standardized protocols and intervention goals across studies, even within the same therapeutic approach, made comparison difficult. For instance, some CBT protocols emphasize establishing the patient’s health locus of control[27, 28]; while others are designed to prevent psychiatric comorbidities[7, 31]

The outcome measures employed constitute a third problem. Most studies depend primarily on subjective self-report inventories with a set of yes-no questions or Likert scale ratings to ascertain constructs that are more complicated than a total score can represent, e.g. “depression” and “quality of life”. Whether these inventories are measuring what they are intended to measure may be questionable, even if the inventory has undergone elaborate validation. It is also doubtful whether statistically significant changes actually carry clinical significance. Moreover, measures are not standardized across studies. For example, a widely measured construct, “depression”, was measured by different scales in different studies. This makes any comparisons across studies difficult.

Most studies assessed therapeutic outcomes immediately after intervention or within short term follow-up duration; long-term response data was lacking. The sustainability of therapeutic effects thus remains uncertain.

Finally, although double-blinded design is not practical because patients’ active participation is required, with only a few exceptions, most of them were not even assessor-blinded. Investigators who conducted both the intervention and the outcome assessments are subject to potential biases in regards to subject selection, implementation of interventions as well as the interpretation of results. To date, only the two studies using the Seizures and Epilepsy Education (SEE) program intervention paradigm were conducted, analyzed and reported by individual third parties[9, 14].

## **5.2 Common confounding factors**

The internalization and application of the therapeutic principles are active processes that differ among individuals and require continuous motivation and compliance. These factors, however, are

difficult to measure. These variations often become a major confounding variable that affect the sustainability of therapeutic effects and the correlation of results with the actual therapeutic mediator.

Moreover, concomitant AEDs usage was not described in most studies; hence it remains unclear if it may have acted as a confounder of the effects of psychobehavioral therapy in regards to both psychological and seizure outcomes. Even if AEDs usage was statistically controlled, the long-term effects of psychotherapy could be difficult to determine if simultaneous changes of drug regimens occur. Likewise, progressive syndromes and fluctuations of seizure manifestation affect the interpretation of long term effects of psychobehavioral therapy.

### **5.3 Proposed research strategies**

Assessor-blinded randomized controlled trials with sufficient statistical power appear to be a necessary design to evaluate the efficacy of specific psychobehavioral paradigms. Control group should be carefully designed to delineate the nonspecific effects of the target therapy to be evaluated. The use of waitlist control, nonspecific supportive treatment or treatment as usual is recommended, which resemble the trial design used in psychotherapy research on traditional psychiatric settings that has demonstrated consistent validity. The relative efficacy of different interventions or their respective components can be separately compared in a second step [78-80].

The measurement approach of outcomes should be optimized. Objective measures of neurobiological correlates following psychobehavioral therapy should be used. For instance, serum levels as a measure of drug compliance could be utilized. Salivary cortisol level may be used as an objective neurobiological correlate of stress[81]. MRI or fMRI could be included as a morphological evidence of the hypothesis of neuroplasticity following psychobehavioral therapy [82-84]. These objective measures are important to determine the extent and mechanisms of psychological agents on PWE in a so called “top-down” process of how our mind changes our brain. Furthermore, confounding factors should be statistically controlled, including changes of clinical characteristics. Patients’ application and practices of the therapeutic skills should be objectively monitored, for instance by journaling and obtaining logs of practices [13, 73].

Randomized controlled trials could be supplemented by other types of research designs such as qualitative studies that could potentially facilitate the correlation of subjective qualia of seizure self-control phenomena and objective measurements, and case reports that might explore specific psychological mediators of seizure self-control[85].

## **Summary Response**

### **1 What psychobehavioral therapies are/may be useful in epilepsy?**

Behavioral approaches, cognitive-behavioral therapy, and mind-body approaches are the most widely applied psychobehavioral approaches for PWE. Some interventions employ a composition of therapeutic elements from multiple approaches.

## **2 What are the hypothesized mechanisms of action of psychobehavioral therapy for epilepsy?**

The major hypothesized mechanisms of action include stress reduction, treating psychiatric comorbidity, self-efficacy promotion including the development of seizure interruption techniques, learning hypothesis and generic adjustment hypothesis.

