

Molecular Tools and Approaches for Optogenetics

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The mammalian brain poses a formidable challenge to the study and treatment of neuropsychiatric diseases, owing to the complex interaction of genetic, epigenetic, and circuit-level mechanisms underlying pathogenesis. Technologies that facilitate functional dissection of distinct brain circuits are necessary for systematic identification of disease origin and therapy. Recent developments in optogenetics technology have begun to address this challenge by enabling precise perturbation of distinct cell types based on molecular signatures, functional projections, and intracellular biochemical signaling pathways. With high temporal precision and reversible neuromodulation, optogenetics promises to improve existing disease models and advance our understanding of psychiatric conditions. In this review, we describe the current state of molecular optogenetic tools and future directions of development.

Key Words: Brain machine interface, channelrhodopsin, gene therapy, neuromodulation, neuropsychiatric disease, optogenetics

The brain is a particularly complicated organ to study because of its sheer diversity and heterogeneity of molecules, cells, and connections. Dysfunctions in specific cell types and circuits can lead to severe neuropsychiatric conditions (1). To dissect the mechanisms of psychiatric conditions, it is essential to investigate the pathogenic mechanisms at every level of brain function: from subcellular molecular signaling and cellular physiology to circuit-level functions. On the basis of clinical observations and insights to neuropsychiatric diseases, it is possible to generate cellular and animal models that recapitulate the multilevel symptoms of human patients (2). Systematic perturbation of the implicated circuit components in these disease models will help identify effective therapeutic targets and strategies. However, precise manipulation of the brain has been a difficult task to achieve. To fully recapitulate biological events, an ideal approach requires highly precise manipulations, including fast temporal control, cell-type specificity, and applicability within awake, behaving animals.

The development of optogenetic technologies enables such precise manipulations (3–6). It combines the temporal and spatial precision of light pulses with cellular specificity of genetic targeting; by coupling cellular behavior with light, it provides a fast, light-controlled approach to manipulate neural activities in intact brain circuits. The general strategy of optogenetics involves introducing a light-sensitive protein to a specific cell type, illuminating the targeted cells with defined parameters, and obtaining reliable readout of the cellular behavior. Compared with electrical microstimulation, optogenetic activation specifically targets the opsin-expressing neurons, whereas electrical stimulation may simultaneously affect diverse local, afferent, and passing axonal fibers (7,8) in addition to cellular somata near the electrode insertion site. Currently, the molecular toolbox of optogenetics (4,6) involves many different light-sensitive constructs capable of controlling a diverse range of neuronal functions, from electrical to biochemical signaling. Through a combination of molecular engineering and genomic discoveries of new light-sensitive proteins, the toolkit of neuron modulators is rapidly improving in specificity and versatility. At its present stage, optogenetics affords millisecond precision in controlling neuron activity, genetically defined cell type targeting, and

the ability to modulate neural circuit function in awake, behaving animals. With these features, optogenetics can be used to causally probe circuitries underlying complex behavior, dissect signaling pathways, and construct models of psychiatric disease through loss- and gain-of-function experiments. This review describes the optogenetic tools available for investigating neural circuit function and future directions in the development of this technology.

Modulating Neural Activity Using Light

The heterogeneous nature of brain tissue presents major challenges for selectively controlling subsets of well-defined neuron types in intact circuits. Most current neuromodulation techniques (i.e., electrical stimulation or pharmacologic intervention) either simultaneously affect surrounding cells and processes in addition to the target population, or have slow kinetics and reversibility. The lack of modulation specificity frequently limits the strength of conclusions drawn from conventional neuromodulation experiments. To overcome the spatial and temporal limitations of electrical, pharmacologic, and genetic neuromodulation approaches, various microbial and engineered opsins (Figure 1A) have been developed recently to control electrical and biochemical activities of neurons with cell-type selectivity, high temporal precision, and rapid reversibility. Because most neurons in the brain are not naturally light-sensitive, selective expression of the opsin genes in targeted neural populations makes it possible to control their activity specifically. Additionally, the fast on–off kinetics of microbial opsins make it possible to evoke or inhibit neural activities within milliseconds, on a time scale relevant to physiologic brain functions.

