

pogil neuron function

pogil neuron function is a fundamental concept in neuroscience education, helping students and learners understand how neurons operate within the nervous system. This approach, often used in POGIL (Process-Oriented Guided Inquiry Learning) activities, emphasizes active engagement and critical thinking to explore the intricate processes that enable neurons to communicate. Understanding neuron function is essential for grasping how the brain processes information, responds to stimuli, and maintains overall bodily functions. In this article, we will delve into the core aspects of neuron function, exploring the structure of neurons, the electrical signals they generate, and how they communicate through synapses.

Understanding the Structure of Neurons

A solid understanding of neuron function begins with recognizing the key structural components of neurons, which are specialized cells designed to transmit information rapidly and efficiently.

1. Cell Body (Soma)

The cell body, or soma, is the central part of the neuron that contains the nucleus. It is responsible for maintaining the neuron's health and metabolic functions. The soma integrates incoming signals and generates outgoing signals to other parts of the neuron.

2. Dendrites

Dendrites are branched extensions from the cell body that receive signals from other neurons or sensory receptors. They serve as the primary input sites, collecting electrical signals (called graded potentials) from synapses.

3. Axon

The axon is a long, slender projection that conducts electrical impulses away from the cell body toward other neurons, muscles, or glands. It can vary in length from a fraction of a millimeter to over a meter in some animals.

4. Myelin Sheath

Many axons are covered by a myelin sheath, a fatty layer formed by Schwann cells in the peripheral nervous system or oligodendrocytes in the central nervous system. The myelin sheath acts as an insulator,

increasing the speed of electrical transmission along the axon.

5. Axon Terminals

At the end of the axon are the axon terminals, which form synapses with target cells. These terminals release neurotransmitters, chemical messengers that carry signals across synapses.

The Generation and Propagation of Electrical Signals

Neurons communicate through electrical signals known as action potentials. Understanding how these signals originate and travel along neurons is crucial in grasping neuron function.

1. Resting Membrane Potential

Neurons maintain a resting membrane potential, typically around -70 mV , due to differences in ion concentrations inside and outside the cell. This voltage difference is maintained by ion pumps and channels.

2. Stimulus and Depolarization

When a neuron receives a stimulus, ion channels in the membrane open, allowing positive ions (such as sodium, Na^+) to enter the cell. This influx causes depolarization, making the membrane potential more positive.

3. Action Potential

If depolarization reaches a specific threshold (around -55 mV), voltage-gated sodium channels open rapidly, leading to a significant influx of Na^+ ions. This rapid change results in the action potential, a brief electrical impulse that travels along the axon.

4. Repolarization and Hyperpolarization

Following the peak of the action potential, sodium channels close, and voltage-gated potassium (K^+) channels open. K^+ ions exit the cell, restoring the negative resting potential. Sometimes, the potential becomes slightly more negative than resting, a phase called hyperpolarization.

5. Refractory Period

During the refractory period, the neuron temporarily cannot fire another action potential, ensuring unidirectional signal propagation and proper timing.

Synaptic Transmission: Communication Between Neurons

Neurons do not connect directly; instead, they communicate across synapses via chemical signals called neurotransmitters.

1. The Synapse Structure

A synapse consists of:

- Presynaptic neuron: Sends the signal
- Synaptic cleft: The gap between neurons
- Postsynaptic neuron: Receives the signal

2. Neurotransmitter Release

When an action potential reaches the axon terminal, voltage-gated calcium (Ca^{2+}) channels open. The influx of Ca^{2+} triggers synaptic vesicles to fuse with the presynaptic membrane, releasing neurotransmitters into the synaptic cleft.

3. Signal Reception and Response

Neurotransmitters bind to specific receptors on the postsynaptic membrane, causing ion channels to open or close. This can result in excitatory or inhibitory postsynaptic potentials, influencing whether the postsynaptic neuron fires its own action potential.

4. Termination of Signal

Neurotransmitter action is terminated through:

- Reuptake into the presynaptic neuron

- Enzymatic degradation in the synaptic cleft
- Diffusion away from the synapse

Types of Neurons and Their Functions

Different neurons are specialized for various roles within the nervous system.

1. Sensory Neurons

Sensory neurons transmit information from sensory receptors (skin, eyes, ears) to the central nervous system. They detect stimuli such as light, sound, touch, and temperature.