## **3 Is psychobehavioral therapy effective in epilepsy?**

Most psychobehavioral therapies aim at enhancing psychological well-being and seizure control; evidences for the former are more established. Because of major methodological deficiencies, reliable evidence on its role on seizure control is scarce.

## **4 What are the methodological challenges associated with the scientific study of psychobehavioral therapies and the proposed strategies for meeting those challenges?**

Major methodological problems include deficiencies in study designs and multiple confounding factors. Blinded randomized controlled trials with sufficient power and carefully defined therapeutic components corresponding with objective and subjective outcome measures are advisable. Confounding factors should also be rigorously controlled. Supplementary types of research designs, such as qualitative research and case reports, could be employed to facilitate the correlation of subjective qualia of seizure self-control phenomena and objective measurements.

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### **References**

1. Jones, N.C. and T.J. O'Brien, *Stress, epilepsy, and psychiatric comorbidity: how can animal models inform the clinic?* *Epilepsy Behav*, 2013. **26**(3): p. 363-9.
2. Kanner, A.M., et al., *Depression and epilepsy: epidemiologic and neurobiologic perspectives that may explain their high comorbid occurrence.* *Epilepsy Behav*, 2012. **24**(2): p. 156-68.
3. Mula, M., *Epilepsy: Bidirectional link between epilepsy and psychiatric disorders.* *Nat Rev Neurol*, 2012. **8**(5): p. 252-3.
4. Ramaratnam, S., G.A. Baker, and L.H. Goldstein, *Psychological treatment for epilepsy [Review].* *Cochrane Database Syst Rev*, 2008(3).
5. Mittan, R.J., *Psychosocial treatment programs in epilepsy: a review.* *Epilepsy Behav*, 2009. **16**(3): p. 371-80.
6. Parrino, J.J., *Reduction of seizures by desensitization.* *Journal of Behavior Therapy and*

- Experimental Psychiatry, 1971. **2**(3): p. 215-218.
7. Goldstein, L.H., et al., *Cognitive behaviour therapy with adults with intractable epilepsy and psychiatric co-morbidity: preliminary observations on changes in psychological state and seizure frequency*. Behav Res Ther, 2003. **41**(4): p. 447-460.
  8. Lundgren, T., et al., *Evaluation of acceptance and commitment therapy for drug refractory epilepsy: a randomized controlled trial in South Africa--a pilot study*. Epilepsia, 2006. **47**(12): p. 2173-9.
  9. Helgeson, D.C., et al., *Sepulveda Epilepsy Education: the efficacy of a psychoeducational treatment program in treating medical and psychosocial aspects of epilepsy*. Epilepsia, 1990. **31**(1): p. 75-82.
  10. May, T.W. and M. Pfafflin, *The efficacy of an educational treatment program for patients with epilepsy (MOSES): results of a controlled, randomized study. Modular Service Package Epilepsy*. Epilepsia, 2002. **43**(5): p. 539-49.
  11. Helde, G., et al., *A structured, nurse-led intervention program improves quality of life in patients with epilepsy: a randomized, controlled trial*. Epilepsy Behav, 2005. **7**(3): p. 451-7.
  12. Schmid-Schönbein, C., *Improvement of seizure control by psychological methods in patients with intractable epilepsies*. Seizure : the journal of the British Epilepsy Association, 1998. **7**(4): p. 261-270.
  13. Michaelis, R., W. Schonfeld, and S.M. Elsas, *Trigger self-control and seizure arrest in the Andrews/Reiter behavioral approach to epilepsy: a retrospective analysis of seizure frequency*. Epilepsy Behav, 2012. **23**(3): p. 266-71.
  14. Shore, C.P., S.M. Perkins, and J.K. Austin, *The Seizures and Epilepsy Education (SEE) program for families of children with epilepsy: a preliminary study*. Epilepsy Behav, 2008. **12**(1): p. 157-64.
  15. Lewis, M.A., et al., *Randomized trial of a program to enhance the competencies of children with epilepsy*. Epilepsia, 1990. **31**(1): p. 101-9.
  16. Grossberg, J.M., *Behavior therapy: A review*. Psychological Bulletin, 1964. **62**(2): p. 73-88.
  17. Rescorla, R.A., *Pavlovian conditioning and its proper control procedures*. Psychological Review, 1967. **74**(1): p. 71-80.
  18. Leitenberg, H., et al., *Contribution of selective positive reinforcement and therapeutic instructions to systematic desensitization therapy*. Journal of Abnormal Psychology, 1969. **74**(1): p. 113-118.
  19. Lavender, A., *A Behavioural Approach to the Treatment of Epilepsy*. Behavioural and Cognitive Psychotherapy (New Series), 1981. **9**(03): p. 231-243.
  20. Schmid-Schönbein, C., *Improvement of seizure control by psychological methods in patients with intractable epilepsies*. Seizure, 1998. **7**(4): p. 261-270.
  21. Iwata, B.A. and A.M. Lorentzson, *Operant control of seizure-like behavior in an institutionalized retarded adult*. Behavior Therapy, 1976. **7**(2): p. 247-251.