Activating Neurons Using Light

The discovery and introduction of algal light-sensitive cation-conducting channelrhodopsins into neurons has enabled fast optical control of genetically defined neuronal populations in intact circuits, both *in vitro* and *in vivo* (9). So far, channelrhodopsin genes from two algal species (*Chlamydomonas reinhardtii* and *Volvox carteri*) have been cloned and characterized in neurons (Figure 1A). Most notably, channelrhodopsin-2 from *Chlamydomonas reinhardtii* (ChR2) (10–12) and channelrhodopsin-1 from *Volvox carteri* (VChR1) (13) have proven to be a powerful pair of ChRs for controlling intact neural circuits using light. ChR2 is maximally activated by blue light around 480 nm, whereas VChR1 remains significantly light sensitive even at 589 nm, a wavelength at which ChR2 is no longer responsive. Therefore, ChR2 and VChR1 can be integrated in combinatorial neural control experiments both *in vitro* and *in vivo* to test the necessity and sufficiency of a variety of neural circuit functions. Both ChR2 and VChR1 have fast onset kinetics with time constants on the order of milliseconds and are therefore able to transduce high frequencies of millisecond-lasting light flashes into reliably evoked action potentials. However, because wildtype VChR1 has lower levels of photocurrent compared with ChR2,

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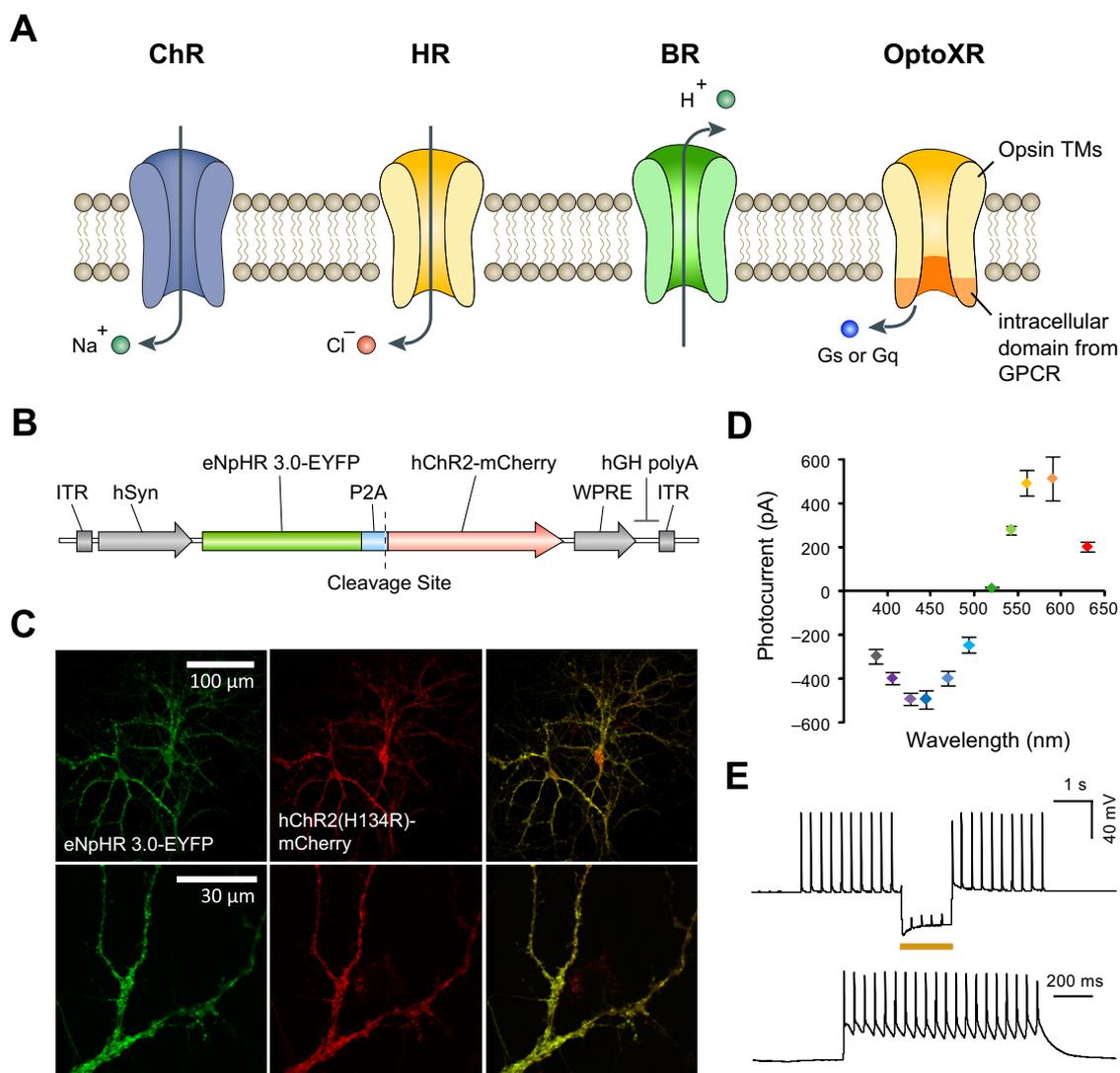


