Sleep and epilepsy: unfortunate bedfellows

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ABSTRACT
The relationship between sleep and seizure disorders is a particularly vicious cycle. Nocturnal seizures can interrupt sleep while a number of factors, including antiepileptics and sleep disorders that cause sleep fragmentation, can worsen seizures. Understanding and managing seizures and related sleep disturbance is therefore an important and treatable intervention target that could potentially improve children’s sleep, but also their learning, mood, behaviour, seizures and parental quality of life.

A PARENT’S ACCOUNT
Our son’s seizures began out of the blue. There was no warning, I was downstairs when his dad called me. Our first thoughts were that he was having a stroke—his mouth dropped only on one side and only one side of his body was affected. We thought he was dying.

We called the ambulance and he stayed in hospital overnight. We waited a week and were then told they thought he had something called benign rolandic epilepsy. It is not a helpful name as we don’t think anything about it is benign.

In one 5-month period, he had 11 seizures from his sleep. Many have been so severe we have needed to call the ambulance to take him to the accident and emergency department which is very stressful. In fact, he has had a good period with no seizures now for a few months that we hope will continue.

There seems to be so much that people do not know. We needed to know about the future, but all the professionals could tell us is ‘he may never have another seizure, or he may have one every day—we don’t know’. ‘It’s just a matter of wait and see’.

For us that was unbearable. We were living on eggshells in anticipation.

In the end, we decided the only way we could cope as a family was to let him co-sleep with his dad. I now sleep in his bed. It’s not ideal, we know, but it’s the only way that we can keep an eye on him. We also know that if he sleeps badly his seizures may be worse and so at least this way we all get a good night’s sleep.

We’ve been told that the seizures don’t cause him damage unless they are very prolonged but that is hard to believe when you see him have one.

Our professionals have discussed about antiepileptic drugs. Again, there were no definite answers and they said “they were there if we wanted it”. We thought a lot about the pros and the cons and decided the risk of side effects were too great. He is happy and intelligent, doing well at school and we didn’t want to change any of that. Anyway, his seizures have stopped on their own for the time being; if he was on medicines we would have thought it was due to them and had to carry on giving them.

No one has ever discussed with us how this affects his sleep and our sleep. No one has asked about our sleeping arrangement. We have had the best support from facebook groups where at least we know other people are going through the same thing.

The impoRTAnCe of epilepsy
Epilepsy is an umbrella term for at least 0.5%–1% of children and often starts in childhood.5 Epilepsy is a common long-term condition affecting 0.5%–1% of children and often starts in childhood.5 Epilepsy is an umbrella term for at least
30 types of seizure disorder with different aetiology, manifestations and prognosis. Children with epilepsy experience seizures. However, the impact of epilepsy for children extends far beyond health, with higher and more persistent levels of emotional, behavioural, social and academic difficulties than children with other chronic health conditions or children in general.\textsuperscript{6,7}

UNDERSTANDING THE VICIOUS CYCLE

The interactions between epilepsy, antiepileptic drugs (AEDs), sleep and learning\textsuperscript{8} are summarised in figure 1.

Children’s sleep passes through different sleep stages conventionally divided into rapid eye movement (REM) and non-REM (NREM) sleep. NREM sleep coordinates electrical brain activity into important slow oscillation and faster bursts called spindles. Sleep research suggests that this slow wave synchrony and spindle activity contribute to the rejuvenation of synapses and overnight learning. Unfortunately, it also results in a vulnerable landscape, primed to allow easy and rapid generation of seizure activity. In this vicious cycle, anything that causes sleep disturbance or fragmentation can further trigger seizures. This includes sleep disorders, antiepileptics or seizures themselves. The association is a powerful one and epilepsy is associated with at least a 12 times higher rate of child-reported and parent-reported sleep disorders, even without nocturnal seizures.\textsuperscript{9,10}

Evidence shows that consolidation of memory during sleep (ie, new learning) is disrupted by seizures and by nocturnal interictal electroencephalograph (brainwave recording) discharges.\textsuperscript{11,12}

All these causes of sleep disturbance also leads to daytime sleepiness and worse behaviour, cognitive functioning and quality of life.\textsuperscript{10,11,14} The impact extends to the whole family; parents of these children wake seven times more often than controls\textsuperscript{15,16} and spend an average of 4 hours asleep, with adverse effects on marital satisfaction and maternal health.\textsuperscript{17}

Within this difficult vicious cycle, sleep disturbance, therefore, is a pivotal target of intervention that could potentially improve child sleep, but also learning, mood, behaviour, seizures, and parental quality of life.

THINGS THAT GO BUMP IN THE NIGHT: SLEEP-RELATED EPILEPSY SYNDROMES

One metaphor is to consider sleep in epilepsy as ‘deep sea diving’, the most dangerous times are during the initial dive and then on surfacing. A number of reviews describe a range of epilepsies that are strongly associated with sleep.\textsuperscript{18} The most striking example of this is the syndrome of electrical status epilepticus during slow sleep (ESES)/continuous spike-wave discharges during sleep (CSWS), an epileptic encephalopathy with behavioural and cognitive deficits, various seizures and continuous spike-wave activity occurring during NREM sleep.

Benign epilepsy with centrotemporal spikes or rolandic epilepsy (RE) is the most common childhood epilepsy. The centrotemporal spike-wave discharges in these epilepsies are activated in sleep, and the focal nature of the seizures (eg, a tingling feeling in the tongue, dysarthria and hemiparesis) means they can be confused with parasomnias if there is no history of secondary generalisation.

Genetic generalised epilepsies such as juvenile myoclonic epilepsy (JME) are sensitive to sleep disturbance and characteristically present with seizures after awakening.

In both structural and genetic frontal epilepsies, there are typically multiple episodes from sleep every night which may initially be difficult to identify as seizures. Parents may not witness their children’s focal onset of a frontal seizure and so miss the characteristic ‘fencing’ posture with head turning. Autosomal dominant nocturnal frontal lobe epilepsy (ADNFLE) is an example of a ‘genetic focal epilepsy’ and can be caused by mutations in a gene coding for acetylcholine receptor subunits. Seizures in ADNFLE may be typically frontal but also may take the form of nightmares, parasomnias and sleepwalking.

The differentiation of rarer nocturnal frontal lobe epilepsies from more common partial arousal parasomnias is important and helped by the history of more stereotyped and frequent (more than three) events in the epilepsies, but often videoelectroencephalography with polysomnography is necessary for their diagnosis. This is a complex area of differentiation, even after conducting appropriate investigations that is addressed in other reviews.\textsuperscript{19}

STRATEGIES TO IMPROVE SEIZURES

Antiepileptic drugs

Most industry trials of AEDs are regulatory and focused on percent seizure reduction and non-inferiority with benchmark drugs.\textsuperscript{20} However, epilepsy-specific quality of life is not determined solely by seizures but rather by factors such as a child’s
learning, mental health, sleep and social support, which are rarely measured.21–23

Patients identify cognitive side effects of AEDs as their most important health issue.24 The available evidence in adults suggests that all AEDs adversely affect learning, older AEDs more than newer.25–27 So far, there have been no randomised trials in children that either assess learning or compare the outcomes of treatment versus no treatment in terms relevant to parents and children.

In one specific example, parents of schoolchildren with nocturnal RE were ambivalent about medicating their children and expressed difficulty in weighing up the potential benefits and harms of using AEDs (http://castlesudy.org.uk/). A recent Cochrane review showed the rationale for antiepileptic treatment versus no treatment is not established in RE.28

Finally, it is well recognised that although improving seizure control might reduce sleep fragmentation, AEDs have differential effects on sleep architecture in epilepsy. This is not a well-studied area and a recent review highlighted differences between AEDs and shortcomings in the way they have been studied to date.29

This review commented that NREM (slow wave sleep (SWS)), for example, is increased by carbamazepine, pregabalin and gabapentin but is reduced by levetiracetam. REM sleep is enhanced by gabapentin but reduced by phenobarbital and levetiracetam. Unfortunately, many AEDs including phenobarbitone, sodium valproate and higher-dose levetiracetam may cause daytime sleepiness.

Several antiepileptic drugs (AEDs) are associated with weight gain such as gabapentin, pregabalin, valproic acid and vigabatrin and to some extent carbamazepine and this can independently increase the risk of conditions such as obstructive sleep apnoea (OSA).

**Ketogenic diet**

Ketogenic diet is used as a strategy to improve certain refractory epilepsies. It might improve nocturnal sleep; its effects on sleep independent of improved seizure frequency have not been analysed.30

**Vaginal nerve stimulation (VNS)**

VNS is used to treat certain refractory epilepsies. Some studies have shown that it can increase the percentage of SWS and reduce daytime sleepiness, although they did not analyse its effects on sleep independent of improved seizure frequency.31 32 It is important to remember that due probably to central and peripheral mechanisms VNS can increase the risk of OSA syndrome (OSAS).

**Surgery**

Improved total sleep time and reduced arousals are seen with epilepsy surgery, if the seizure frequency is also improved. It is obviously difficult to parse out the effects of improved seizure control from improved sleep.33

**STRATEGIES TO IMPROVE SLEEP DISORDERS**

**Sleep apnoea**

Sleep-related breathing disruptions in children with epilepsy are common and can range from primary snoring to OSA. Untreated OSA can lead to significant morbidity. The prevalence of OSAS in children with epilepsy is estimated at between 30% and 60%. Although increased weight is a possible risk factor, the risk of OSAS is greatest in those children with poorly controlled epilepsy requiring polypharmacy.34

It is thought that the presence of OSA itself may exacerbate epilepsy. The potential mechanisms are thought to be via changes in sleep architecture, sleep deprivation or hypoxia. In this way, another vicious cycle is seen whereby epilepsy may increase prevalence of OSA, and OSA may worsen epilepsy. Encouragingly, breaking this cycle seems possible and in adults treating the OSAS with positive airways pressure therapy has been reported to lead to a reduction in their seizure frequency.

**Disorders of sleep initiation and maintenance**

Parent-based sleep (PBS) interventions are effective in randomised controlled trials for typically developing (TD) younger children, attention-deficit hyperactivity disorder and autism populations35–37 and could easily be modified for children with epilepsy.

Many of the same sleep problems experienced by children with epilepsy are commonly seen in TD children and the same sleep interventions are effective in both TD and neurodevelopmental disorder populations.38 36 PBS interventions may therefore be effective in children with epilepsy. However, a ‘one-size-fits-all’ approach to behavioural management of child sleep problems fails to acknowledge additional relevant factors for a child with epilepsy (eg, nocturnal seizures, anxiety about seizures, anxiety about sudden unexpected death in epilepsy (SUDEP)) and their parents (eg, concerns about the appropriateness of using some behavioural techniques with a child who might have seizures) and therefore may disenfranchise parents. A recent sleep and seizure Delphi core outcome study has started to prioritise the sleep and seizure outcomes that matter most for parents and professionals.38 39 Acknowledging and addressing these needs and concerns needs to explicitly form part of any behavioural intervention delivery as is the case with the current CASTLE e-learning sleep module http://castlesudy.org.uk/castle-projects/sleep-intervention/.

Integrating SUDEP evidence-based advice into any sleep programme is challenging and the content needs to be developed with parents and professional groups in tandem. Generalised tonic–clonic seizures are clear risk factors for SUDEP, but nocturnal seizures may also increase the risk.30 These findings, in conjunction with the observation that postictal respiratory depression is a major mechanism in SUDEP suggest that witnessed nocturnal seizures and postictal respiratory depression can cause SUDEP.

Epidemiological studies showed that the presence of another individual at least 10 years of age and of normal intelligence in the bedroom was associated with a decreased SUDEP risk.40 These results might imply that a bedroom observer could detect seizures, check on the child and provide sufficient stimulation to prevent respiratory arrest, although this association does not prove that these interventions directly treat the mechanism that causes SUDEP. There are considerable concerns about the burden of such monitoring on caregivers and the degree of intrusiveness for the young person. Such pressure on parents to co-sleep can trigger inappropriate sleep associations that are often the beginning of more pervasive sleep difficulties. An intermediate step can be to use a listening device (baby monitor with or without video signal) to allow remote monitoring. The evidence for this is, however, weak.

Behavioural sleep modifications are arguably more difficult in adolescence, an age where uncontrolled seizures have implications for driving and employment deprivation. Sleep fragmentation can significantly reduce seizure threshold, especially, for generalised tonic–clonic seizures, to such an extent that it has been suggested that ‘treatment of JME is a lifestyle choice’. The social curfews experienced by these young people have been
described as ‘Cinderella syndrome’ with a constant imperative to be ‘home by midnight’.

THE FUTURE

There are many emerging studies across different domains with potential to help parents and professionals better understand and manage sleep and seizures in children.

Delphi studies involving all stakeholders are starting to allow consensus around the core sleep and seizure outcomes that matter most to young people, their parents and professionals. These need to be routinely incorporated in all future sleep and epilepsy intervention studies. For many such studies, where there is equipoise about treatment options, a ‘no-treatment’ arm is essential.

While high-tech body worn devices might soon allow accurate monitoring and management of sleep and seizure disorder, at present they lack the necessary precision, and their effectiveness needs proper testing in varied patient groups to avoid the consequences of either false reassurance or missing true events. In the interim, there are ‘low-tech’ solutions involving evidence-based parent-led sleep interventions that are underused and have the potential to improve sleep of affected children and their carers and enable a virtuous cycle of sleep and seizures.

Contributors PG conceived the content of the review. PG and FMG contributed to writing the review. EM wrote the parent account that begins the whole review and read and checked the rest of the review.

Funding Professor Gringas and Dr Gibbon are investigators on the Changing Agendas in Sleep and Treatment in Childhood Epilepsy (CASTLE) Project— independent research funded by the National Institute for Health Research (NIHR) under its Programme Grants for Applied Research Programme (RP-PG-0615-20007). The views expressed are those of the authors and not necessarily those of the NHS, the NIHR or the Department of Health.

Competing interests None declared.

Patient consent Obtained.

Provenance and peer review Commissioned; externally peer reviewed.

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