Despite the availability of more than 20 different antiseizure drugs and the provision of appropriate medical therapy, 30% of people with epilepsy continue to have seizures.\(^1,2\) The approval of many new antiseizure drugs during the past two decades, including several with novel mechanisms of action, has not substantially reduced the proportion of patients with medically refractory disease.\(^1\) The safety and side-effect profile of antiseizure drugs has improved, but side effects related to the central nervous system are common and affect quality of life.\(^3\) Patients need new treatments that control seizures and have fewer side effects. This treatment gap has led patients and families to seek alternative treatments. Cannabis-based treatment for epilepsy has recently received prominent attention in the lay press\(^4\) and in social media, with reports of dramatic improvements in seizure control in children with severe epilepsy. In response, many states have legalized cannabis for the treatment of epilepsy (and other medical conditions) in children and adults (for a list of medical marijuana laws according to state, see www.ncsl.org/research/health/state-medical-marijuana-laws.aspx).

Cannabis has been used medicinally for millennia and was used in the treatment of epilepsy as early as 1800 B.C.E. in Sumeria.\(^5\) Victorian-era neurologists used Indian hemp to treat epilepsy and reported dramatic success.\(^5,6\) The use of cannabis therapy for the treatment of epilepsy diminished with the introduction of phenobarbital (1912) and phenytoin (1937) and the passage of the Marijuana Tax Act (1937). The discovery of an endogenous cannabinoid-signaling system in the 1990s\(^7\) rekindled interest in therapies derived from constituents of cannabis for nervous system disorders such as epilepsy (see ClinicalTrials.gov numbers, NCT02091375, NCT02224690, NCT02324673, NCT02318537, and NCT02318563). This review addresses the current preclinical and clinical data that suggest that compounds found in cannabis have efficacy against seizures. The pharmacokinetic properties of cannabinoids and related safety and regulatory issues that may affect clinical use are also discussed, as are the distinct challenges of conducting rigorous clinical trials of these compounds.

More than 545 distinct compounds have been isolated from cannabis species; the most abundant are the cannabinoids, a family of molecules that have a 21-carbon terpenophenolic skeleton and includes numerous metabolites.\(^8\) The best studied of these cannabinoids (termed “phytocannabinoids” if derived from the plant) are \(\Delta^9\)-tetrahydrocannabinol (\(\Delta^9\)-THC) and cannabidiol and their metabolites. (See Fig. 1 for the structure of \(\Delta^9\)-THC, cannabidiol, and one other cannabinoid, cannabidiicarvin, as well as their targets in the central nervous system, and their actions.) Most of the psychoactive effects of cannabis are mediated by \(\Delta^9\)-THC. Many of the noncannabinoid molecules in cannabis plants may have biologic activity. This review focuses on cannabinoids, since other cannabis-derived compounds have been less well studied.

Dan L. Longo, M.D., Editor

Cannabinoids in the Treatment of Epilepsy

Daniel Friedman, M.D., and Orrin Devinsky, M.D.
The major cannabinoid receptor in the central nervous system is cannabinoid receptor 1 (CB₁R), a presynaptic, G-protein–coupled receptor that activates voltage-gated calcium channels and enhances potassium-channel conduction in presynaptic terminals. The cloning of CB₁R, the confirmation that Δ⁹-THC binds CB₁R, and the discovery of two endogenous ligands — 2-arachidonoylglycerol (2-AG) and anandamide — that bind CB₁R² has stimulated investigations intended to elucidate the role of the endocannabinoids both in normal brain function and in disease states. CB₁R is activated by the activity-dependent synthesis of 2-AG and is involved in the retrograde control of synaptic transmission. Anandamide can also affect excitability in neuronal networks by activating the transient receptor potential cation channel, subfamily V, member 1.¹¹ As modulators of neuronal excitability, endogenous cannabinoids are well poised to affect the initiation, propagation, and spread of seizures.

Preliminary studies have identified defects in
the endocannabinoid system in persons with epilepsy. In one study, patients with newly diagnosed temporal-lobe epilepsy had significantly lower levels of anandamide in cerebrospinal fluid than healthy controls. In another study, tissue resected from patients undergoing surgery for epilepsy had lower levels of CB1R messenger RNA, particularly in the glutamatergic terminals in the dentate gyrus, than did specimens obtained post mortem from persons without epilepsy. There was also reduced expression of diacyl glycerol lipase α (DAGL-α), the enzyme responsible for the “on-demand” synthesis of 2-AG in postsynaptic neurons. These studies support the suggestion that the endocannabinoid system plays a role in the inhibition of seizures in humans with epilepsy.