Figure 1. Microbial opsins as optogenetic tools. **(A)** A variety of natural and engineered opsins can be used to control the membrane potential and intracellular biochemical signaling of neurons. Channelrhodopsins (ChR) are light-activatable cation-conducting channels capable of activating neurons by depolarizing the membrane potential. Halorhodopsins (HR) and bacteriorhodopsins (BR) are light-activatable chloride and proton pumps capable of inhibiting neurons by hyperpolarizing the membrane potential. OptoXRs are engineered rhodopsin-G protein-coupled receptor (GPCR) chimeras useful for controlling intracellular G-protein-coupled signaling cascades. OptoXRs are engineered by replacing the intracellular domain of vertebrate rhodopsin with the intracellular loops from GPCRs. **(B)** A bi-cistronic vector used to express channelrhodopsin-2 (ChR2) and halorhodopsin (eNpHR3.0) in the same neuron. The two opsin genes are linked using a 2A self-cleavage linker. **(C)** Images of neurons coexpressing ChR2 and eNpHR3.0. **(D)** Activation spectra of neurons coexpressing ChR2 and eNpHR3.0. **(E)** Bidirectional control of neurons coexpressing ChR2 and eNpHR3.0. Blue light (470 nm) flashes induced action potential firing and yellow light (583 nm) facilitated spike inhibition. Panel A adapted with permission from Zhang *et al.* (4), and panels B through E are reprinted with permission from Gradinaru *et al.* (22), with permission from Elsevier, copyright 2010. Cl, chloride; Gq, G-protein Gq; Gs, G-protein Gs; hSYN, human synapsin I promoter; ITR, inverted terminal repeat; Na, sodium; TMs, transmembrane domains.

a chimera between *Chlamydomonas* ChR1 and *Volvox* ChR1 has been generated called C1V1. In conjunction with ChR2, C1V1 forms a powerful pair of opsins to control the activity of distinct groups of neurons *in vivo* simultaneously (14).

Mutagenesis studies could expand the function of channelrhodopsins by altering the spectral properties, conductance, or kinetics of the channels. Indeed, one mutant ChR2(H134R) has been shown to exhibit enhanced photocurrent (15), and three point-mutants of humanized ChR2 convert a brief pulse of light into a period of stable current influx that can last for many minutes (16). These latter mutant channels were generated by substitution of the C128 position in the original ChR2 sequence. All three mutants are activated by blue light (470 nm). Photocurrents generated by the opening of

ChR2(C128A) and ChR2(C128S) can be effectively terminated by a pulse of green light (542 nm). Other slow mutants were generated by mutating residue D156; this mutation has been shown to result in similar stabilization of the conducting state (14,17).