22. Sterman, M.B., *Basic concepts and clinical findings in the treatment of seizure disorders with EEG operant conditioning*. Clinical electroencephalography, 2000. **31**(1): p. 45-55.
23. Sterman, M.B. and T. Egner, *Foundation and Practice of Neurofeedback for the Treatment of Epilepsy*. Applied Psychophysiology and Biofeedback, 2006. **31**(1): p. 21-35.
24. Sterman, M.B., L.R. Macdonald, and R.K. Stone, *Biofeedback Training of the Sensorimotor Electroencephalogram Rhythm in Man: Effects on Epilepsy*. Epilepsia, 1974. **15**(3): p. 395-416.
25. Tan, G., et al., *Meta-Analysis of EEG Biofeedback in Treating Epilepsy*. Clinical EEG and Neuroscience, 2009. **40**(3): p. 173-179.
26. Reiter, J.M. and D.J. Andrews, *A neurobehavioral approach for treatment of complex partial epilepsy: efficacy*. Seizure : the journal of the British Epilepsy Association, 2000. **9**(3): p. 198-203.
27. Au, A., et al., *Cognitive-behavioral group treatment program for adults with epilepsy in Hong Kong*. Epilepsy & Behavior, 2003. **4**(4): p. 441-446.
28. Andrews, D.J., J.M. Reiter, and C. Janis, *Taking control of your epilepsy. A workbook for patients and professionals*. . 1987, Santa Rosa: The Basics Publishing Company.
29. Forman, E.M., et al., *A Randomized Controlled Effectiveness Trial of Acceptance and Commitment Therapy and Cognitive Therapy for Anxiety and Depression*. Behavior Modification, 2007. **31**(6): p. 772-799.
30. Walker, E.R., et al., *Formative and process evaluations of a cognitive-behavioral therapy and mindfulness intervention for people with epilepsy and depression*. Epilepsy Behav, 2010. **19**(3): p. 239-46.
31. Macrodimitris, S., et al., *Group cognitive-behavioral therapy for patients with epilepsy and comorbid depression and anxiety*. Epilepsy & Behavior, 2011. **20**(1): p. 83-88.
32. Martinović, Ž., P. Simonović, and R. Djokić, *Preventing depression in adolescents with epilepsy*. Epilepsy & Behavior, 2006. **9**(4): p. 619-624.
33. Lundgren, T., et al., *Acceptance and Commitment Therapy and yoga for drug-refractory epilepsy: a randomized controlled trial*. Epilepsy Behav, 2008. **13**(1): p. 102-8.
34. Lundgren, T., J. Dahl, and S.C. Hayes, *Evaluation of mediators of change in the treatment of epilepsy with acceptance and commitment therapy*. J Behav Med, 2008. **31**(3): p. 225-35.
35. Dahl, J. and T. Lundgren, *Analysis and treatment of epilepsy using mindfulness, acceptance, values, and countermeasures*, in *Mindfulness and acceptance in behavioral medicine*, L.M. McCracken, Editor. 2011, New Harbinger Publications, Inc.: Oakland, CA.
36. Thompson, N.J., et al., *Distance delivery of mindfulness-based cognitive therapy for depression: project UPLIFT*. Epilepsy Behav, 2010. **19**(3): p. 247-54.
37. Sathyaprabha, T.N., et al., *Modulation of cardiac autonomic balance with adjuvant yoga therapy in patients with refractory epilepsy*. Epilepsy Behav, 2008. **12**(2): p. 245-52.
38. Ramaratnam, S. and K. Sridharan, *Yoga for epilepsy*. Cochrane Database Syst Rev, 2000. **3**.
39. Yardi, N., *Yoga for control of epilepsy*. Seizure, 2001. **10**(1): p. 7-12.