The endocannabinoid system is strongly activated by seizures, and the upregulation of CB1R activity has antiseizure effects. In mice, hippocampal anandamide levels rise after seizures induced by the intraperitoneal injection of kainic acid. In cultures of neurons from the hippocampus, CB1R antagonists induce prolonged, seizure-like discharges, whereas CB1R agonists eliminate these discharges. Conditional knockout mice that lack pyramid-cell CB1R in their forebrain have more severe and prolonged seizures than wild-type mice in response to kainic acid; in contrast, viral-vector–mediated overexpression of CB1R in hippocampal pyramidal cells is protective. Reducing the metabolic degradation of endocannabinoids ameliorates experimentally induced seizures.

**Preclinical Evidence of Antiseizure Effects**

The activation of CB1R receptors with the use of Δ9-THC or synthetic agonists in experimentally induced seizures has been studied in various animal models (see Hill et al., for a summary). In most studies, CB1R agonists reduced seizures, but in others no effect was observed, and in four studies CB1R activation was associated with convulsive effects at some doses. CB1R antagonists reduced the threshold for seizure in some studies in animals, a finding that further supports the possibility that CB1R activation has anticonvulsant effects.

Other plant cannabinoids have also been studied in animal models of seizures and epilepsy. Cannabidiol, the most abundant nonpsychoactive cannabinoid, has shown antiseizure effects in several in vivo and in vitro models of epilepsy. Unlike Δ9-THC, cannabidiol does not exert its main neural effects through the activation of CB1R. At high levels, cannabidiol may function as an indirect CB1R antagonist. Cannabidiol alters neuronal excitability by other means. These include binding to members of the TRP family of cation channels at low levels, which antagonizes the G-protein–coupled receptor 55, leading to decreased presynaptic release of glutamate; activating 5-hydroxytryptophan 1A receptors; and inhibiting adenosine reuptake through multiple mechanisms. In addition, cannabidiol may exert antioxidant and anti-inflammatory effects. Cannabidiol's lack of psychoactive effects and the preclinical evidence of antiseizure effects has generated interest in its potential as an antiseizure drug in humans.

Cannabidivarin, the propyl variant of cannabidiol, has also shown antiseizure effects in both in vitro and in vivo models. Like cannabidiol, cannabidivarin has antiseizure effects that are independent of the endocannabinoid system and may function by means of its influence on TRP channels or by lowering 2-AG synthesis through the inhibition of DAGL-α. Little is known about the antiseizure effects of other phytocannabinoids. Cannabinoil and Δ1-THCV, the propyl variant of Δ9-THC, have been shown to have anticonvulsant effects in a few small studies.

**Evidence of Antiseizure Effects in Humans**

Despite the preclinical data and anecdotal reports on the efficacy of cannabis in the treatment of epilepsy that include reports from epileptologists, a recent Cochrane review concluded that “no reliable conclusions can be drawn at present regarding the efficacy of cannabinoids as a treatment for epilepsy” owing to the lack of adequate data from randomized, controlled trials of Δ9-THC, cannabidiol, or any other cannabinoid (Table 1). This assessment was confirmed in a recent systematic review by the American Academy of Neurology.

Limited epidemiologic evidence supports the view that cannabinoids have antiseizure properties in humans. In a case–control study of illicit drug use and new-onset seizures in Harlem,
New York, men who used cannabis within 90 days before hospital admission were at a significantly lower risk for presenting with new-onset seizures than men who did not use cannabis (odds ratio, 0.36; 95% confidence interval, 0.18 to 0.74).\textsuperscript{48}

Several patient and caregiver surveys have examined the effects of cannabis in epilepsy. In one survey, 28 of 136 patients in an epilepsy center that provided tertiary care reported cannabis use. Most of these patients associated use with a reduction in seizure frequency and severity.\textsuperscript{45} A 2013 survey of caregivers of 19 children with severe epilepsy who were receiving cannabidiol-enriched cannabis extracts indicated that 2 of the children had become seizure-free and 8 others had a reduction in the frequency of seizures of 80% after taking the extract.\textsuperscript{42} In a 2015 survey of 75 parents whose children were treated with oral cannabis extracts in Colorado, the parents reported that one third of the children had a reduction in seizures of more than 50%.\textsuperscript{34} However, electroencephalograms were obtained for 8 of these children before and after the administration of cannabis, and none showed improvement in background activity.