Substituting ChR2 residue E123 with T or A was found to accelerate channel closure kinetics and significantly increase the fidelity of fast optogenetic control (18). These E123 mutations can be combined with other gain-of-function modifications such as the H134R and T159C mutations (18,19) or membrane trafficking signals (14,20–22). The E123 mutations appear unique thus far because they eliminate the sensitivity of channel kinetics to membrane potential (19). Opsins of this class (E123 mutations alone or in combination with other modifications (18) are termed ChETAs (ChR

E1237/A). ChETA tools have been shown to deliver improved performance (up to 200 Hz) within intact mammalian brain tissue (18). A caveat is that faster deactivation is accompanied by reduced light sensitivity for long pulses of light.

Inhibiting Neurons Using Light

A number of light-activated anion pumps have been explored for neural activity silencing. In particular, the chloride-pumping halorhodopsin from *Natronomonas pharaonis* (NpHR) (23) has been shown to hyperpolarize neurons upon illumination with green or yellow light (Figure 1A). Also, NpHR and ChR2 have sufficient spectral separation and can be simultaneously expressed in the same neurons to facilitate bidirectional control of neural activity, to test both the necessity and sufficiency of a circuit component in neural function (Figure 1B–E). eNpHR3.0, an enhanced version of NpHR, was used along with bilateral optical fiber devices to inhibit the cholinergic neurons of the nucleus accumbens and elucidate a causal role for these rare cells in implementing cocaine conditioning in freely moving mice, which appears to operate via enhancing inhibition of inhibitory striatal medium spiny neurons (24).

Ecological prospecting has led to the discovery of additional neural silencers to eNpHR3.0, such as archaeorhodopsin-3 (Arch), a green light (~550 nm)-activated proton pump isolated from the archaeobacteria *Halorubrum sodomense* (25,26). Arch is well expressed in neurons and can drive large inhibitory currents. Other prominent channels for silencing neurons include bacteriorhodopsin (BR) and *Guillardia theta* rhodopsin-3 (GtR3). Although the peak absorbance of BR is approximately 560 nm (27), GtR3 is maximally excited by approximately 470 nm light (22). The diversity in their spectral properties renders them useful in selective silencing of neurons using multiple wavelengths.

Neuron Activity Sensors

Optical neural activity actuators can be combined with genetically encoded neural activity reporters to greatly enhance our ability to probe complex circuit-level mechanisms underlying neuropsychiatric diseases. By genetically targeting activity actuators and reporters into specific cell types, one can selectively perturb a specific cell population and simultaneously observe its effect on downstream circuit components. Two major classes of genetically encoded activity reporters have been developed to detect changes in intracellular calcium or membrane potential.

Calcium Sensors

Because of the involvement of voltage-gated calcium channels in propagating action potentials, much effort has been spent in developing fast and accurate calcium sensors to report neural activity. Traditional synthetic calcium dyes cannot be targeted to specific cell types and exhibit fast photobleaching. To overcome these limitations, genetically encoded calcium indicators (GECI) have been developed (28). One class of GECI, called GCaMP, consists of a circularly permuted green fluorescent protein that fluoresces in response to calcium influx (29,30). Modifications of the original GCaMP led to new variants, the best of which is GCaMP3, which contains amino acid mutations T116V, M66K, and N363D (28).

Voltage Sensors

Another route to measure neural activity is by monitoring the membrane potential changes in neurons. Genetically encoded voltage-sensitive fluorescent proteins (VSFPs) have been constructed by fusing fluorescent reporters with voltage-sensing domains. Upon voltage fluctuations, the voltage-sensing domain undergoes

conformational change, thereby shifting the spectral response of the reporter protein (31).

More recently, Kralj *et al.* (32) showed that Arch isolated from *Halorubrum sodomense* can also be used as a genetically encoded voltage sensor (32). After virally delivering Arch to rat hippocampal neurons cultures, they demonstrated that Arch exhibits high sensitivity and speed, both of which exceed that of existing VSFPs by a factor of 10. These properties allow Arch to detect single action potentials in vitro, which is a major step forward in achieving precise optical detection of neural circuit dynamics (32).

Modulation of Biochemical Signaling

Optogenetic control of well-defined biochemical signaling events can be achieved by constructing chimeras (33) between vertebrate rhodopsin and conventional ligand-gated G protein-coupled receptors (GPCR) (34,35). Many of the membrane receptors involved in neurotransmission and neuromodulation, such as dopaminergic, serotonergic, and adrenergic receptors belong to the GPCR family. When used as optogenetic tools, these vertebrate opsin chimeras are referred to as optoXRs (Figure 1A), which allow for optically controlled intracellular signaling with temporal resolution suitable for modulating behavior in freely behaving animals (34). OptoXRs provide speed and cellular precision that are not achievable with pharmacologic and genetic tools.

Targeting Optogenetic Modulators to Specific Neural Circuits

Neuropsychiatric diseases affect complex neural circuits. To precisely pinpoint the circuit-level mechanism of pathogenesis, we must be able to target brain tissue with cellular- and circuit-level specificity. Opsins can be selectively introduced into subsets of neurons based on their molecular signature, projection pattern, anatomic organization, and functional activity.

Targeting Specific Cell Types Using Molecular Signatures

The cellular substrate of pathology for many neuropsychiatric diseases tends to involve molecularly defined cell types. For example, parvalbumin, dopaminergic, and orexinergic neurons are affected in schizophrenia, Parkinson's disease, and narcolepsy, respectively. Many cell types can be targeted to express microbial opsins using the promoters of their signature genes, such as parvalbumin (36), tyrosine hydroxylase (37), and hypocretin (38) for the cell types described above. Depending on the genomic structure, some promoters are compact enough to be packaged into viral vectors so that specific cell types can be used to target across a wide range of animal models. Cell types with promoters that are too large to be packaged into viral vectors can be targeted using transgenic technologies, although this practice is mostly limited to mice and rats.

Viral Gene Delivery. Viral vectors are versatile tools for delivering genetic constructs into neurons. Both lentivirus and adeno-associated virus (AAV) vectors have been used to deliver optogenetic constructs into genetically defined cell populations. By fusing microbial opsin genes to cell-type-specific promoters, one can obtain restricted gene expression in the cell type of interest. For in vivo applications, stereotaxic injection can be used to deliver concentrated viral vectors into target brain regions. The advantages of using lentiviral and AAV vectors include stable long-term expression and high transgene levels (39). Although lentiviral vectors are permanently integrated into the genome of target cells, AAV vectors are mostly maintained extra-chromosomally. One major constraint of the viral delivery method is the limited packaging capacity: AAV can accommodate transgene constructs up to 5 kb, whereas lentiviral vectors

are limited to 10 kb for the complete packaged genome. Because of the package length constraint, AAV and lentiviruses can only be used to deliver relatively small promoters (~2 kb for AAV and ~5 kb for lentiviral vectors; microbial opsin gene plus a fluorescent tag can take up to 2 kb of sequence length). This exclusion of long promoters limits the expression specificity because some cell-type-specific promoters have much longer regulatory elements. Another caveat is that the spatial precision of stereotaxic injection is limited, especially when targeting small brain nuclei, and multiple injections are required to achieve efficient expression in large regions of brain tissue (4).

Transgenic Technology. For promoters that exceed the packaging limit of viral vectors, transgenic animal lines can be generated to express optogenetic constructs in a cell-type-specific fashion. Transgenic constructs can be introduced into transgenic mice either through bacterial artificial chromosomes or using short transgene cassettes containing recombinant cell-type-specific promoters (40). Several transgenic mouse lines expressing channelrhodopsin and/or halorhodopsin under the pan-neuronal Thy-1 promoter have been developed (41,42). Compared with viral delivery, one disadvantage is the amount of time needed to generate and breed transgenic animal lines. In addition, because of the untargeted nature of the transgene insertion, each transgenic animal line needs to be carefully characterized to avoid undesirable gene interruptions.

Conditional Expression

A major challenge with generating transgenic animals constitutively expressing microbial opsins is that some cell-type-specific promoters do not drive strong expression of the downstream gene. Because many of the opsins have low conductance or pumping properties (6), high levels of opsin expression are required for achieving reliable neural activation or silencing. To amplify the expression level, inducible AAV vectors carrying double-floxed inverse orf (DIO) of microbial opsin genes (DIO AAV vectors) (36,37) can be delivered into the brain of transgenic animals expressing Cre recombinase under the control of cell-type-specific promoters. Cre recombinase will recognize the double-floxed opsin construct and enable transgene expression after recombination (36,37). The DIO AAV system allows the gene of interest to be expressed using strong ubiquitous promoters, while deriving cell-type specificity from Cre recombinase expression. The availability of numerous constitutive and inducible Cre transgenic mouse (and rats in the near future) lines from the Allen Brain Institute for Brain Science, Jackson Laboratory, Gene Expression Nervous System Atlas (GENSAT), and other transgenic animal repositories provides a convenient solution for targeting a wide variety of cell types in the brain.

Circuit-Specific Targeting

Even cell populations identified via a common molecular signature can represent a heterogeneous mixture of cell types. For example, the tyrosine hydroxylase-positive dopaminergic neurons in the midbrain project to multiple downstream brain regions, including the striatum, nucleus accumbens, prefrontal cortex, habenula, and many other areas. Although dopaminergic neurons projecting to the dorsal striatum play a critical role in motor function, dopaminergic neurons projecting to the nucleus accumbens are involved in reward conditioning (37,43). The fact that a single molecularly defined cell population can project to multiple downstream brain regions and affect diverse brain processes warrants the need to improve cell-type-specific targeting based on their wiring. Circuit-specific targeting also holds the potential for uncovering the therapeutic mechanisms of deep brain stimulation, which has demonstrated therapeutic efficacy for neuropsychiatric diseases such as

Parkinson's disease (44) and major depressive disorder (45,46). However, because implanted electrodes nonselectively modulate all the cells as well as processes in the vicinity of implantation site, optogenetic stimulation can be used to identify the specific circuit components involved in the symptomatic rescue and facilitate more efficacious therapies.

Another approach to target specific neural projections can be achieved using trans-synaptic viral vectors or proteins. A number of trans-synaptic proteins and viral vectors with unique anterograde- or retrograde-transporting properties (47–49) may be engineered with recombinases to activate gene expression in subpopulations of neurons with cell-type and circuit-specificity. For example, fusion proteins containing Cre and either wheat germ agglutinin or tetanus toxin fragment C can be expressed in the cell bodies of one brain region and allow the recombinase to be transsynaptically delivered to the postsynaptic or presynaptic neurons in another brain region (22). Similarly, retrograde- and anterograde-transporting viral vectors (50,51) such as rabies virus, herpes simplex virus 1, and vesicular stomatitis virus can also be used to deliver recombinases or transgene cassettes in a projection specific fashion. When combined with conditional expression systems, either Cre-dependent transgenic mice or viral vectors, these projection-specific strategies will allow circuit-specific gene expression in a variety of animal models.

Future Developments

Optogenetics provides a basis for linking behavioral functions with cellular and circuit-level activity patterns. Systematic optogenetic perturbation of distinct cell types and brain projections has already begun to shed light on the cellular substrates of neurologic and psychiatric disorders including Parkinson's disease, anxiety, retinal degeneration, cocaine conditioning, social dysfunction, and depression (14,24,52–56). Tantalizing possibilities for the future of optogenetics lie in several directions, including the extension of optogenetic tools with distinct wavelengths of sensitivity and novel modes of control beyond electrical signaling, the emergence of gene expression technologies capable of targeting optogenetic tools to functionally defined cell populations and circuit elements, and the continued intersection of optogenetics with a variety of readout modalities and especially with high-throughput system-wide electrical and genome-wide molecular profiling technologies.