40. Polak, E.L., et al., *Behavioral intervention as an add-on therapy in epilepsy: Designing a clinical trial*. *Epilepsy Behav*, 2012. **25**(4): p. 505-510.
41. Hermann, B. and A. Jacoby, *The psychosocial impact of epilepsy in adults*. *Epilepsy Behav*, 2009. **15 Suppl 1**: p. S11-6.
42. Novakova, B., et al., *The role of stress as a trigger for epileptic seizures: A narrative review of evidence from human and animal studies*. *Epilepsia*, 2013: p. n/a-n/a.
43. Fried, R., *Breathing training for the self-regulation of alveolar CO<sub>2</sub> in the behavioral control of idiopathic epileptic seizures*, in *The Neurobehavioral Treatment of Epilepsy*, D.I. Mostofsky and Y. Loyning, Editors. 1993, Lawrence Erlbaum: Hillsdale, NJ. p. 19-66.
44. Rousseau, A., B. Hermann, and S. Whitman, *Effects of progressive relaxation on epilepsy: analysis of a series of cases*. *Psychol Rep*, 1985. **57**(3 Pt 2): p. 1203-12.
45. Dahl, J.A., L. Melin, and L. Lund, *Effects of a Contingent Relaxation Treatment Program on Adults with Refractory Epileptic Seizures*. *Epilepsia*, 1987. **28**(2): p. 125-132.
46. Puskarich, C.A., et al., *Controlled examination of effects of progressive relaxation training on seizure reduction*. *Epilepsia*, 1992. **33**(4): p. 675-80.
47. Nagai, Y., et al., *Clinical efficacy of galvanic skin response biofeedback training in reducing seizures in adult epilepsy: a preliminary randomized controlled study*. *Epilepsy Behav*, 2004. **5**(2): p. 216-23.
48. Passaro, E.A., *Psychiatric comorbidity in epilepsy*. *Primary Psychiatry*, 2003. **10**(10): p. 72-79.
49. Gaitatzis, A., M.R. Trimble, and J.W. Sander, *The psychiatric comorbidity of epilepsy*. *Acta Neurol Scand*, 2004. **110**(4): p. 207-20.
50. Chang, Y.T., et al., *Bidirectional relation between schizophrenia and epilepsy: a population-based retrospective cohort study*. *Epilepsia*, 2011. **52**(11): p. 2036-42.
51. Hesdorffer, D.C., et al., *Depression and suicide attempt as risk factors for incident unprovoked seizures*. *Ann Neurol*, 2006. **59**(1): p. 35-41.
52. Kanner, A.M., et al., *A lifetime psychiatric history predicts a worse seizure outcome following temporal lobectomy*. *Neurology*, 2009. **72**(9): p. 793-9.
53. Devinsky, O., *Psychiatric comorbidity in patients with epilepsy: implications for diagnosis and treatment*. *Epilepsy & Behavior*, 2003. **4**: p. 2-10.
54. Kanner, A.M., *Depression in epilepsy: prevalence, clinical semiology, pathogenic mechanisms, and treatment*. *Biological Psychiatry*, 2003. **54**(3): p. 388-398.
55. Kanner, A.M., *Depression and epilepsy: Do glucocorticoids and glutamate explain their relationship?* *Current Neurology and Neuroscience Reports*, 2009. **9**(4): p. 307-12.
56. Kanner, A.M. and A. Balabanov, *Depression and epilepsy: How closely related are they?* *Neurology*, 2002. **58**(8 suppl 5): p. S27-S39.
57. Gopinath, B., et al., *A questionnaire survey about doctor-patient communication, compliance and locus of control among South Indian people with epilepsy*. *Epilepsy Research*, 2000. **39**(1): p. 73-82.

