Epilepsy affects more than 50 million people worldwide, or roughly 1% of the population, with a frequency of 4 to 10 per 1000 people per year. Treatment with antiepileptic drugs is frequently associated with poorly tolerated side effects, and the disease has proven to be medically refractory in more than 20% of patients. Resection of portions of the temporal lobe, amygdala, and hippocampus has been found to be an excellent option in the small fraction of patients suffering from temporal lobe seizures, with 86%–90% of patients experiencing freedom from seizures at 2 years. However, not all patients experience seizures emanating from the mesial temporal lobe, and of those that do, not all are candidates for resection. Therefore, there is a continued need for therapies that address focal epilepsies currently not amenable to existing techniques.

There are two general approaches to controlling seizures. The first approach, which is the basis of most antiepileptic medications, is to alter the excitability of the neural network to prevent seizures from ever occurring. This pharmacological approach has led to many new antiepileptic drugs over the past 5 decades, but the rate of intractable seizures has remained stable. A second approach for those intractable seizures is to abort them just as they begin, which has led to a search for medical devices that can stop seizures on demand. These “closed-loop” devices, using a strategy similar to implanted cardioverter defibrillators, would confirm the presence of a seizure and respond within milliseconds of onset, correcting the aberrant neurons while leaving normal neurons unaffected. Closed-loop devices have the advantage of limiting the intervention to specific times, which conserves power and leaves the normal neuronal or glial physiology unaltered during interictal periods. In the last decade, electrical stimulation of seizure foci via subdural or depth electrodes has gained attention. Two recent multicenter, double-blind, randomized controlled trials showed significantly reduced seizure frequencies when stimulation was applied. While the results are promising, perhaps offering a novel treatment modality for epilepsy, electrical stimulation lacks the precision of cell-type specificity.

Optogenetics is the ability to use optical systems for the stimulation or inhibition of a specific subset of neurons by introduction of genes for light-gated channels, known as opsins, neurons that initiate or propagate seizures may be silenced. To date, studies using this technique have been performed in animal models, and current efforts are moving toward more sophisticated nonhuman primate models. In this paper, the authors present a brief overview of the development of optogenetics and review recent studies investigating optogenetic modification of circuits involved in seizures. Further developments in the field are explored, with an emphasis on how optogenetics may influence future neurosurgical interventions.

### Optogenetic Tools

Optogenetics is a novel molecular intervention that relies on naturally occurring opsins, or light-gated ion channels, first identified in the early 1970s. The ion channel bacteriorhodopsin, found in the plasma membrane of...
*Halobacterium halobium*, undergoes a conformational change in response to the absorption of light of a specific wavelength, resulting in the outflow of protons. This efflux of ions generates a life-sustaining transmembrane gradient and electric potential that the bacterial cell then uses to drive adenosine 5'-triphosphate synthesis. At a time when the intricacies of the chemiosmotic theory were being clarified, opsins provided insight into a prevalent biochemical mechanism. However, these channels would not be applied to neuroscience until three decades later. In 2004, researchers effectively incorporated a proton channel (channelrhodopsin-1) and a cation channel (channelrhodopsin-2) into mammalian neurons. These opsins responded to flashes of light within milliseconds, resulting in neuronal depolarization in a precise manner. Genetic engineering improved upon several aspects of the channels, with increases in photocurrent amplitudes, protein expression, and action potential reliability. Additionally, opsins were identified that were able to permit depolarizing inflow of ions in a “red-shifted” spectrum, important for neural application, as red light has increased penetrance in brain tissue.

With the ability to turn on circuits, efforts were shifted to identification of a channel to inhibit them, which was achieved with a hyperpolarizing opsin from the halophilic archaeabacterium *Natronomonas pharaonis*. This pump, known as halorhodopsin, a heptahelical membrane protein, converts all-trans retinal to 13-cis retinal, exchanging 1 photon for 1 chloride ion. The addition of a C-terminal endoplasmic reticulum export motif onto the halorhodopsin protein resulted in improved plasma membrane localization, and further optimization increased the photocurrent amplitude. This version of the channel, termed eNpHR3.0, has been shown to be safely and reliably expressed in freely moving animals. Continued modifications have led to a variety of engineered opsins, and recently, a tandem construct has been developed that will enable expression of both an excitatory channelrhodopsin and inhibitory halorhodopsin in 1 vector. The number of available opsins continues to grow and will likely continue to be optimized for experimental and human application. Figure 1 depicts overall function of these channels.

Opsins are delivered to cells of interest by a variety of methods. Currently, injection of viral vectors into various regions may be most applicable to clinical scenarios. Subsets of cells within a region are selected using cell-type specific promoters, and several of these cell types are involved in epilepsy. The opsin genes themselves are carried in viral vectors, typically adenoassociated virus or lentivirus. Another method of introduction involves crossing Cre-recombinase knockin animals expressing Cre in specific neurons with animals transfected with the gene of interest. This results in expression of the opsin channel only in those neurons expressing Cre-recombinase in the cell type (for review, see Madisen et al.). An example of the gene is shown in Fig. 2, including the opsin gene itself and the regions needed for effective translocation.