2. Motor Neurons

Motor neurons carry signals from the central nervous system to muscles and glands, initiating actions like muscle contraction or gland secretion.

3. Interneurons

Interneurons connect sensory and motor neurons within the central nervous system. They process information, integrate signals, and coordinate responses.

Factors Affecting Neuron Function

Various factors can influence how neurons operate and communicate.

1. Ion Channel Functionality

Proper functioning of sodium, potassium, calcium, and chloride channels is vital for action potential generation and synaptic transmission.

2. Neurotransmitter Availability

The synthesis, release, and reuptake of neurotransmitters determine the strength and duration of synaptic signals.

3. Myelination

Myelin sheaths increase conduction velocity, enabling rapid responses. Demyelinating diseases like multiple sclerosis impair neuron function.

4. Neuroplasticity

The nervous system's ability to reorganize itself by forming new connections affects learning, memory, and recovery from injury.

Summary

In understanding **neuron function**, it is essential to grasp how neurons are structurally designed to transmit electrical and chemical signals efficiently. From the resting membrane potential to action potential propagation and synaptic transmission, each step plays a critical role in the overall function of the nervous system. Recognizing how different neuron types operate and how various factors influence their activity provides insight into the complex yet fascinating world of neural communication. This knowledge is fundamental not only for students studying neuroscience but also for anyone interested in understanding the biological basis of behavior, sensation, and cognition.

Frequently Asked Questions

What is the primary function of neurons in the human body?

Neurons are responsible for transmitting electrical and chemical signals throughout the nervous system, enabling functions like sensation, movement, and cognition.

How do POGIL activities enhance understanding of neuron functions?

POGIL activities promote active learning through exploration, collaboration, and application, helping students grasp complex concepts like neuron structure and signaling processes effectively.

What are the main parts of a neuron involved in its function?

The main parts include the cell body (soma), dendrites (receive signals), axon (transmits signals), and axon terminals (communicate with other neurons).

How does an action potential propagate along a neuron?

An action potential is generated when a neuron depolarizes, causing a wave of electrical charge to travel along the axon, transmitting the nerve impulse rapidly.

What role do neurotransmitters play in neuron communication?

Neurotransmitters are chemical messengers that cross synapses to transmit signals from one neuron to another, facilitating communication within the nervous system.

Why is the myelin sheath important for neuron function?

The myelin sheath insulates the axon, increasing the speed of electrical signal transmission and ensuring efficient communication between neurons.

How do POGIL activities help students understand the concept of neuron signaling?

POGIL activities allow students to explore and model how neurons generate and transmit signals, reinforcing their understanding through hands-on, collaborative learning.

What is the significance of the resting potential in neuron function?

The resting potential is the electrical charge difference across the neuron's membrane when inactive, serving as the baseline state necessary for initiating action potentials.

Additional Resources

Pogil Neuron Function: Decoding the Brain's Electrical Symphony

Understanding how neurons operate is central to unraveling the complexities of the nervous system, and the Pogil neuron function offers a fascinating glimpse into this intricate biological machinery. As a cornerstone of neural communication, neurons are specialized cells that transmit information across the nervous system through electrical and chemical signals. In this detailed exploration, we delve into the anatomy, physiology, and the dynamic processes that define Pogil neuron function, presenting insights that are essential for students, educators, and neuroscience enthusiasts alike.

Introduction to Pogil Neuron Function

The Pogil (Process-Oriented Guided Inquiry Learning) approach emphasizes active student engagement, critical thinking, and collaborative learning. When applied to neuroscience, Pogil strategies facilitate a deeper understanding of neuron function by breaking down complex concepts into manageable, inquiry-driven segments. At the heart of this methodology lies the detailed study of how neurons operate — their structure, signaling mechanisms, and role within the nervous system.

Understanding Pogil neuron function involves exploring key features such as:

- Neuron anatomy
- Resting potential
- Action potential generation
- Synaptic transmission
- Neural integration

Each component plays a pivotal role in ensuring efficient communication within the nervous system.

