Epilepsy and its Management in Relation to Psychiatry
Amir Mufaddel*
Community Mental Health Services, Behavioural Sciences Institute, Al Ain Hospital, United Arab Emirates University, United Arab Emirates

Abstract
The relation between psychiatry and epilepsy remains one of the topics that have been continuously attracting attention in medical literature since the time of Hippocrates. Forced normalization was later conceptualized when biological treatment for psychiatric disorders was developed. Recently, several reports were published regarding possible adverse psychiatric effects of antiepileptic medications. The picture is further complicated by the fact that psychotropic medications, which may be used for treatment of psychiatric disorders in epilepsy, can have negative effects on seizure control.

This paper aimed to review psychiatric presentations in individuals with epilepsy and the adverse psychiatric effects of antiepileptic medications. The paper also briefly discusses available treatment options with regards of better psychotropic treatment options.

Recent developments in this field led to the observation that psychiatric symptoms, in relation to seizure, can be categorized into the following types pre-ictal, post-ictal, inter-ictal and ictal symptoms. Depression and anxiety are commonly associated with epilepsy and can occur as pre-ictal, ictal, post-ictal or inter-ictal symptoms. Depression can be severe and suicidal behavior has been reported as one of the common psychiatric associations with epilepsy and has also been linked to use of some antiepileptic drugs. Psychosis can occur as an acute or chronic form. Transient pleomorphic postictal psychosis particularly observed in those with hippocampal sclerosis. There is an increased risk of developing schizophrenia and schizophrenia-like psychosis in patients with epilepsy. Psychiatric symptoms can also emerge during use of some of the antiepileptic drugs. Adverse psychiatric events include depression, psychosis, suicidal risk, and cognitive impairment. Some of the psychotropic medications are known to be epileptogenic. Therefore, selection of psychotropic medications should be based on their possible effects on seizures and those known to be epileptogenic should be avoided.

Keywords: Epilepsy; Psychosis; Mood disorders; Anxiety; Psychotropic; Antiepileptic

Introduction
Epilepsy is a chronic disorder characterized by recurrent seizures, or a seizure of a paroxysmal brain dysfunction due to excessive neuronal discharge. The Greek physician Hippocrates wrote the first book on epilepsy more than 2000 years ago, and he stated that “Most melancholics usually also become epileptics and epileptics melancholics” [1].

The association between psychological symptoms and epilepsy can be related to several factors such as the possibility of shared etiology, stigma and psychosocial factors related to epilepsy, and the adverse psychiatric reactions of antiepileptic drugs. Some of the psychotropic medications are epileptogenic; and this should also be considered when prescribing a psychotropic medication for an epileptic patient. Psychiatric disorders in epilepsy can be classified, based on their relationship to seizures, into the following categories: pre-ictal symptoms (occur before seizure/ prodromal states and mood disturbance), inter-ictal psychiatric disorders occurring between seizures; ictal psychiatric symptoms occurring during the seizure events; and post-ictal psychiatric symptoms following seizures (Table 1). Occasionally, an altered mental state can be the only sign of non-convulsive status epilepticus such as that of complex partial or a seizure of a paroxysmal brain dysfunction due to excessive epileptogenic should be avoided.

Psychiatric comorbidity is common among patients with epilepsy with rates increasing at least two-folds in that attending specialist care. Psychiatric conditions that are commonly associated with epilepsy are depressive disorder, anxiety disorder, attention-deficit hyperactivity disorder and psychoses [3]. For example, a recent study including 319 patients with focal epilepsy suggested that about 58% of patients with focal epilepsy have either a current or past history of a psychiatric disorder. Depression was found to be the most common psychiatric diagnosis occurring in 32.6% of patients. Other psychiatric diagnoses were psychotic disorders (7.2%), anxiety (6.9%), substance misuse/dependence (3.1%), somatoform disorders (4.7%), and personality disorders (13.8%) [4]. Data from the California Health Interview Survey conducted in 2005, including data from 604 adult participants with history of epilepsy, indicated that 27% had psychological distress associated with epilepsy, and 84% of them needed mental health care, but only 57% have been seen by a mental health professional [5]. A recent population-based cohort study comparing 938 patients whom were newly diagnosed with epilepsy and 518,748 subjects without epilepsy revealed that the incidence of psychiatric disorders for people with epilepsy was 94.1 per 1000 persons-year and for those without epilepsy were 22.6 per 1000 persons-year. The epilepsy cohort had the highest risks in mental retardation bipolar disorder and alcohol or drug induced psychosis. The highest risk occurred in the first year following the diagnosis of epilepsy [6].

