Kainate Receptors Are the Key to Understanding Synaptic Plasticity, Learning and Memory (Review)

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Glutamatergic signaling is one of the main types of excitatory synaptic transmission in the brain. It plays a key role in the normal brain function and the cognitive performance. Glutamatergic signaling failure is associated with brain disorders; therefore, this system is considered an essential target of therapeutic interventions. Glutamatergic synaptic transmission is mediated by a set of ionotropic and metabotropic glutamate receptors including the kainate receptors. These receptors (both ionotropic and metabotropic) are involved in the process of synaptic transmission by modulating the excitation/inhibition balance. The modulatory effect of kainate receptors is mediated by the mechanisms that involve the presynaptic and postsynaptic endings, the rhythmic activity of the neural network, the function of the astroglial network, and the neuron-glial interaction. Thus, a dysfunction of kainate receptors can lead to deviations in the balance between excitation and inhibition, disorders of the neuronal networks, and even epileptiform manifestations. The present report reviews the major mechanisms of ionotropic and metabotropic activation of kainate receptors involved in the regulation of synaptic transmission, plasticity, learning, and memory.

Key words: kainate receptors; synaptic transmission; metabotropic effect; G-proteins; long-term potentiation; brain rhythms; cognitive functions; epilepsy.

According to the current concept, there are three major groups of ionotropic glutamate receptors, named after their selective agonists NMDA (N-methyl-D-aspartate), AMPA (α-amino-3-hydroxy-5-methyl-4-isoxazole propionic acid) and kainate (kainic acid), and three groups of metabotropic receptors (mGluR), associated with the G proteins (the GTPase family); these two types of glutamate receptors differ by their structure and mechanism of action. The ionotropic glutamate receptors mediate most of the excitatory transmission in the CNS and play a major role in the regulation of synaptic plasticity. The glutamatergic signaling is highly important for synaptic transmission, normal brain functioning, and cognitive performance; therefore, interruptions in the normal signaling process can manifest in clinical disorders. Studying the neurotransmission pathways mediated by glutamate receptors is one of the main research tasks of neurobiology today.

Kainate receptors are unique among the family of glutamate receptors: they represent both the classical ionotropic and the non-classical metabotropic functions. The role of kainate receptors (KAR) is not known well enough as compared with the AMPA and NMDA receptors (AMPAR and NMDAR). A major reason for that is that in addition to ionotropic effects, the kainate receptors also have a metabotropic effect, which is not typical. Another problem is that both KAR and AMPAR show some overlapping sensitivity to most competitive antagonists such as CNQX and NBQX; in addition, a selective AMPA agonist can activate kainate receptors. Despite the fact that new selective agonists for KAR are being developed, not all of them have due affinities. This issue complicates studying the receptor functions at the full scale.

It is known that the long-term potentiation requires the activation of NMDA receptors and the increase in expression of AMPA receptors on the postsynaptic membrane surface and/or the increase in the dendritic spine size. However, there is also an independent NMDA receptor that employs the metabotropic signaling pathways and stimulates the AMPAR incorporation into the membrane; this one is activated by kainic acid — a KAR agonist.

A good object for studying the presynaptic functions of kainate receptors is the synapses of unmyelinated axons, called mossy fibers, formed by granular cells of
the hippocampal dentate fascia, where a high level of expression of the receptor subunits is found. The kainate receptors participate in the neurotransmission between the cerebral fascia cells and the pyramidal neurons of the hippocampal CA3 area, showing a positive response to the release of the neurotransmitter from the presynaptic terminal and thus depolarizing the axons. Most of the studies on the synapses of mossy fibers in the hippocampal CA3 area in vertebrates address the presynaptic kainate receptors; however, the postsynaptic functions of these receptors are less studied.

The KAR are expressed mostly in the CNS neuronal cells. Along with that, glial cells also show the presence of all subunits of this receptor (mRNA or proteins), often in co-expression with the AMPA receptor subunits. In astrocytes, the receptor subunits are located throughout the entire cell, whereas in oligodendrocytes, they are mainly limited to the soma. In this case, the glutamate-induced membrane current in oligodendrocytes is fully generated via the AMPA and kainate receptors in both cell culture and brain sections.