The optogenetics toolbox of the future will enable multiplexed control of several populations of neurons within the same experimental setup and will enable the combinatorial testing of signal processing roles of distinct neural circuit elements in complex behavior. Genome prospecting and molecular engineering of opsin genes from diverse microbial species have already expanded the optogenetic toolbox with a diverse palette of neuronal activators and inhibitors with distinct wavelengths of excitation, ion selectivity, and kinetics (16,18,26,57,58). The search continues for proteins sensitive to far-red wavelengths that will not only enable probing of deeper brain regions but will also facilitate the simultaneous investigation of multiple neural circuits when used in conjunction with other opsins. Recent successes in obtaining an atomic resolution structure of channelrhodopsin will catalyze opsin engineering efforts (59,60). In addition, proteins responsive to wavelengths from the ultrasound or magnetic frequency ranges will possibly enable the remote control of neural tissue without the necessity of any surgical implants. New opsins with selective conductance for potassium and calcium, ions important in synaptic events and intraneuronal signaling pathways, will result in precise dissection of the role of specific signaling processes in neural function.

Optogenetic control will also expand beyond electrical signaling by enabling precise modulation of endogenous transcription within specific neural populations. The development of light-sensing protein domains from plant and microbial species (61–64) will enable the design of photoactivatable enzymes and transcription factors. These light-sensing domains can be coupled with programmable DNA binding domains, including designer zinc finger proteins (65) and transcription activator-like effectors (66–69), to achieve optically controlled targeted modulation of endogenous transgene and RNA expression in mammalian genomes. The ability to regulate transcription within specific circuits in awake, behaving animals will be important for dissecting the molecular and epigenetic basis of neuropsychiatric diseases.

To fully understand the circuit-level mechanisms underlying neuropsychiatric disease, it is important to pinpoint the specific circuits underlying behavioral dysfunctions. The targeting of optogenetic tools into specific cells based on their activity during behavioral testing allows the identification of functional circuits. This can be achieved using molecular markers of neural activity (70) including immediate early genes such as *c-Fos*, *Zif268*, *Arc*, and *Npas4*. Leveraging the activity-dependent transcription regulatory elements of immediate early genes (71,72), it is possible to target the expression of opsin genes to distinct groups of neurons that are active during behavioral testing. For example, in an animal model of posttraumatic stress disorder (PTSD), activity-dependent targeting of opsins will enable the identification of cells involved in the phenotype. Subsequent activation of the targeted cells in the absence of trauma-inducing stimuli can be used to determine the necessity or sufficiency of specific cell populations in the manifestation of the PTSD phenotype. The epigenetic profile of these cells can be further analyzed to identify molecular signatures of PTSD.

A third major avenue for the future will be to integrate the use of optogenetics with other system-level technologies to facilitate global analysis of brain functions. A combination of optogenetics and functional magnetic resonance imaging (73) or large-scale multi-electrode arrays (74) will allow us to identify target brain regions and activity patterns corresponding to specific behavioral functions. Subsequent genome-wide transcriptome (74) and epigenome (75) analysis of the identified circuits will bridge molecular signatures with circuit dynamics and behavior. The integration of optogenetics with a diverse range of readout modalities will further bridge our understanding of nervous system functions and disease processes from molecules and cells to circuits and behavior.

As the optogenetics technology matures, it will continue to augment our ability to dissect the mammalian brain with increasing precision and expanding modes of control. The ability to specifically modulate the electrical, biochemical, and transcriptional activity of specific neuronal circuits will bring us one step closer to understanding the mechanism of normal and pathologic nervous system function. The intersection of optogenetics with other disciplines of biology and engineering will lead to novel therapeutic targets and innovative therapeutic interventions for neuropsychiatric diseases.

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