Case reports support the antiseizure effects of cannabis in patients with epilepsy\textsuperscript{6,32-34,49} and show exacerbation of seizures after abrupt discontinuation.\textsuperscript{50} However, in a survey conducted in Germany among adults with epilepsy who used cannabis, the substance had no apparent effect on seizure control,\textsuperscript{46} and some case reports have shown an exacerbation of seizures among patients who used cannabis\textsuperscript{51} or a synthetic cannabinoid.\textsuperscript{52}

Few prospective therapeutic trials have been performed that involve the isolated use of cannabinoids to treat epilepsy. A study conducted in 1949 indicated that two of five institutionalized children with refractory epilepsy achieved seizure control after receiving treatment with a Δ9-THC analogue.\textsuperscript{36} To our knowledge, only four placebo-controlled studies of the use of cannabinoids for the treatment of epilepsy have been performed (reviewed in Gloss and Vickrey\textsuperscript{35}). All the studies were considerably underpowered and had methodologic problems, including the lack of blinding. Two studies showed a reduction in the number of seizures in patients treated with cannabidiol, whereas the other two studies showed no effect.

Since 2013, a consortium of 10 epilepsy centers has been collecting prospective data on children and young adults with severe epilepsy who are receiving Epidiolex, a purified cannabis extract containing 99% cannabidiol and less than 0.10% Δ9-THC (GW Pharmaceuticals), through an expanded-access program authorized by the Food and Drug Administration (FDA). A preliminary report from this open-label study, initiated by investigators to assess the safety and dosing of cannabidiol, noted that among 137 patients who had received at least 12 weeks of treatment, the median reduction in the number of seizures was 54%.\textsuperscript{46} Randomized clinical trials of Epidiolex are now being conducted for the treatment of two forms of severe, childhood-onset epilepsy: Dravet’s syndrome (a severe myoclonic epilepsy of infancy) (NCT02091375) and the Lennox–Gastaut syndrome (a childhood-onset, treatment-resistant epilepsy characterized by multiple types of seizures and developmental delay) (NCT02224690). Although some of the anecdotal evidence described above suggests that cannabidiol-rich treatments may ameliorate seizures in patients with these disorders, no evidence suggests that the antiseizure effects of cannabidiol are limited to the treatment of these conditions. The clinical development of synthetic forms of cannabidiol is also in progress (NCT02318563). Table 1 summarizes the current clinical evidence for the use of cannabidiol-containing compounds in the treatment of epilepsy.

### Safety in Humans

Much of the available data regarding the safety and side-effect profile of cannabinoids, especially with long-term use, come from studies examining the effects of recreational use.\textsuperscript{53,54} The short-term side effects of cannabis use may include impairment of memory, judgment, and motor performance. High levels of Δ9-THC are associated with psychosis and an increased risk of motor-vehicle accidents. With long-term use there is a risk of addiction, which occurs in approximately 9% of long-term users. Other effects of long-term use include cognitive impairment, decreased motivation, and an increased risk of psychotic disorders.

Cannabis-based treatment with Δ9-THC may have irreversible effects on brain development.
### Table 1. Clinical Trials, Case Series, and Case Reports on Cannabinoids in the Treatment of Epilepsy.

<table>
<thead>
<tr>
<th>Compound and Study Type</th>
<th>Dose*</th>
<th>No. of Participants</th>
<th>Results</th>
<th>Reference</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Isolated oral cannabinoids</strong></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td><strong>THC isomers</strong></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Case series of institutionalized children with intellectual disability and epilepsy treated for 3–7 wk</td>
<td></td>
<td>5</td>
<td>One patient was seizure-free and one patient nearly seizure-free</td>
<td>Davis and Ramsey36</td>
</tr>
<tr>
<td><strong>Cannabidiol</strong></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Prospective, placebo-controlled, 3-mo trial involving adults with treatment-resistant epilepsy</td>
<td>200 mg/day</td>
<td>Treatment: 4 Placebo: 5</td>
<td>Two patients in the cannabidiol group were seizure-free and one showed partial improvement No report of baseline seizure measurement</td>
<td>Mechoulam and Carlini37</td>
</tr>
<tr>
<td>Prospective, placebo-controlled trial involving teenagers and adults with treatment-resistant convulsive seizures (at least 1 per wk) in which participants had 8–18 wk of exposure</td>
<td>200–300 mg/day</td>
<td>Treatment: 8 Placebo: 7</td>
<td>Four patients in cannabidiol group and 1 in placebo group were seizure-free; somnolence was a reported side effect Clinicians were not masked to group assignment; one patient switched groups for unknown reasons</td>
<td>Cunha et al.38</td>
</tr>
<tr>
<td>Prospective, placebo-controlled, 3-wk trial involving institutionalized adults with intellectual disability and epilepsy</td>
<td>200–300 mg/day</td>
<td>Treatment: 6 Placebo: 6</td>
<td>No significant difference between groups Somnolence was a reported side effect</td>
<td>Ames and Cridland39</td>
</tr>
<tr>
<td>Prospective randomized, double-blind placebo-controlled, 6-mo crossover study involving adults with treatment-resistant epilepsy</td>
<td>300 mg/day</td>
<td>12</td>
<td>No significant difference between cannabidiol and placebo Somnolence was a reported side effect</td>
<td>Trembly and Sherman40</td>
</tr>
<tr>
<td><strong>Purified oral cannabidiol extract</strong></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Prospective, open-label, 12-wk trial involving children and young adults with severe, childhood-onset epilepsy</td>
<td></td>
<td>137</td>
<td>Median reduction in weekly rate of convulsive seizures of 54% Somnolence and diarrhea were the most common side effects</td>
<td>Devinsky et al.41</td>
</tr>
<tr>
<td><strong>Oral cannabis extracts</strong></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Cannabis indica extract</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Case report of a 40-yr-old man with focal epilepsy who was resistant to bromides</td>
<td>32 mg/day</td>
<td>1</td>
<td>Seizure-free for 6 mo followed by recurrence when cannabis extract was discontinued; seizure control resumed with resumption of cannabis several months later</td>
<td>Gowers6</td>
</tr>
<tr>
<td><strong>Cannabidiol–Δ9-THC–containing extracts of varying composition</strong></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Survey among participants in a Facebook group for parents of children with severe epilepsies</td>
<td>Cannabidiol: Up to 28 mg/kg body weight/day Δ9-THC: Up to 0.8 mg/kg body weight/day</td>
<td>19</td>
<td>Improvement with cannabidiol–Δ9-THC reported by 16 patients (84%); 2 patients (11%) became seizure-free</td>
<td>Porter and Jacobson42</td>
</tr>
<tr>
<td>Compound and Study Type</td>
<td>Dose</td>
<td>No. of Participants</td>
<td>Results</td>
<td>Reference</td>
</tr>
<tr>
<td>-------------------------</td>
<td>------</td>
<td>---------------------</td>
<td>---------</td>
<td>-----------</td>
</tr>
<tr>
<td>Oral cannabis extract, with high ratio of cannabidiol to Δ9-THC</td>
<td>1053</td>
<td>Reduction of &gt;90% in frequency of generalized tonic–clonic seizures, which allowed for reduction of other drugs taken for epilepsy</td>
<td>Maa and Figi</td>
<td></td>
</tr>
<tr>
<td>Case report of 5-yr-old girl with Dravet’s syndrome</td>
<td>1</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Oral cannabis extracts</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Retrospective case series of children with refractory epilepsy at a center in Colorado</td>
<td>75</td>
<td>Reduction of &gt;50% in frequency of seizures in 25 patients (33%)</td>
<td>Press et al.</td>
<td></td>
</tr>
<tr>
<td>Smoked cannabis</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Cannabis</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Case report of 20-yr-old man with refractory tonic–clonic seizures whose seizures were well-controlled</td>
<td>1</td>
<td>Seizures were exacerbated after smoking marijuana</td>
<td>Keeler and Reifler</td>
<td></td>
</tr>
<tr>
<td>Case report of 24-yr-old man with refractory generalized epilepsy</td>
<td>1</td>
<td>Nearly seizure-free after daily cannabis use</td>
<td>Consroe et al.</td>
<td></td>
</tr>
<tr>
<td>Case report of 29-yr-old man with refractory focal epilepsy</td>
<td>1</td>
<td>Suppression of complex partial seizures with cannabis use and exacerbation of seizures on withdrawal</td>
<td>Ellison et al.</td>
<td></td>
</tr>
<tr>
<td>Case report of 45-yr-old man with cerebral palsy and refractory focal epilepsy</td>
<td>1</td>
<td>Reduction of &gt;90% in nocturnal seizures and tonic–clonic seizures</td>
<td>Mortati et al.</td>
<td></td>
</tr>
<tr>
<td>Survey of active users at center for patients with tertiary epilepsy</td>
<td>28</td>
<td>Reduction in severity of seizures reported by 19 patients (68%); 15 patients (54%) reported reduction in frequency of seizures</td>
<td>Gross et al.</td>
<td></td>
</tr>
<tr>
<td>Survey of cannabis users seen at center for patients with tertiary epilepsy</td>
<td>Active users: 13 Former users: 297</td>
<td>Reduction in frequency of seizures reported by 2 active users (15%); increase in frequency and severity of seizures reported by 7 former users (0.2%)</td>
<td>Hamerle et al.</td>
<td></td>
</tr>
</tbody>
</table>

* Data on dosage has been provided when available.
The New England Journal of Medicine

The endocannabinoid system undergoes development in childhood and adolescence; long-term exposure to endocannabinoids, especially Δ9-THC, may lead to cognitive and behavioral changes. Imaging studies of the brain reveal altered structure and function in long-term adult users, including impaired connectivity of the prefrontal cortices and precuneus and decreased volume.54 Long-term use of cannabis in childhood may be associated with lower-than-expected IQ scores (although socioeconomic status may be a confounding factor; see Rugeberg57). It is unknown whether adverse effects on the brain are mediated solely by psychoactive cannabinoids, such as Δ9-THC, or whether long-term exposure to cannabidiol and cannabidivarin also have deleterious effects. Until more data become available, the neurodevelopmental risks of cannabinoid-based therapies should be weighed against the potential benefits for seizure control, since seizures also affect brain development. Notably, scientific data on the potential long-term developmental effects of FDA-approved antiseizure drugs are also limited.

Many antiseizure drugs are associated with teratogenicity and neurodevelopmental impairments in children who are exposed in utero. Little is known about the effects of fetal exposure to cannabinoids. Studies of children born to parents who are recreational cannabis users have not shown an increased risk of congenital abnormalities, but difficulties with attention, impulse control, and executive function have been reported.58 However, potential confounding factors, such as socioeconomic status and coexisting maternal psychiatric illness, limit the extent to which these findings can be interpreted.

Data regarding the outcomes of short-term and long-term exposure to cannabinoids in recreational users are often confounded by the factors that drive a person to use cannabis. More valid data regarding the safety of short-term use comes from randomized clinical trials of cannabinoid-containing medications, including purified cannabis extracts (Cannador, Society for Clinical Research, Germany; 2:1 ratio of Δ9-THC and cannabidiol),59 nabiximols (Sativex, GW Pharmaceuticals, 1:1 ratio of Δ9-THC and cannabidiol),60 and the synthetic Δ9-THC analogues dronabinol (Marinol, Unimed Pharmaceuticals),61 and nabiximols (Cesamet, Valeant Pharmaceuticals).62 These trials involved the systematic collection of data on safety. In a pooled analysis that included 1619 patients in short-term placebo-controlled studies who received cannabinoids for the treatment of pain and tremor and for spasticity related to multiple sclerosis, 6.9% withdrew because of adverse effects, as compared with 2.2% who withdrew in the placebo groups.47 Adverse effects that occurred in more than one study included nausea, weakness, mood changes, psychosis, hallucinations, suicidal ideation, dizziness or light-headedness, fatigue, and feeling of intoxication. No deaths from overdose were reported in association with cannabinoid-containing medications. In small studies of cannabidiol use in healthy volunteers and in patients with multiple disease conditions, serious side effects have been associated with either long-term or short-term administration of doses of up to 1500 mg daily.63 In the preliminary results of an open-label study of the use of cannabidiol oral solution for severe, refractory, childhood-onset epilepsy, the most common side effects were somnolence (occurring in 21% of the participants), diarrhea (17%), fatigue (17%), and decreased appetite (16%). Increased frequency or severity of seizures, weight loss, diarrhea, pneumonia, and abnormal results on tests of liver function were less common, occurring in 1 to 7% of patients.41

Long-term recreational use of cannabis is associated with a risk of dependence.54 Little is known regarding the potential for the abuse of cannabinoid-based treatments when they are administered in a clinical setting. A single-dose, double-blind, crossover study involving 23 recreational cannabis users showed higher scores on scales of drug preference for dronabinol and high-dose nabiximols but not for low-dose nabiximols,64 which suggests that there may be a potential for abuse associated with cannabinoid-based therapies, at least when the compounds used contain Δ9-THC or its analogues. Few data are available on the effects of other cannabinoids, although the relative absence of psychoactive effects reported for cannabidiol and cannabidivarin suggests that the potential for abuse of these compounds is low.

Some safety concerns have been raised with regard to the pharmacokinetic interactions of cannabinoids in patients with epilepsy who are long-term users. Cannabinoids can inhibit cytochrome P-450 (CYP) enzymes. Both Δ9-THC and cannabidiol inhibit the CYP2C family of isozymes.
at low micromolar concentrations and CYP3A4 at higher concentrations. These enzymes help to metabolize many antiseizure drugs, and inhibition can potentiate drug toxicity and efficacy. Both cannabidiol and Δ⁹-THC are metabolized through the P-450 system, especially through CYP2C9 and CYP3A4. These isozymes are induced by commonly prescribed antiseizure drugs, such as carbamazepine, topiramate, and phenytoin, and are inhibited by others, such as valproate, and the potential for drug–drug interactions between antiseizure drugs and cannabinoids is bidirectional. Preliminary evidence suggests that cannabidiol can raise the serum levels of the N-desmethyl metabolite of clobazam, which can have antiseizure and sedative effects.

As is the case with any medication, accidental ingestion of cannabis by children is a concern, and with cannabis preparations, the concern is particularly great because these preparations are not packaged in childproof containers and because some are made in formulations that may be appealing to children (gummies, brownies, or other edible forms). Finally, there are safety concerns related to the preparation of cannabis for medicinal use. Although many states have approved the use of “medical” marijuana, patients or caregivers often process the plant for therapeutically use. Reliance on recipes pulled from the Internet that use butane or high-proof alcohols to extract cannabinoids from plant material has resulted in more than 30 home explosions in a 5-month period in Colorado.

**PERCEIVED THERAPEUTIC BENEFIT**

Another obstacle to scientific inquiry into cannabinoids for the treatment of epilepsy is the perception among many patients and caregivers that sufficient evidence of their safety and efficacy already exists. The gap between patient beliefs and available scientific evidence highlights a set of factors that confound cannabinoid research and therapy, including the naturalistic fallacy (the belief that nature’s products are safe), the conversion of anecdotes and strong beliefs into facts, failure to appreciate the difference between research and treatment, and a desire to control one’s care, including access to therapies of perceived benefit. In one study of children with epilepsy in Colorado, the rate of response to therapy reported by parents who had moved their family to the state to receive cannabinoid therapy was more than twice as high as that reported by parents who were already residing in the state (47% vs. 22%). This finding...
suggests that the stronger the belief that the drug will be beneficial and the greater the sacrifice involved to obtain the drug, the greater the reported response. In the future, randomized, controlled studies of cannabinoids will have to contend with large placebo effects that may actually prevent researchers from demonstrating the efficacy of cannabinoids over placebo.

The currently planned randomized clinical trials of cannabidiol will target primarily children with severe epilepsy. Placebo response rates are high among children and adolescents with a wide variety of conditions, including pain-related disorders (e.g., migraines and gastrointestinal disorders), medical disorders (e.g., asthma), and psychiatric disorders (e.g., anxiety, major depression, obsessive-compulsive disorder, and attention-deficit disorder). The issue of high response rates to placebos in studies of children is especially relevant to epilepsy and emphasizes the importance of placebo-controlled trials. A meta-analysis showed that among patients with treatment-resistant focal epilepsy, children had more improvement with placebo than did adults (19.9% vs. 9.9%), although there was no significant difference in the response to active treatment. Children with intellectual disability and severe epilepsy are especially prone to elevated response rates to placebo. For instance, in a clinical trial of clobazam in children with the Lennox–Gastaut syndrome (mean age, 12.4 years), the response rate (defined as a decline of more than 50% in the number of drop seizures [brief seizures associated with a sudden increase or decrease in muscle tone, often causing a fall if the person is standing]) in the placebo group was 31.6%, a rate similar to that in the group receiving clobazam. However, the average weekly frequency of seizures was significantly lower in the clobazam group.

CONCLUSIONS AND FUTURE DIRECTIONS

Preclinical and preliminary data from studies in humans suggest that cannabidiol and Δ⁹-THC may be effective in the treatment of some patients with epilepsy. However, current data from studies in humans are extremely limited, and no conclusions can be drawn. Relaxation of the regulatory status of cannabis-derived drugs, especially those containing a high proportion of nonpsychoactive cannabinoids, for which the potential for abuse is low, could help to accelerate scientific study. Despite the power of anecdote and the approval of medical cannabis by many state legislatures, only double-blind, placebo-controlled, randomized clinical trials in which consistent preparations of one or more cannabinoids are used can provide reliable information on safety and efficacy. The use of medical cannabis for the treatment of epilepsy could go the way of vitamin and nutritional supplements, for which the science never caught up to the hype and was drowned out by unverified claims, sensational testimonials, and clever marketing. If randomized clinical trials show that specific cannabinoids are unsafe or ineffective, those preparations should not be available. If studies show that specific cannabinoids are safe and effective, those preparations should be approved and made readily available.

Dr. Devinsky reports receiving grant support from GW Pharmaceuticals and Novartis and serving on the scientific advisory board of MiaMed; and Dr. Friedman, receiving fees for serving on an advisory board for Marinus Pharmaceuticals and consulting fees from Eisai, Marinus Pharmaceuticals, SK Biopharmaceuticals, Upsher-Smith Laboratories, and Pfizer, all of which were paid to the Epilepsy Study Consortium. No other potential conflict of interest relevant to this article was reported.

Disclosure forms provided by the authors are available with the full text of this article at NEJM.org.

REFERENCES

10. Pertwee RG, Cascio MG. Known pharmacological actions of delta-9-tetrahydrocannabinol and of four other chem-
Cannabinooids in the Treatment of Epilepsy

23. Pertwee RG. The diverse CB1 and CB2 receptor pharmacology of three plant can-
nabinoids: delta9-tetrahydrocannabinol, cannabidiol and delta9-tetrahydrocannabin-
cannabinoid-enriched Cannabis extracts on TRP channels and endocannabinoid
25. Sylantyev S, Jensen TP, Ross RA, Rusa-
kov DA. Cannabinoid- and lysophosphatidyl-
diinositol-sensitive receptor GPR55 boosts
neurotransmitter release at central syn-
apses. Proc Natl Acad Sci U S A 2013;110:
5193-8.
26. Campos AC, Ferreira FR, Guimarães FS. Cannabidiol blocks long-lasting be-
havioral consequences of predator threat stress: possible involvement of SHT1A
receptors. J Psychiatr Res 2012;46:1501-
10.
27. Carrier EJ, Auchampach JA, Hillard CJ. Inhibition of an equilibrative nucleo-
side transporter by cannabidiol: a mecha-
nism of cannabinoid immunosuppres-
ion. Proc Natl Acad Sci U S A 2006;103:
7895-900.
28. Hampson AJ, Grimaldi M, Axelrod J, Wink D. Cannabidiol and (−)Delta9-tetra-
hydrocannabinol are neuroprotective anti-
oxidants. Proc Natl Acad Sci U S A 1998;
95:8268-73.
29. Hill TD, Cascio MG, Romano B, et al. Cannabidiervarin-rich cannabis extracts are
anticonvulsant in mouse and rat via a CB1
receptor-independent mechanism. Br J
30. Iannotti EA, Hill CL, Leo A, et al. Non-
psychotropic plant cannabinoids, canna-
bidiavin (CBDV) and cannabidiol (CBD),
activate and desensitize transient receptor
potential vanilloid 1 (TRPV1) channels in
vitro: potential for the treatment of ne-
uronal hyperexcitability. ACS Chem Neuro-
sci 2014;5:1131-41.
31. Ellison JM, Gelwan E, Oglettre I. Complex partial seizure symptoms affect-
32. Mortati K, Dworetzky B, Devinsky O. Mar-
ijana: an effective antiepileptic treat-
ment in partial epilepsy? A case report
and review of the literature. Rev Neurol
33. Maa E, Figi P. The case for medical mari-
34. Gross DW, Hamm J, Ashworth NL, Quigley D. Marijuana use and epilepsy:
prevalence in patients of a tertiary care
35. Hamerle M, Ghaeni L, Kowski A, Weissinger F, Holtkamp M. Cannabis and
36. Davis JP, Ramsey HH. Anti-epileptic
37. Mechoulam R, Carlini EA. Toward
drugs derived from cannabis. Naturwissen-
38. Cunha JM, Carlini EA, Pereira AE, et al. Chronic administration of cannabidiol
39. Ames FR, Girdland S. Anticonvulsant
effect of cannabidiol. S Afr Med J 1986;69:
14.
40. Trembly B, Sherman M. Double-blind
clinical study of cannabidiol as a second-
ary anticonvulsant. Presented at the Mari-
ijuana ’90 International Conference on
Cannabis and Cannabinoids, Kolymvari,
Crete, July 8–11, 1990. abstract.
41. Devinsky O, Sullivan J, Friedman D,
et al. Epidiolax (cannabidiol) in treatment
resistant epilepsy. Presented at the an-
ual meeting of the American Academy of
Neurology, Washington, DC, April 18–25,
2015. abstract.
42. Porter BE, Jacobson C. Report of a
parent survey of cannabidiol-enriched can-
nabis use in pediatric treatment-resistant
43. Keeler MH, Reifler CB. Grand mal
convulsions subsequent to marijuana use:
44. Consroe PF, Wood GC, Buchsbaum H.
45. Gross DW, Hamm J, Ashworth NL, Qui-
gley D. Marijuana use and epilepsy:
prevalence in patients of a tertiary care
46. Hamerle M, Ghaeni L, Kowski A, Weissinger F, Holtkamp M. Cannabis and
47. Koppel BS, Brust JC, Fife T, et al. Sys-
tematic review: efficacy and safety of
medical marijuana in selected neurologic
disorders: report of the Guideline Devel-
opment Subcommittee of the American
Academy of Neurology. Neurology 2014;
82:1536-63.
48. Brust JC, Ng SK, Hauser AW, Susser M. Mar-
ijana use and the risk of new onset
seizures. Trans Am Clin Climatol Assoc
49. Gordon E, Devinsky O. Alcohol and
marijuana: effects on epilepsy and use by
patients with epilepsy. Epilepsia 2001;42:
1266-72.
50. Hegde M, Santos-Sanchez C, Hess CP,
Kabir AA, Garcia PA. Seizure exacerbation
in two patients with focal epilepsy following
51. Feeney DM. Marijuana use among
52. Tofghi B, Lee JD. Internet highs –
seizures after consumption of synthetic
cannabinoids purchased online. J Addict

The New England Journal of Medicine
Downloaded from nejm.org by PASCAL BESMAN on September 9, 2015. For personal use only. No other uses without permission. Copyright © 2015 Massachusetts Medical Society. All rights reserved.
57. Rogeberg O. Correlations between cannabis use and IQ change in the Dundein cohort are consistent with confounding from socioeconomic status. Proc Natl Acad Sci U S A 2013;110:4251-4.
69. Mathern GW, Beninsig L, Nehlig A. Fewer specialists support using medical marijuana and CBD in treating epilepsy patients compared with other medical professionals and patients: result of Epilepsia’s survey. Epilepsia 2015;56:1-6.
71. Mathern GW, Beninsig L, Nehlig A. Fewer specialists support using medical marijuana and CBD in treating epilepsy patients compared with other medical professionals and patients: result of Epilepsia’s survey. Epilepsia 2015;56:1-6.
72. Mathern GW, Beninsig L, Nehlig A. Fewer specialists support using medical marijuana and CBD in treating epilepsy patients compared with other medical professionals and patients: result of Epilepsia’s survey. Epilepsia 2015;56:1-6.

Copyright © 2015 Massachusetts Medical Society.