Light delivery to activate the opsins is another area of research, with many different systems undergoing trials. Thin, cylindrical “optrodes” with a tip containing an optical fiber and recording electrode are reminiscent of deep brain stimulation electrodes. Other versions use an optrode implanted in 1 area, with a separate micro-LED array for recording. Before human use, these systems will need to meet the standards currently used for other implantable devices, with modifications made for patient tolerance and safety.

**Optogenetics in Epilepsy**

Optogenetics has now been applied to numerous neurological diseases as well as some nonneurological applications, including stimulation of heart muscle. Other cell types being targeted include medium spiny neurons in the basal ganglia for studying Parkinson disease, midbrain ventral tegmental area dopaminergic neurons in depression models, and dopamine D1-expressing neurons of the prefrontal cortex to investigate a circuit underlying temporal control of behavior. Probing various physiological mechanisms has also become a focus of optogenetic manipulation, with recent studies elucidating a role of ventral surface astrocytes in control of breathing. In epilepsy, optogenetics has become an attractive option to silence hyperactive neurons, with targeting of those believed to initiate or propagate seizures.

Epilepsy presents a difficult entity to study because of the large number of network pathologies that initiate or propagate seizures. However, there are several recent studies that aim to leverage the selectivity of optogenetics to analyze and control network dynamics in several animal models of seizures. Epilepsy is characterized by overactivity of various circuits, and consequently, many of the current investigations have focused on ways to silence or inhibit overactive circuits. The ability to selectively target mammalian hippocampal neurons was a first step toward probing this disease pathway, and was achieved using the inhibitory opsin halorhodopsin in organotypic cultures of mouse hippocampi. Following this achievement, halorhodopsin was used in selected principal glutamatergic neurons in the hippocampus. In this mouse model, the injection of a lentiviral vector into CA1 and CA3 pyramidal cells resulted in hyperpolarization in response to orange light (wavelength 573–613 nm). Action potentials were attenuated and paroxysmal depolarizing shifts were curbed, suppressing epileptiform activity. Importantly, the presence of the virally transfected opsins into neurons did not affect normal physiological properties of cells. To test this, organotypic hippocampal slices were subjected to complete darkness, and various properties of the cells were tested. Parameters including input resistance, resting membrane potential, action potential threshold and duration, amplitudes in response to hyperpolarizing and depolarizing pulses, and accommodation of cells to action potentials were similar in the halorhodopsin neurons in comparison with the controls.

Recently, this technology has been demonstrated in 3 different in vivo rodent models of epilepsy. In the rat tetanus toxin injection model, principal neurons were transfected with a lentiviral vector carrying halorhodopsin 2.0. Illuminating the transfected cells with 561-
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![Fig. 1. Type I microbial rhodopsins. Bacteriorhodopsins (BRs) and proteorhodopsins pump protons from the cytoplasm to the extracellular medium, and halorhodopsins (HRs) pump chloride into the cytoplasm; all 3 hyperpolarize the cell. Sensory rhodopsins (SRs) lack transmembrane ion transport in the presence of the His (histidine) kinase transducer protein Htr, and algal channelrhodopsins (ChRs) conduct cations across the membrane in both directions but always along the electrochemical gradient of the transported ions. In SRs and ChRs, proton translocation within the protein is linked to efficient photocycle progression, but these protons are not necessarily exchanged between the intra- and extracellular spaces. Reprinted from Cell 147: Zhang F, Vierock J, Yizhar O, et al., The microbial opsin family of optogenetic tools, pp. 1446–1457, 2011, with permission from Elsevier.](image1)

nm wavelength light resulted in decreased epileptiform EEG activity. Decreased high-frequency power was also observed, which correlate to bursts of activity in human epilepsy, and event frequency also diminished. The other two examples used a closed-loop strategy to abort seizures. In the first, a cortical stroke model was used in rats to promote seizures induced by photothrombosis, in which animals were first sensitized by injection of Rose Bengal dye, after which illumination resulted in increased light absorption and focal thrombosis. Intracranial EEG electrodes confirmed spontaneous epileptiform oscillations in the thalamus and cortex of injured rats but not in controls, correlating with thalamic network hyperexcitability. To determine whether these hyperactive thalamic cells could respond to light and silence seizure activity, a viral vector was injected consisting of the cell-type specific promoter Camk2a, the halorhodopsin eNpHR3.0, optimized with membrane trafficking signal motifs, and a reporter (eYFP; yellow fluorescent protein). The promoter ensured that only excitatory thalamic cells would be subject to viral transduction. Injections were performed stereotactically into the ventrobasal thalamus, followed by slice preparation for in vitro whole-cell recordings, as well as chronic recordings and stimulation in freely moving animals. To perform the chronic recordings, a device containing multiple EEG electrodes and chronic multisite optrode was implanted into the injected thalamus and peristroke cortex. Yellow light of a wavelength 594 nm was delivered, which resulted in silencing of epileptic EEG activity in thalamus and cortex and correlated with abotion of behavioral seizures as well. Lower intensity light was not effective, and the 594-nm light did not affect interictal EEG tracings or behaviors. The authors also developed a method of detecting and interrupting seizures online by routing to a real-time processor capable of determining seizure onset, which triggered light stimulation and resulted in seizure abortion within 1 second.