Behavioral aspects of epilepsy were well-studied in temporal lobe epilepsy and mesial temporal lobe epilepsy compared with other types of epilepsy such as frontal lobe epilepsy. This may be due to the fact that temporal lobe epilepsy represents the majority of cases with

*Corresponding author: Amir Mufaddel, Community Mental Health Services, Behavioural Sciences Institute, United Arab Emirates University, Al Ain Hospital, P.O. Box 1006, United Arab Emirates, Tel: 00971556292529; E-mail: khaliaamir@yahoo.co.uk
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Prodromal states that occur several hours or days before seizures may include confusion, disorientation, irritability, and agitation.

Ictal psychiatric symptoms that are directly related to seizure include cognitive symptoms, delirium, impaired consciousness, and automatisms. Violence is rare.

Post-ictal psychiatric symptoms that occur hours following seizures may include cognitive symptoms and delirium. Psychosis is rare.

Inter-ictal psychiatric disorders occur between seizures and may include psychiatric disturbances that occur between seizures.

Mood and Anxiety Disorders

Depression and anxiety are common findings in patients with epilepsy occurring in 9-37% and 11-25% of patients respectively. Higher rates of depression are particularly associated with uncontrolled epilepsy [8].

Symptoms of anxiety are commonly associated with temporal lobe epilepsy, and they can occur as inter-ictal, ictal, or post-ictal anxiety states. Anxiety and fear are the most common form of ictal affect and can be associated with palpitations, rising epigastric sensation, mydriasis, diaphoresis, and pallor [1]. A nationally representative population-based study, recently conducted in England, documented that about one-third of patients with epilepsy had anxiety or depressive disorders, and the conditions which were particularly associated with epilepsy included social phobia, agoraphobia, generalized anxiety disorder, depression, and suicidality [9].

The presentation of depression in patients with epilepsy may occur with the same criteria of primary depressive disorders, but may also occur with atypical pleomorphic features [10]. Barry, in his review of mood disorders in epilepsy, identified some risk factors for developing mood disorders in people with epilepsy. These include neurobiological factors, iatrogenic factors (effect of antiepileptic drugs), and psychosocial factors. The depletion of biogenic amines and GABA plays a role in the pathogenesis of both depression and epilepsy. Additionally, frontal lobe dysfunction and decrease in metabolic activity has been recognized in epilepsy as well as in idiopathic depression. Use of antiepileptic drugs may be associated with developing symptoms of mood disorders. For example, depression can be induced by barbiturates, topiramate, tiagabine, vigabatrin, and felbamate. Hypomanic states has also been reported with antiepileptic medications such as zonisamide. Psychosocial factors that may predict developing depression in patients with epilepsy include level of social support, stigma, external locus of control, and poor vocational adjustment [11]. The relationship between mood disorders and epilepsy seems to be bidirectional. Patients with epilepsy have increased risk of depression, and patients with depression may also have a risk of developing epilepsy. Both conditions share common pathogenic mechanisms including changes in neurotransmitters such as serotonin, norepinephrine, glutamate, and GABA [10].

Patients with history of depression and epilepsy may have more frequent focal epilepsies arising from the temporal lobe. There was no difference between patients with depression and controls with respect to severity of epilepsy [12].

Mood disorders may also occur following surgery for epilepsy. Therefore, psychiatric follow-up and evaluation is suggested after surgery to avoid complications such as risk of suicide. Study including 38 patients with history of surgery for intractable temporal lobe epilepsy suggested a relation between history of preoperative postictal psychosis and with developing manic or depressive episodes following surgery particularly for those who had left sided lobectomy [13].

Risk of suicide is four times greater than the general population and deliberate self harm is 6 times more frequent in patients with epilepsy particularly in those who have temporal lobe epilepsy and following surgical treatment [2].

Psychosis

Psychotic symptoms in epilepsy usually occur in the inter-ictal or post-ictal states, and they are rarely occurring in the ictal states. Therefore, psychosis in patients with epilepsy can be divided into three types including chronic and acute interictal psychoses and postictal psychosis. The mentioned types represent about 95% of psychotic symptoms that occur in patients with epilepsy [14].

Transient postictal psychotic episodes are more common in patients with unilateral hippocampal sclerosis than those who have negative MRI findings [15]. About 88% of patients presenting with postictal psychosis were found to have a relative increase of right temporal perfusion and SPECT findings suggested a trend of right-sided temporal predominance regardless of the location (right sided or left sided) of the pathology suspected during a non-psychotic state [16]. Psychotic symptoms usually occur 12-72 hours following seizure and can be delayed up to one week. The Postictal psychosis has pleomorphic...
presentation which may include delusions of persecution, grandiosity, and reference. Delusions can also be of somatic and religious types. Catatonia and hallucinations can also occur; and mood symptoms are often prominent [17].

Chronic psychotic disorders such as schizophrenia and related phososes, should be considered as a possible differential diagnosis for psychotic symptoms in patients with epilepsy. There is an increased risk of schizophrenia and schizophrenia-like psychosis in patients with epilepsy, which increases with age, presence of family history of psychosis, family history of epilepsy and developing epilepsy at a later age. The increased risk for schizophrenia or schizophrenia-like psychosis was not related to gender or type of epilepsy [18]. Patients who have history of Schizophrenia and epilepsy were found to have an earlier onset and more severe forms of epilepsy characterized by history of status epilepticus, and multiple seizure types compared to controls who have epilepsy but no psychiatric symptoms. They have shown various EEGs abnormalities including temporal lobe discharges but there was no lateralization to either side [21].

**Aggression**

In relation to seizures, aggression in people with epilepsy can be ictal, post-ictal or inter-ictal, and it is a kind of affective aggression that is usually associated with high emotional arousal, anger or fear [19]. One large series suggest that ictal violence is extremely rare and was found to be less than 0.3 % [20]. Post-ictal aggression is rare but more common than aggression occurring during the seizure episodes. Post-ictal aggression may be related more to male gender and psychotic experiences and can be recurrent and stereotyped following a cluster of seizures than after a single ictus. Patients are usually remorseful in the inter-ictal period [21].

Aggressive behavior in patients with temporal lobe epilepsy is associated with male gender, early onset of seizures, long-standing behavioral problems, low IQ scores and poor occupational records [22]. Brain structures that mediate aggression include periaqueductal grey matter, hypothalamus, amygdala, limbic structures, and frontal lobes [19.22-24].

Examination of 50 patients with temporal lobe epilepsy of which 25 patients with and 25 without history of intermittent explosive disorder, revealed no higher prevalence of amygdala sclerosis in the aggressive patients. However, 20% of patients with temporal lobe epilepsy and aggression had severe amygdala atrophy, and 28% had left temporal lobe lesions affecting the amygdala and periamygdaloid structures [19].

Frontal lobe epilepsy may present with disinhibition, loss of impulse control, and impulsive aggression. Such presentations can be related to the orbitofrontal cortex which connects the frontal and limbic structures which are involved in the control of behavior. Deviant social behavior and affective symptoms can also be associated with damage to the anterior cingulated gyrus which has connections to the amygdala [7].

**Cognitive Symptoms and Pervasive Developmental Disorders**

Cognitive symptoms are relatively common in patients suffering from epilepsy, particularly chronic and refractory forms of epilepsy [25]. Study of learning disabilities in children with epilepsy indicated that 48% had learning disability reflected in at least one academic area by using the IQ-achievement discrepancy definition. By using low-achievement definitions, 41% to 62% of children had difficulties in at least one academic area [26]. Progressive cognitive decline has been reported in patients with epilepsy. Factors that can be implicated include the nature, timing, and course of cognitive impairment in addition to factors related to epilepsy such as chronic symptoms, and refractory epilepsy [27].

There is an increased incidence of Attention Deficit Hyperactivity Disorder (ADHD) in patients with epilepsy which is estimated to be about 7.76 cases per 1000 person-years compared with an incidence of 3.22 in patients without history of epilepsy. Similarly, the incidence of epilepsy is greater in patients suffering from ADHD approaching 3.24 cases per 1000 person-year, compared with an incidence of 0.78 in those without history of ADHD [28].

There is a high prevalence of autism in children with epilepsy and there is also a high prevalence of epilepsy in children diagnosed with autism. Both conditions are also associated with intellectual disabilities and this observation suggests that intellectual disability can be a possible connection between the two disorders [29]. Study of 150 individuals who were previously diagnosed with childhood autism indicated that epilepsy developed in 22% of participants when they were later assessed at the age of 22 years and above. The onset of seizures was after 10 years of age for most of the cases; and they were predominantly generalized tonic-clonic seizures (88%) [30].

**Other Psychiatric Comorbidities**

Example of other psychiatric problems that may exist in patients with epilepsy are the dissociative symptoms which can manifest as psychogenic non-epileptic seizures, personality changes and sexual dysfunction such as decreased libido and loss of sexual interest.

Dissociative symptoms manifesting as seizures were reported in 9% to 50% of patients in specialist epilepsy centers [31]. Psychogenic non-epileptic seizures (pseudoseizures) and dissociative states are very difficult to differentiate from epileptic seizures. Video electroencephalogram is the gold standard for proper diagnosis, but this procedure is expensive and time-consuming [32]. Diagnosis of pseudoseizure is suggested by identifiable psychosocial precipitating factors, history of physical or sexual abuse, family history of psychiatric problems, unusual or variable pattern of seizures, and absence of autonomic signs [2].

There is a debate about the presence of certain personality characteristics linked to epilepsy. Historically, the epileptic personality has been described by the following characteristics: egocentricity, religiosity, irritability, quarrelsome, and tricky thought processes. Later surveys suggest that only minorities of patients with epilepsy (particularly those with temporal lobe lesions) have serious difficulties related to personality [2]. Some studies suggest that personality disorders can also occur in other types of epilepsy and not only temporal lobe epilepsy. For example, one study including 43 patients with juvenile myoclonic epilepsy has shown that personality disorders were present in 23% of patients [33].

Although reproductive and sexual lives are normal in most of the patients with epilepsy, around 20% to 30% of females have dysfunctions in libido, arousal, and orgasm. Sexual problems in males with epilepsy
Etiology may be related to forced normalization or

Both epilepsy and antiepileptic drugs may be increased risk with rapid dose titration (e.g.,

Limited evidence

May be related to forced normalization (psychotic symptoms emerge when frequency of seizures diminished).

### Psychiatric Side Effects of Antiepileptic Drugs

Despite the fact that some antiepileptic medications have positive effects in improving psychiatric symptoms (commonly used as mood stabilizers, such as valproate, carbamazepine, lamotrigine; and some can improve anxiety symptoms), but they can also be associated with adverse psychiatric events including behavioral problems (most commonly), affective symptoms and rarely psychosis [35]. Table 2 summarizes some of the adverse psychiatric events that can be associated with antiepileptic drug treatment. New onset of psychiatric symptoms such as psychosis and depression can occur with anticonvulsant drugs. Paradoxical onset of depression in patients who have well-controlled seizures by antiepileptic drug treatment may be explained by forced normalization and the effect of antiepileptic medications on lowering the levels of folic acid [36].

Few cognitive symptoms can be associated with vigabatrin, topiramate, tiagabine, gabapentin, lamotrigine and levetiracetam [37].

Psychotic symptoms can be induced by ethosuximide, vigabatrin, phenytoin, topiramate carbamazepine, phenobarbital, primidone and benzodiazepines. Psychosis can occur in 6% of patients of patients receiving topiramate, 0.8% of patients with tiagabine treatment and 0.3% of patients taking lamotrigine. Between 2-4% of patients taking vigabatrin can develop schizophrenia-like psychosis [37-40].

Depression can be associated with the following antiepileptic drugs: acetazolamide, barbiturates, ethosuximide, carbamazepine, gabapentin, felbamate, levetiracetam, piracetam, phenytoin, tiagabine, topiramate, vigabatrin and zonisamide. There is limited evidence of self-harm and suicidal behavior linked to levetiracetam, tiagabine, topiramate and vigabatrin [41].

In 2008, the U.S. Food and Drug Administration (FDA) published a meta-analysis of data including all clinical trials involving antiepileptic drugs. Suicidal risk was found to be 0.43 per 1000 patients in active drug arms compared with a rate of 0.22 per 1000 in the placebo arm. However, risk of suicide due to antiepileptic drugs remains a controversial issue. Careful psychiatric assessment and screening for depressive symptoms and suicidal thoughts is advisable both before and during treatment with antiepileptic medications [42-44].

#### Table 2: Adverse Psychiatric Events Associated with Antiepileptic Drugs

| Depression             | Acetazolamide, barbiturates, ethosuximide, carbamazepine, gabapentin, felbamate, levetiracetam, piracetam, phenytoin, tiagabine, topiramate, vigabatrin and zonisamide. | Etiology may be related to forced normalization of decreasing folic acid levels. |
|------------------------|-------------------------------------------------------------------------------------------------|---------------------------------------------------------------------------------|
| Suicidal behavior      | Levetiracetam, tiagabine, topiramate and vigabatrin.                                           | Increased risk with rapid dose titration (e.g., topiramate)                     |
| Cognitive symptoms     | Vigabatrin, topiramate, tiagabine, gabapentin, lamotrigine and levetiracetam.                  | Limited evidence                                                               |
| Psychotic symptoms     | Ethosuximide, vigabatrin, phenytoin, topiramate, carbamazepine, phenobarbital, primidone and benzodiazepines. | Both epilepsy and antiepileptic drugs may be implicated                         |

Use of Psychotropic Medications for Patients with Epilepsy

Some of the psychotropic medications are associated with a dose-related risk of reducing seizure threshold, particularly if they are inducing severe hyponatraemia. Therefore, treatment should be started with lower doses and increased gradually [41].

Increased incidence of seizures is associated with antipsychotics, particularly clozapine and olanzapine, and with the antidepressant clomipramine. Other psychotropic medications which have high incidence of seizures include alprazolam, bupropion immediate release form, and quetiapine [45].

Some studies found no statistically significant difference in frequency of seizures between the period before using psychotropics and treatment period. This finding has been documented in a retrospective assessment of 57 consecutive patients with epilepsy who are also using different psychotropic medications indicated that seizure frequency decreased in 33%, was unchanged in 44%, and increased in 23% of patients during psychotropic drug therapy [46].

Good choices of antipsychotic medications for patients with epilepsy include: trifluperazine, haloperidol, and sulpiride which have low proconvulsive effects. Care is required when using risperidone, olanzapine (may affect EEG and myoclonic seizures reported), quetiapine (may reduce seizure threshold up to two-fold), and aripiprazole. Antipsychotics that are known as being epileptogenic should be avoided such as clozapine, chlorpromazine and zotepine. Good choices among antidepressants include moclobemide, and Selective Serotonin Re-uptake Inhibitors (SSRIs). Mirtazapine, reboxetine, venlafaxine and duloxetine should be used with caution. Bupropion and most of the tricyclic antidepressants (such as amitrptyline and clomiramine) are epileptogenic and should be avoided in patients with epilepsy [41].

Conclusion

Epilepsy and its treatment are associated with a wide range of psychiatric symptoms including mood disorders, psychosis and cognitive impairment. On the other hand, some of the psychotropic medications used for treatment of emerging psychiatric symptoms in patients with epilepsy may reduce seizure threshold. Evaluation of psychiatric symptoms in relation to seizures is essential for decision regarding the necessity of using psychotropic medications and the choice of psychotropics, if needed, depends on their effect on seizure control.
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