Neuronal Anatomy and Its Role in Function

Key Structures of a Neuron

A neuron consists of several specialized parts, each contributing uniquely to its overall function:

- Cell Body (Soma): Contains the nucleus and organelles vital for cell maintenance and metabolic activity.
- Dendrites: Branched extensions that receive incoming signals from other neurons.
- Axon: Long projection that transmits electrical impulses away from the cell body toward target cells.
- Myelin Sheath: Insulating layer surrounding the axon, facilitating rapid signal conduction.
- Nodes of Ranvier: Gaps in the myelin sheath that enable saltatory conduction.
- Axon Terminals (Synaptic Boutons): Structures that release neurotransmitters to communicate with downstream neurons or effector cells.

Understanding how these parts work together is fundamental to grasping Pogil neuron function, especially in processes like signal initiation and propagation.

Functional Significance of Neuronal Structures

- Dendrites act as the primary receivers of signals, converting chemical messages into electrical signals.
- The axon transmits these signals over long distances, often covering significant body areas.
- Myelin sheaths increase conduction velocity by enabling saltatory conduction, where action potentials leap between Nodes of Ranvier.
- Axon terminals facilitate chemical communication via neurotransmitter release, bridging electrical and chemical signaling.

Resting Potential: The Neuron's Electrical Baseline

Understanding Resting Membrane Potential

The resting potential is the electrical potential difference across the neuron's membrane when the neuron is not actively sending a signal. Typically, this is around -70 millivolts (mV), with the interior of the cell being negatively charged relative to the exterior.

Key factors maintaining resting potential include:

- Sodium-Potassium Pump: An active transporter that moves 3 Na^+ ions out and 2 K^+ ions in, maintaining concentration gradients.
- Leak Channels: Passive channels allowing K^+ ions to move freely, stabilizing the membrane potential.
- Negatively Charged Proteins: Large molecules inside the cell contribute to the negative internal charge.

Proper functioning of these elements ensures neurons are ready to respond swiftly to stimuli, forming the basis for all neural activity.

Action Potential: The Neural Signal in Motion

What Is an Action Potential?

An action potential is a rapid, transient change in membrane potential that propagates along the axon, serving as the fundamental message carrier in neurons. It involves a sequence of voltage changes driven by the opening and closing of specific ion channels.

Stages of an Action Potential

1. Depolarization: Triggered when a stimulus causes the membrane potential to reach the threshold (~ -55 mV), leading to opening of voltage-gated Na^+ channels. Na^+ influx causes a rapid rise in positive charge inside the neuron.
2. Repolarization: Following the peak ($\sim +30$ mV), Na^+ channels inactivate, and voltage-gated K^+ channels open. K^+ exits the cell, restoring negative internal charge.
3. Hyperpolarization: The K^+ channels remain open slightly longer than needed, making the membrane potential more negative than resting potential.
4. Return to Resting Potential: Na^+/K^+ pump and leak channels restore the original resting potential, preparing the neuron for subsequent signals.

Mechanisms Ensuring Unidirectional Signal Propagation

- Refractory Periods: During the absolute refractory period, Na^+ channels are inactivated, preventing backward propagation.
- Saltatory Conduction: In myelinated neurons, the action potential jumps between Nodes of Ranvier, increasing speed and efficiency.

Synaptic Transmission: Chemical Communication Between Neurons

From Electrical to Chemical Signal

Once an action potential reaches the axon terminal, it triggers the release of neurotransmitters—chemical messengers stored in synaptic vesicles. These molecules cross the synaptic cleft to bind to receptors on the postsynaptic neuron, initiating a new electrical signal.

Steps in Synaptic Transmission

1. Arrival of Action Potential: Depolarizes the axon terminal membrane.
2. Calcium Influx: Voltage-gated Ca^{2+} channels open, allowing calcium ions into the terminal.
3. Neurotransmitter Release: Elevated calcium causes synaptic vesicles to fuse with the membrane, releasing neurotransmitters.
4. Receptor Binding: Neurotransmitters bind to specific receptors on the postsynaptic membrane.
5. Post-Synaptic Response: Ion channels open or close, leading to excitatory or inhibitory postsynaptic potentials.
6. Termination: Neurotransmitter removal occurs via enzymatic degradation, reuptake, or diffusion.