Following this report, optogenetic silencing of seizures was tested in a model of temporal lobe epilepsy. Here, mice were injected with kainic acid unilaterally into the dorsal hippocampus, and 2 methods of seizure control were explored: inhibition of excitatory principal cells and stimulation of inhibitory GABAAergic cells. Opsins were expressed by using transgenic animals, crossing mice expressing the Cre-lox recombinase in either principal cells or parvalbumin-containing GABAAergic interneurons with those expressing halorhodopsin or channelrhodopsin-2, respectively. With this method, only cells containing Cre were able to express the opsin of interest, ensuring specificity. As in previous studies, a reporter fluorescent protein was also expressed. Seizure detection was achieved via depth electrodes and tuning of seizure “signatures” (magnitude, amplitude, width, rate, and frequency) with custom software. For inhibition of principal neurons with halorhodopsin, amber light of wavelength 589 nm was delivered via an optical fiber implanted in the ipsilateral hippocampus upon seizure activity detection. All animals (n = 6) responded with silencing of electromagnetic seizures, with a majority (57% ± 12%) aborted within 1 second. The percentage of seizures aborting within 5 seconds was approximately 70%, and was unaffected by light duration (30 seconds vs 10 seconds). Further interrogation was made into the effect of red-shifted light, which has increased tissue penetrance, but is suboptimal for channelrhodopsin-2. This 635-nm light aborted 46% ± 12% seizures within 1 second of illumination. One animal in particular responded with 100% of seizures aborting within 1 second. The results from this portion of the experiment are shown in Fig. 3.

Activation of parvalbumin-GABAAergic neurons us-
ing channelrhodopsin-2 also resulted in reduced seizure duration. Blue light delivery (wavelength 473 nm) aborted 59% ± 11% of seizures within 5 seconds. This was a significant finding, as these cells make up less than 5% of hippocampal neurons, suggesting that even in a small population of targeted cells, optical stimulation of transgenic cells may be efficacious.

### Future Directions

These studies form the basis of our understanding of how to use light-activated channels as a therapeutic option for seizure control. Much remains to be learned from the use of opsins and light delivery, and many technical aspects must still be optimized before application to humans, several of which may prove to be substantial challenges. Among these aspects are investigation of the human immune response to these foreign proteins, the stability of viral vectors in neural tissue, and optimization of light delivery with chronically implanted devices. Thus far, application of this powerful tool has been successful in elucidating normal and pathological mechanisms, and some inquiry has been made into therapeutic applications, such as those presented here concerning epilepsy. The development of a nonhuman primate model is one recent achievement that may increase understanding of how optogenetics may be applied to humans.14

### Conclusions

Epilepsy is a prevalent disease conferring a significant burden to the people affected. Despite best pharmacological treatment, medical therapies leave more than 20% of patients with refractory disease. Resection of a seizure focus is one alternative treatment, but with the large number of patients in whom surgery is not an option, there continues to be a need for novel therapies. Recent progress has been made in the field of neurostimulation via subdural or depth electrodes, but efficacy has not yet been definitively borne out in clinical trials. One limitation to these previous therapies is the inability to precisely target aberrant cells and circuits while leaving others unaltered. For this reason, optogenetics has emerged as a possible technique, and several recent studies have investigated its usefulness in animal models. Although still in the early stages, introduction of light-gated ion channels into neurons via transgenic methods has shown to be possible. Light delivery to the region of interest, with the resultant opening and closing of these channels, leading to depolarization or hyperpolarization, has also been shown in numerous studies. Much remains to be understood before application to humans, including optimization of opsin introduction, light delivery, and device implantation. Furthermore, the safety of these channels will need to be determined. However, the ability to specifically alter

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**Fig. 3.** Seizure control in mice expressing halorhodopsin (HR) in principal cells in a model of temporal lobe epilepsy. a: Crossing CamK-Cre and Cre-dependent HR mouse lines generated mice expressing the inhibitory opsin HR in excitatory cells (Cam-HR mice). b: Experimental timeline. c–e: Example electrographic seizures detected (vertical green bars), activating amber light (589 nm) randomly for 50% of events (light: amber line, example in d; no light example in e). f: Typical example distribution of postdetection seizure durations (5-second bin size) during light (solid amber) and no-light internal control conditions (hatched gray). Inset: First 5-second bin expanded, 1-second bin size. Note that most seizures stop within 1 second of light delivery. Adapted from Nat Commun 4: Krook-Magnuson E, Armstrong C, Oijala M, Soltesz I, On-demand optogenetic control of spontaneous seizures in temporal lobe epilepsy, p. 1376, 2013, with permission from Elsevier. KA = kainic acid.
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a defined circuit in a reversible but reliable manner may provide insight into the mechanism of epilepsy as well as a possible therapy.

Disclosure

The authors report no conflict of interest concerning the materials or methods used in this study or the findings specified in this paper.

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