The modulating function of kainate receptors is based on the mechanisms involving the presynaptic and postsynaptic endings, the rhythmic activity of the neural network, the function of the astroglial network, and the neuron-glial interactions. Thus, a dysfunction of kainate receptors can distort the excitation/inhibition balance and provoke one or more CNS disorders including manifestations of epileptiform brain activity.

**Classification and functions of ionotropic glutamate receptors**

There are three main types of ionotropic glutamate receptors (iGluRs), named after selective agonists: NMDA, AMPA, and kainate [1]. These receptors mediate most of the excitatory neurotransmission throughout the CNS and play a major role in synaptic plasticity. Another type of receptors is represented by δ receptors; those are also part of the iGluRs group, since they share homologous fragments with the subunits of NMDAR, AMPAR, and KAR. Notably, they do not bind glutamate, and no specific endogenous ligands have been identified [2].

The AMPA receptors are tetrameric ion channels, which largely conduct Na⁺ and K⁺ ions; they can also be permeable to Ca²⁺, which depends on the combination of their subunits [3]. The NMDA receptors are tetrameric complexes that can serve both voltage-dependent and ligand-dependent ion channels [4]. The kainate receptors are composed from five subunits of GluK1-5 [5] that combine to form tetrameric ion channels, which are mainly permeable to Na⁺ and K⁺ [6].

In most cases, the permeability of KAR to Ca²⁺ is very low and depends on the composition of the given receptor complex; the specific combination of subunits determines the KAR function [7]. The KAR subunits are similar to the AMPAR and NMDAR subunits [8], but their presence in the CNS is more limited. The conductance of the KAR channel is similar to that of the AMPAR channel, which is about 20 ps, but the rise and decay of postsynaptic potentials generated by KAR are slower than those of AMPAR [9].

To activate KAR [10–12], kainic acid or domotic acid are needed; these two are strong high affinity agonists of KAR and AMPAR [13]. High concentrations of kainate cause recurrent continuing limbic seizures in rodents; their manifestations resemble those in human temporal lobe epilepsy [14]. Kainate provokes behavioral changes, mitochondrial dysfunction, neuron degeneration, and can eventually lead to death [11, 15]. This detrimental effect of kainate (a non-decomposable analogue of glutamate that is 30 times more toxic) is called excitotoxicity [15].

Kainic acid causes depolarization of virtually all types of brain neurons in mammals; this effect is mediated by the activation of KAR and AMPAR and depends on kainate concentration. The currents induced by high concentrations of kainic acid (>100 μM) occur mainly through AMPAR, since these receptors populate the neuron membranes at a higher density than the kainate receptors [10]. Also, KAR and AMPAR show some overlapping sensitivity to most competitive antagonists, such as CNQX and NBQX [16]. In addition, AMPA — the major AMPAR agonist — can activate various KAR [8]. For a long time, the lack of specific antibodies to different KAR subunits made it difficult to study the distribution of these receptors. Specific antibodies against the KAR — GluK2, 3 and GluK5 subunits are now available, although not all of them show sufficient selectivity [8]. The low affinity GluK1–3 subunits with their low affinity for kainic acid can assemble with each other and form functional homomeric receptors. In contrast, the high affinity subunits of GluK4, 5 have to combine with GluK1–3 to become functional heteromeric receptors [16–18]. The GluK1 subunit is mainly expressed in the hippocampal and cortical interneurons, and the GluK2 subunit is largely present in pyramidal cells of the hippocampus, cerebellum and cortical pyramidal cells [19]. The GluK1 and GluK2 subunits undergo editing at the mRNA level. The replacement of glutamine with arginine (most frequent substitute) brings about an almost complete loss of Ca²⁺ permeability [20]. The GluK3 subunit is less common; it appears in the IV layer of the neocortex and in the dentate gyrus of the hippocampus [21]. GluK4 is mainly expressed in pyramidal neurons of the hippocampal CA3 area, the dentate gyrus, the neocortex and Purkinje cells, whereas GluK5 is abundant throughout the entire brain [22].