Types of Neurotransmitters and Their Roles

- Acetylcholine: Involved in muscle activation and attention.
- Dopamine: Modulates mood, reward, and motor control.
- Serotonin: Regulates mood, appetite, and sleep.
- GABA: Primary inhibitory neurotransmitter.
- Glutamate: Main excitatory neurotransmitter.

Neural Integration and Signal Processing

Summation of Postsynaptic Potentials

Neurons integrate multiple signals via:

- Temporal Summation: Multiple signals arrive at the same synapse in quick succession.
- Spatial Summation: Simultaneous signals arrive from different synapses.

The net effect determines whether the neuron reaches the threshold to fire an action potential.

Neuronal Pathways and Networks

Neurons form complex circuits, enabling:

- Reflexes
- Sensory processing
- Motor coordination
- Cognitive functions

Understanding Pogil neuron function in these contexts involves analyzing how individual neuron properties contribute to larger system behaviors.

Implications and Applications of Pogil Neuron Function

- Educational Impact: The Pogil approach promotes active learning, encouraging students to explore neuron functions through inquiry, diagrams, and problem-solving.
- Clinical Relevance: Insights into neuron function underpin treatments for neurological disorders such as epilepsy, multiple sclerosis, and neurodegenerative diseases.
- Technological Advances: Understanding neuronal signaling informs the development of neural prosthetics, brain-computer interfaces, and neuropharmacology.

Conclusion

The Pogil neuron function encapsulates the elegant choreography of electrical and chemical processes that enable the nervous system to operate seamlessly. From the structural intricacies of neurons to the dynamic processes like action potentials and synaptic transmission, each element is vital for neural communication. By adopting a Pogil-based inquiry approach, learners can deepen their understanding of these complex mechanisms, fostering a more comprehensive and engaged appreciation of neuroscience.

In essence, neurons are not just biological cells but are the fundamental units of intelligence, sensation, and action. Their function, as explored through Pogil strategies, exemplifies the marvel of biological engineering—a finely tuned system that continues to inspire scientific discovery and innovation.

Pogil Neuron Function

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pogil neuron function: *Basic Concepts of Neuronal Function* Don L. Jewett, 1984

pogil neuron function: Netrin-1 Modulation of Hippocampal Neuron Structure and Function Ian Beamish, 2018 Axon guidance cues such as netrin-1 are known to be important for the development of neural circuits, however, little is known regarding possible roles in adulthood. Recent work from our lab has demonstrated that the netrin-1 receptor deleted-in-colorectal-cancer (DCC) regulates the structure and function of mature synapses in the mammalian brain. Here, we demonstrate that netrin-1 protein is readily detectable in the adult brain and is necessary for the expression of long-term potentiation (LTP) at the Schaffer collateral hippocampal synapse. Next, using genetic, biochemical, and imaging approaches, we show that netrin-1 is secreted from cultured hippocampal neurons in an activity-dependent manner. Bath application of netrin-1 results in an increase in intracellular calcium levels within hippocampal neurons and leads to a dramatic potentiation of synaptic responses at the CA3-CA1 synapse. Classic long-term synaptic potentiation (LTP) induces phosphorylation of GluA1 by PKC and CaMKII at S831, resulting in increased channel conductance, and by PKA at S845, that can promote trafficking of GluA1 to synaptic sites. Here, we show that netrin-1-induced LTP depends on exocytosis of GluA1-containing AMPARs. Moreover, we show elevated GluA1 levels are accompanied by significant increases in phosphorylation of both S831 and S845 sites, as well as increases in phospho-CaMKII[alpha] levels. Lastly, we demonstrate that exogenous application of netrin-1 specifically increases the volume of thin type dendritic spines in organotypic hippocampal cultures. These results point to a critical role for activity-dependent netrin-1 release in the regulation of synaptic transmission and plasticity via netrin-1 mediated modulation of serine residues on GluA1. --

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pogil neuron function: The Retinal Müller Cell Vijay Sarthy, Harris Ripps, 2014-02-02 The human brain contains more than a billion neurons which interconnect to form networks that process, store, and recall sensory information. These neuronal activities are supported by a group of

accessory brain cells collectively known as neuroglia. Surprisingly, glial cells are ten times more numerous than neurons, and occupy more than half the brain volume (Hydén, 1961). Although long considered a passive, albeit necessary, component of the nervous system, many interesting and unusual functional properties of glial cells are only now being brought to light. As a result, the status of these cellular elements is approaching parity with nerve cells as a subject for experimental study. The term glia (or glue) seems today to be a misnomer in view of the diverse functions attributed to glial cells. Experimental studies in the last three decades have clearly established that the behavior of glial cells is far from passive, and that they are at least as complex as neurons with regard to their membrane properties. In addition, glial cells are of importance in signal processing, cellular metabolism, nervous system development, and the pathophysiology of neurological diseases. The Müller cell of the vertebrate retina provides a splendid example of an accessory cell that exhibits features illustrating every aspect of the complex behavior now associated with glial cells.

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Nerve Membranes: A Study of the Biological and Chemical Aspects of Neuron-Glia Relationships presents the various aspects of neuronal and glial structure and function. This book provides an interdisciplinary approach to the analysis of neuron-glia relationships and of membranes in the nervous system. Comprised of seven chapters, this book begins with an overview of the function of the biological membranes to improve, retard, and regulate the rate of cellular reactions. This text then determines the differences in the organization of the cells in the nervous system in the vertebrates and the invertebrates. Other chapters examine the role of certain intermolecular forces and of water in the organization of lipid-protein and lipid-lipid associations. This book reviews as well the theories of biological membrane structure and considers how these contribute towards understanding the methods by which membranes perform their role. This book is a valuable resource for neuroscientists, neurochemists, and researchers.

pogil neuron function: Purinergic Signalling in Neuron-Glia Interactions Derek J. Chadwick, Jamie A. Goode, 2006-05-01 ATP, the intracellular energy source, is also an extremely important cell-cell signalling molecule for a wide variety of cells across evolutionarily diverse organisms. The extracellular biochemistry of ATP and its derivatives is complex, and the multiple membrane receptors that it activates are linked to many intracellular signalling systems. Purinergic signalling affects a diverse range of cellular phenomena, including ion channel function, cytoskeletal dynamics, gene expression, secretion, cell proliferation, differentiation and cell death. Recently, this class of signalling molecules and receptors has been found to mediate communication between neurons and non-neuronal cells (glia) in the central and peripheral nervous systems. Glia are critical for normal brain function, development and response to injury. Neural impulse activity is detected by glia and purinergic signalling is emerging as a major means of integrating functional activity between neurons, glia and vascular cells in the nervous system. These interactions mediate effects of neural activity on the development of the nervous system and in association with injury, neurodegeneration, myelination and cancer. Bringing together contributions from experts in diverse fields, including glial biologists, neurobiologists and specialists in purinergic receptor structure and pharmacology, this book considers how extracellular ATP acts to integrate communication between different types of glia, and between neurons and glia. Beginning with an overview of glia and purinergic signalling, it contains detailed coverage of purine release, receptors and reagents, purinergic signalling in the neural control of glial development, glial involvement in information processing, and discussion of the interactions between neurons and microglia.

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pogil neuron function: Glial ↔ Neuronal Signaling Glenn I. Hatton, Vladimir Parpura, 2004-05-31 Glial Neuronal Signaling fills a need for a monograph/textbook to be used in advanced courses or graduate seminars aimed at exploring glial-neuronal interactions. Even experts in the field will find useful the authoritative summaries of evidence on ion channels and transporters in glia, genes involved in signaling during development, metabolic cross talk and cooperation between astrocytes and neurons, to mention but a few of the timely summaries of a wide range of

glial-neuronal interactions. The chapters are written by the top researchers in the field of glial-neuronal signaling, and cover the most current advances in this field. The book will also be of value to the workers in the field of cell biology in general. When we think about the brain we usually think about neurons. Although there are 100 billion neurons in mammalian brain, these cells do not constitute a majority. Quite the contrary, glial cells and other non-neuronal cells are 10-50 times more numerous than neurons. This book is meant to integrate the emerging body of information that has been accumulating, revealing the interactive nature of the brain's two major neural cell types, neurons and glia, in brain function.

pogil neuron function: The Functional Roles of Glial Cells in Health and Disease

Rebecca Matsas, Marco Tsacopoulos, 2012-12-06 Thirty-five years ago, when Stephen Kuffler and his colleagues at Harvard initiated a new era of research on the properties and functions of neuroglial cells, very few neuro scientists were impressed at the time with the hypothesis that neuroglial cells could have another, though more subtle, role to play in the nervous system than to provide static support to neurons. Today, very few neuroscientists are unaware of the fact that multiple interactions between neurons and glial cells have been described, and that they constitute the basis for understanding the function and the pathology of the nervous system. Glial cells outnumber neurons and make up about one-half of the bulk of the nervous system. They are divided into two major classes: first, the macroglia, which include astrocytes and oligodendrocytes in the central nervous system, and the Schwann cells in the peripheral nervous system; and second, the microglial cells. These different classes of glial cells have different functions and contribute in different ways in the development, function, and the pathology of the nervous system.

pogil neuron function: The Neuron in Context Vanessa Lux, 2024-04-26 Neuroscience has largely abandoned its localizationist and mechanistic framework of the 20th century. The plastic, embodied, and network character of our nervous system is widely acknowledged and systems theory approaches to consciousness dominate the field. However, the underlying neuron theory has not changed. The neuron doctrine, conceptualizing the single neuron as atomistic, one-directional source of neural function, still provides the template for our understanding of these basic elements of our nervous system and the material foundation of consciousness. Yet, the single neuron does not exist as an isolated unit. It is embedded within multiple cellular, structural, and functional contexts, and highly depends on them for its development, neural activity, and survival. The book discusses the constraints of the neuron doctrine and its pragmatic reductionism in the light of the growing knowledge about the brain's connectivity, plasticity, and systemic and embodied nature. To overcome these constraints, the author argues for a new neuron theory, depicting the neuron as bidirectional hub which is at the same time source and product of neural function. This bidirectionality is further characterized by spatial and time dimensions, placing the neuron within a multi-level pathway model of psychobiological development from the perspective of Developmental Embodiment Research. Furthermore, the author discusses the potential of neuroepigenetic markers to characterize the neuron and its range of plasticity within this developmental perspective. With its focus on neuroepigenetics, the book addresses a knowledge gap in the current study of the neural foundations of psychological functions. The multi-level and bidirectional perspective is already realized in approaches coming from developmental systems theory, which model neural function at the connectome level, and it also fits with approaches investigating feedback loops underlying neural activity at the single cell level. At both these levels, the spatial and the time dimensions are well characterized, either as changing connectivity patterns across different age groups, or as synaptic feedback loops underlying neural activation patterns. However, for the intermediate level of small neural populations, which is currently the main target for studies investigating the neural basis of specific psychological functions, this characterization turned out to be more challenging. Multi-cell recordings have provided a first glimpse into the complex interaction patterns of these small neural networks, but they are limited to the recording period and do not provide information about the long-term developmental and activation history. Here, neuroepigenetic markers could be of use. Due to their relative stability and, at the same time, environmental sensitivity,

neuroepigenetic markers represent an additional layer of information in which, to a certain degree, the cell's metabolic and activation history is aggregated over time. This information is available at the single neuron level but could also be modeled as aggregated information for small neural populations and the supporting cellular context. Looking through this "epigenetic lens" adds to our understanding of the neuron as bidirectional hub by emphasizing the molecular correlates of functional stabilization and their contextual prerequisites. These prerequisites reach from the immediate cellular context to the social-cultural contexts which shape the culturally specific modes of acquisition of psychological functions throughout the lifespan. Accounting for this multilayered contextuality of the neuron and its function affords to repositions the relationship between neuroscience and psychology in their joint effort to unravel the material basis of consciousness. This provides new challenges but also new perspectives for theoretical psychology. The book presents these current developments and debates to researchers, graduate students, and interested professionals and practitioners working in neuroscience, epigenetics, psychiatry, psychology and psychotherapy. It also provides a basic introduction into neuroepigenetics, its mechanisms, and first findings for graduate students as well as interested professionals and practitioners working in psychiatry, psychology, and psychotherapy.

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pogil neuron function: *Light-Controlled Modulation and Analysis of Neuronal Functions* Carlo Matera, Piotr Bregestovski, 2023 Light is an extraordinary tool allowing us to read out and control neuronal functions thanks to its unique properties: it has a great degree of bioorthogonality and is minimally invasive; it can be precisely delivered with high spatial and temporal precision; and it can be used simultaneously or consequently at multiple wavelengths and locations.

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