58. Velissaris, S.L., et al., *Psychological trajectories in the year after a newly diagnosed seizure*. *Epilepsia*, 2012. **53**(10): p. 1774-81.
59. Whitlock, J.R., et al., *Learning Induces Long-Term Potentiation in the Hippocampus*. *Science*, 2006. **313**(5790): p. 1093-1097.
60. Lockard, J. and A. Ward, *Epilepsy: a Window to Brain Mechanism*. . 1980, New York: Raven Press. 51-68.
61. Koutsogiannopoulos, S., et al., *Stressors at the onset of adult epilepsy: implications for practice*. *Epileptic Disord*, 2009. **11**(1): p. 42-7.
62. Englot, D.J. and H. Blumenfeld, *Consciousness and epilepsy: why are complex-partial seizures complex?* *Prog Brain Res*, 2009. **177**: p. 147-70.
63. Danielson, N.B., J.N. Guo, and H. Blumenfeld, *The default mode network and altered consciousness in epilepsy*. *Behav Neurol*, 2011. **24**(1): p. 55-65.
64. Yu, L. and H. Blumenfeld, *Theories of impaired consciousness in epilepsy*. *Ann N Y Acad Sci*, 2009.
65. Blumenfeld, H. and J. Taylor, *Why do seizures cause loss of consciousness?* *Neuroscientist*, 2003. **9**(5): p. 301-10.
66. Markoula, S., et al., *De novo psychogenic nonepileptic attacks after adult epilepsy surgery: An underestimated entity*. *Epilepsia*, 2013: p. n/a-n/a.
67. Piazzini, A., et al., *Coping strategies in epilepsy: 50 drug-resistant and 50 seizure-free patients*. *Seizure*, 2007. **16**(3): p. 211-217.
68. Wright, L., *Aversive conditioning of self-induced seizures*. *Behavior Therapy*, 1973. **4**(5): p. 712-713.
69. Dahl, J., L. Melin, and L. Lund, *Effects of a contingent relaxation treatment program on adults with refractory epileptic seizures*. *Epilepsia*, 1987. **28**(2): p. 125-32.
70. McLaughlin, D.P. and K. McFarland, *A randomized trial of a group based cognitive behavior therapy program for older adults with epilepsy: the impact on seizure frequency, depression and psychosocial well-being*. *J Behav Med*, 2011. **34**(3): p. 201-7.
71. Wolf, P. and N. Okujava, *Possibilities of non-pharmacological conservative treatment of epilepsy*. *Seizure : the journal of the British Epilepsy Association*, 1999. **8**(1): p. 45-52.
72. Joy Andrews, D., et al., *A neurobehavioral treatment for unilateral complex partial seizure disorders: a comparison of right- and left-hemisphere patients*. *Seizure*, 2000. **9**(3): p. 189-97.
73. Elsas, S.M., et al., *Aura interruption: the Andrews/Reiter behavioral intervention may reduce seizures and improve quality of life - a pilot trial*. *Epilepsy Behav*, 2011. **22**(4): p. 765-72.
74. Tan, S.Y. and J. Bruni, *Cognitive-behavior therapy with adult patients with epilepsy: a controlled outcome study*. *Epilepsia*, 1986. **27**(3): p. 225-33.
75. Davis, G.R., et al., *Cognitive-behavioral treatment of depressed affect among epileptics: Preliminary findings*. *Journal of Clinical Psychology*, 1984. **40**(4): p. 930-935.
76. Walker, E.R., et al., *Formative and process evaluations of a cognitive-behavioral therapy and*