The kainate receptors play a role in fine tuning of the balance between excitation and inhibition in the CNS [12]. They participate in synaptic transmission both at presynaptic [23, 24] and post-synaptic sites [25, 26], modulating the excitatory and inhibitory synaptic transmission [27–29]. Despite their role in neurotransmission, inhibiting the kainate receptors (or
Role of presynaptic kainate receptors in synaptic transmission

Presynaptic KAR play a crucial role in regulating the process of neurotransmission [33–34]. Exogenous KAR agonists regulate the release of the neurotransmitter both in the excitatory and inhibitory synapses (in the biphasic regimen), depending on the type of the synapse and the concentration of the agonist [35–37]. They modulate neurotransmission between the granular cells of the dentate fascia and the pyramidal neurons of the hippocampal CA3 area. Granular neurons of the dentate fascia receive information from the neocortex and transmit it to the hippocampus through their axons — mossy fibers. The bundles of these unmyelinated axons run along the layer of apical dendrites of pyramidal neurons of the CA3 area and form synaptic complexes on them [38]. These synaptic complexes are controlled by high affinity KAR that are activated by the release of glutamate from axons, so the presynaptic receptors can actively depolarize presynaptic boutons or axons, thus showing a positive response to the release of the neurotransmitter [39–42]. A similar effect can be caused by high-frequency stimulation (25–100 Hz) of moss fibers. This positive feedback mechanism, which leads to enhanced activation of KAR and release of glutamate does not depend on activation of NMDA receptors [43]. It has been shown that the presynaptic KAR on mossy fibers are permeable to Ca2+ and that the influx of Ca2+ through these receptors facilitates the release of glutamate from the presynaptic terminal and the release of Ca2+ from internal calcium stores [44]. The released glutamate binds to glutamate receptors on the postsynaptic membrane, initiating the release of Mg2+ and blocking NMDA receptors. The influx of Ca2+ through NMDAR channels initiates a chain of events that can lead to long-term potentiation [45]. However, the permeability of presynaptic KAR to Ca2+ has not been confirmed; besides, if the synaptic transmission is susceptible to blockade by KAR antagonists at low Ca2+ concentrations, then high concentrations of Ca2+ would overcome this block, indicating an alternative pathway for Ca2+ to enter the presynaptic bouton [43]. Similar Ca2+ dependence has been found for the kainate receptor-mediated regulation of the inhibitory synaptic transmission in the medial prefrontal cortex [46].

In the hippocampal CA1 area, presynaptic KAR, when activated by external glutamate or glutamate released from the Shaffer collaterals, facilitate the release of the CNS inhibitory mediator γ-aminobutyric acid (GABA). The endogenous glutamate can boost the inhibitory postsynaptic currents (IPSC), whereas exogenous agonists can either facilitate or inhibit the transmission depending on their concentrations [32, 47]. It has been shown that by depolarizing the interneurons with kainic acid, presynaptic KAR can increase the extracellular concentration of GABA. This is a secondary decrease in the GABAergic inhibition of pyramidal cells by the negative feedback mechanism [48–52].

The role of presynaptic receptors in the glutamatergic excitatory postsynaptic currents (EPSC) induced in the mossy fibers was also studied [29, 53, 54]. High concentrations of kainic acid suppress the synaptic transmission [55–57]. Low kainate concentrations, on the contrary, enhanced the synaptic transmission via both AMPAR and NMDAR [58]. The latter effect was blocked by KAR antagonists, which indicates that KAR participate in this bi-directional mechanism [59]. According to some data, kainic acid increases the frequency of spontaneous IPSC in pyramidal hippocampal cells [35, 50–52, 60–61], which is explained by depolarization of interneurons [35, 50, 51] or increased axonal excitability [61]. Other studies, on the contrary, show a decrease in the amplitude of IPSC induced in the hippocampal pyramidal neurons [28, 61], which is challenged by other investigators [61, 62] who failed to observe such a decrease in IPSC [38].

Thus, studies of the interactions of synaptic-related cells (interneuron-interneuron and interneuron-pyramidal neuron/dentate fascia cell) in CA1, CA3 and the dentate gyrus of the hippocampus showed that the effect of KAR activation on the inhibitory synaptic transmission depends on the type of pre- or post-synaptic cells, concentration of the agonist, synapse type, expression of subunits and receptor specificity [38].

The non-classical metabotropic activation of kainate receptors

It is known that the long-term potentiation requires the activation of NMDA receptors [63] and can include both the recirculation-dependent increase in the surface expression of AMPAR on the postsynaptic membrane and the increase in size of the dendritic spine [64]. However, there is also an NMDAR-independent mechanism that stimulates the incorporation of AMPA receptors [65, 66] into the membrane and is mediated by the metabotropic action of KAR in response to kainic acid. This mechanism of KAR signaling via G-proteins, in particular through the Go family and other secondary messengers, distinguish the KAR from the family of glutamate-directed ion channels [67]. This signaling mechanism is not well studied because KAR have a typical topology of ligand-dependent ion channels that do not contain normal motifs on C-terminal domains that would support direct binding to G proteins. This may indicate the existence of additional intermediary proteins acting within the receptor–G protein signaling complex [9]. This non-classical pathway involves the activation of protein kinase C and phospholipase C, which induce the NMDAR-independent hippocampal long-term potentiation, and cause structural changes (in
size and shape) in neuronal spines of the hippocampal CA1 region [65]. There is also a KAR-mediated NMDAR-independent long-term suppression, which can be caused by prolonged low-frequency stimulation or postsynaptic depolarization [66]. This kind of synaptic plasticity, independent of NMDAR activation, exists also in other areas of the CNS, which indicates the importance of such non-classical signaling. The above studies show that KAR have a variety of mechanisms for fine tuning of neuron activity; therefore, more information on the physiological role of this metabotropic KAR signaling is necessary to better understand its role in the CNS [67–69].

The role of postsynaptic kainate receptors in synaptic transmission

Post-synaptic kainate receptors have been found in several locations of the CNS, including the hippocampus [27, 70], the spinal cord [71], the somatosensory cortex [72], the cerebellum [73], and the medial entorhinal cortex [74].

Studies on dissociated neuronal cultures demonstrate that the activation of postsynaptic kainate receptors stimulates the surface expression of receptors containing the GluK2 subunit [74], and also the development of filopodia as well as the axonal and dendritic growth [75–77]. This occurs with the help of metabotropic signaling, which facilitates the recycling of KAR in the spines with the help of Rab11-dependent release of the recycling endosomes towards the head of the spine. These endosomes are tubular membrane structures associated with microtubules that mediate the recirculation in the dendritic spines [78, 79]; it is important to note that these membranes contain the small Rab11 protein from the GTPase family [80, 81]. This non-classical metabotropic pathway mediates a positive feedback system that leads to an increase in the surface-expressed postsynaptic KAR associated with GluK2; the latter exemplifies a previously unknown self-regulated pathway. This pathway provides additional flexibility to synaptic regulation and is likely to have important physiological and pathophysiological implications for controlling the neuron excitability and synaptic transmission. A short-term exposure to kainic acid causes externalization of KAR, whereas a longer stimulation by kainate leads to endocytosis and receptor degradation [75, 82]. This bi-directional feedback system demonstrates an elegant mechanism of scaling, which increases the number of KAR on low-active synapses and decreases it on highly active synapses. Low or moderate activation of KAR increases endosomal recycling in the spine through the metabotropic pathway, which involves the activation of protein kinase C, G protein and Rab11 [74].

The metabotropic action of postsynaptic KAR also induces NMDA-independent long-term potentiation. This can occur due to increased activity of voltage-dependent calcium channels, the influx of extracellular Ca	extsuperscript{2+}, which in turn stimulates a calcium release from the calcium depot [83, 84]. It is also known that the metabotropic effect of postsynaptic KAR increases the excitability of neurons by inhibiting hyperpolarization caused by potassium current in pyramidal cells of the CA1 region of the hippocampus [85].

The kainate receptors have a wide functional spectrum of postsynaptic generation of exciting internal currents in the hippocampus [86]. It has been shown that a short-term high-frequency stimulation of mossy fibers causes slow EPSC mediated by postsynaptic high affinity kainate receptors in CA3 neurons [87, 88]. Some researchers suggest that the GluK1 subunit can modulate these EPSC, since a GluK1 antagonist reduced these currents. Similar to the EPSC associated with presynaptic KAR, increased concentrations of kainic acid also led to a decrease in the amplitude of EPSC [85]. EPSC mediated by postsynaptic KAR were found in pyramidal and fast-specific cells in the second, third, and fifth layers of the rat motor cortex [89], thalamic cortical synapses, and pyramidal neurons of the fifth layer of the neocortex [90, 91].

Kainate receptors and the astroglia function

Immunohistochemical studies demonstrated the expression of all KAR subunits (mRNA or protein) in glial cells and their co-expression with subunits of the AMPA receptor [92–94]. The GluK1–3 and GluK5 subunits are present in 50% of astrocytes and 40% of oligodendrocytes. In astrocytes, the subunits are distributed throughout the cell body, whereas in oligodendrocytes they are mainly limited to the soma. It has been reported that the glutamate-induced membrane current in oligodendrocytes is generated totally by AMPA and kainate receptors, both in culture [38, 95, 96] and in brain sections [97, 98]. In addition, the kainate receptors identified in oligodendrocytes can mediate glutamate excitotoxicity [99, 100]. Kainate receptors containing the GluK1 subunits are essential for the transmission of astroglia–neuron signals in the hippocampus, where glutamate released from the astroglia sends a signal to inhibitory neurons by activating the neuronal kainate receptors [101, 102]. This addition of glutamate sensors, expressed in the glial cell membranes, allows the glia to decipher the neuronal activity, synchronize and integrate the neuronal, neuron-glial and glia-glial interactions [103–105].

Kainate receptors and the rhythmic activity of neural networks

It is known that KAR can facilitate the synchronous rhythmic electrical activity. An example of this mechanism in the healthy brain is gamma-oscillations (20–80 Hz) in the hippocampal and neocortical networks, which play an important role in learning and memory. In brain disorders, epileptiform electrographic seizures
(i.e. periodic high-frequency high-amplitude oscillations) can be observed. The induction of rhythmic activity by chemically activating the kainate receptors can generate stable gamma-oscillations [106, 107] independent of NMDAR, mGluRs or AMPAR [108]; kainate injections able to provoke epileptogenic bursts are used to model epileptogenesis in animals [109].

The GluK1 and GluK2 subunits play an important role in the generation of gamma-oscillations and epileptiform bursts; small changes in the overall activity in the hippocampal CA3 area can change the balance between excitation and inhibition, causing the neural network to switch from gamma-oscillations to epileptiform activity [108, 110]. The absence of GluK1 or GluK2 subunits manifests in the phenotypes with gamma-oscillations. Thus, in the absence of GluK1, an increased susceptibility to epileptiform bursts develops whereas GluK2 ablation did not induce either gamma–oscillations or epileptiform bursts. The results suggest that the GluK1 and GluK2 subunits play different roles in the rhythmic activity induced by kainate (see Figure) [110–113].
Kainate receptors differ from glutamate receptors as they have both ionotropic and metabotropic effects. They perform a number of important functions, such as mediation and modulation of synaptic transmission to maintain the balance of excitation and inhibition. The modulatory effect of kainate receptors is mediated by the mechanisms controlling the neuron activity and the function of glial cells. Kainate receptors are commonly expressed throughout the CNS, mainly in neurons, and also in glial cells. The kainate receptors act on both presynaptic and postsynaptic neuron endings, controlling the excitatory and inhibitory synaptic transmission and the glia function.

Presynaptic kainate receptors participate in controlling the release of inhibitory and excitatory neurotransmitters in the concentration-depending biphasic mode; the latter represents a homeostatic mechanism. These receptors act via several pathways to finely tune the neuron activity; these pathways include the non-classical metabotropic effect on synaptic transmission through G-proteins that induces the NMDAR-independent long-term potentiation. In addition, kainate receptors contribute to the morphological plasticity and expression in the dendritic spines of the GluK2-containing receptors. The glial cell receptors are sensitive to glutamate as AMPA and NMDA, the kainate receptors are less well known in the literature but worth further research. Since compared with other glutamate receptors, such as AMPA and NMDA, the kainate receptors are less known in the literature but worth further research. Since the non-classical metabotropic signaling via kainate receptors is involved in the regulation of synaptic transmission, plasticity, learning and memory, it is especially important to study their morphological and functional aspects together with their physiological and pathological roles in the CNS. This will allow us to proceed to the study of a new class of pharmacological targets for the treatment of epilepsy and other CNS diseases. Therefore, studying the mechanisms mediated by the kainate receptors of glutamate is one of the main tasks of neurobiology today.

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