- mindfulness intervention for people with epilepsy and depression*. *Epilepsy & Behavior*, 2010. **19**(3): p. 239-246.
77. Olley, B.O., H.O. Osinowo, and W.R. Brieger, *Psycho-educational therapy among Nigerian adult patients with epilepsy: a controlled outcome study*. *Patient Educ Couns*, 2001. **42**(1): p. 25-33.
  78. Bisson, J. and M. Andrew, *Psychological treatment of post-traumatic stress disorder (PTSD)*. *Cochrane Database Syst Rev*, 2007. **18**(3).
  79. Furukawa, T.A., N. Watanabe, and R. Churchill, *Combined psychotherapy plus antidepressants for panic disorder with or without agoraphobia*. *Cochrane Database Syst Rev*, 2007. **24**(1).
  80. Hunot, V., et al., *Psychological therapies for generalised anxiety disorder*. *Cochrane Database Syst Rev*, 2007. **24**(1).
  81. Stawski, R.S., et al., *Associations among daily stressors and salivary cortisol: Findings from the National Study of Daily Experiences*. *Psychoneuroendocrinology*, 2013. **38**(11): p. 2654-65.
  82. Yoshimura, S., et al., *Cognitive behavioral therapy for depression changes medial prefrontal and ventral anterior cingulate cortex activity associated with self-referential processing*. *Soc Cogn Affect Neurosci*, 2013. **11**: p. 11.
  83. Guleria, A., et al., *Effect of "SOHAM" meditation on the human brain: An fMRI study*. *Psychiatry Research: Neuroimaging*, (0).
  84. Allen, M., et al., *Cognitive-affective neural plasticity following active-controlled mindfulness intervention*. *J Neurosci*, 2012. **32**(44): p. 15601-10.
  85. Petitmengin, C., V. Navarro, and V. Quyen Mle, *Anticipating seizure: pre-reflective experience at the center of neuro-phenomenology*. *Conscious Cogn*, 2007. **16**(3): p. 746-64.

Fig 1: The seizure event as the end-point of seizure precipitating factors.

Fig 1 conceptualizes the hypothesized interaction between the seizure threshold and the accumulation of seizure activity in the brain of an individual with an increased epileptic disposition. The seizure event is pictured as the end point of proceeding factors that appear in the schematic order of temporal proximity to the actual seizure event. The proceeding factors can be correlated with various stages of the electrophysiological build-up of seizure activity: The disposition to experience epileptic seizures may increase due to alterations on a cellular or macroscopic level that can originate from various etiologies e.g. traumatic brain injury, genetic mutations. Stress is conceptualized to act as a non-specific seizure precipitant that corresponds with various neuroendocrinologic changes. Seizure risk factors are non-specific seizure precipitants that affect the brain as a whole, e.g. missed sleep, missed meal, omitted medication. Seizure triggers are specific seizure precipitants that induce epileptic activity by being physiologically processed in an area adjacent to or overlapping with epileptogenic neurons. The nature of the neuronal network in which the seizure trigger is processed may be simple (e.g. visual stimuli) or complex (e.g. cognitive or emotional triggers). Seizure warning signs include all subjective phenomena that are reported by individuals to occur reliably prior to a generalized or complex partial seizure event. Those seizure warning signs may appear days and hours (usually referred to as “prodrome”) or minutes to seconds (usually referred to as “aura”) prior to the seizure. The latter is usually regarded as the subjective correlate of the initial appearance of confined epileptic activity in the brain. If the accumulated seizure activity crosses the seizure threshold, a seizure occurs. All or some of these factors may operate for a given seizure in an individual.

Fig 2a: Model of interaction between multifactorially conditioned accumulation of seizure activity, seizure threshold and seizure occurrence

Fig 2a shows the conceptualized interaction between the seizure threshold and fluctuating levels of seizure activity in the brain of an individual with an increased epileptic disposition. The rising and falling levels of seizure activity are hypothesized to depend on the individual's reactions to multiple seizure precipitating factors as well as to seizure warning signs.

Fig 2b: Model of hypothesized effects of psychobehavioral interventions for epilepsy

Fig 2b illustrates the hypothesized short-term and long-term effects of psychobehavioral interventions. Individuals with an increased epileptic disposition learn to proactively avoid the build-up of seizure activity per various strategies that are being employed by psychobehavioral interventions. This is hypothesized to reduce non-specific seizure precipitants and specific seizure precipitants in the short-term. Long-term practice of the psychobehavioral strategies leads to habituation results in use-dependent changes of neuronal circuits. This change of brain structure is hypothesized to correlate with a decrease of the individual brain's epileptic disposition.

Fig 1:

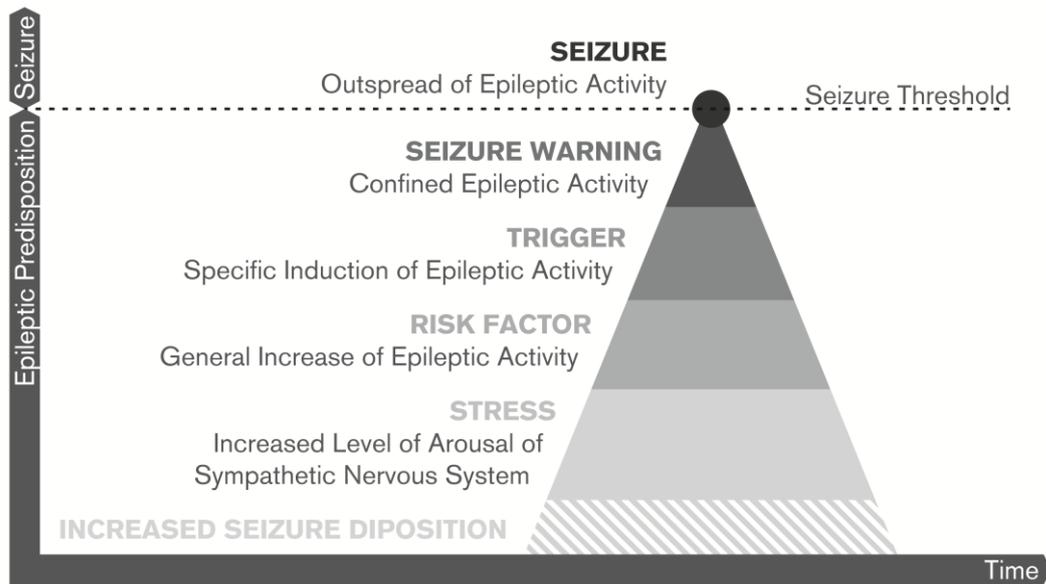


Fig 2a:

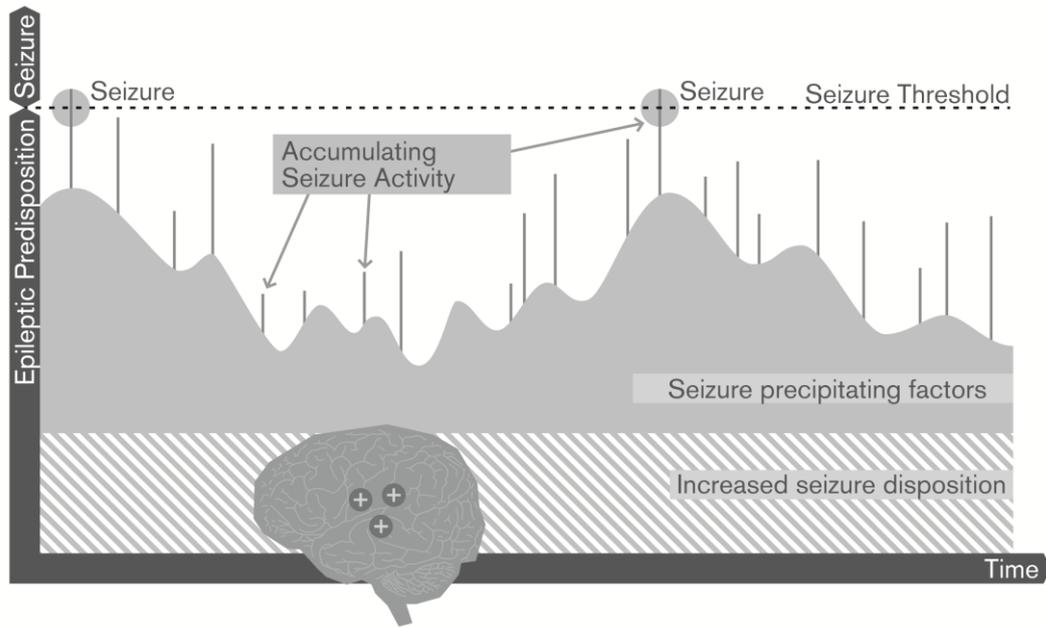


Fig 2b:

