



Published in final edited form as:

Physiol Rev. 2011 July ; 91(3): 1009–1022. doi:10.1152/physrev.00015.2010.

The dynamic modulation of GABA_A receptor trafficking and its role in the formation of inhibitory synapses

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Abstract

γ -aminobutyric acid type-A receptors (GABA_ARs) mediate the majority of fast synaptic inhibition and are the principle sites of action for anxiolytic, sedative, hypnotic and anti-convulsant agents that include benzodiazepines, barbiturates, neurosteroids and some general anesthetics. GABA_ARs are hetero-pentameric ligand-gated ion channels that are found concentrated at inhibitory postsynaptic sites where they mediate phasic inhibition. Specialized populations of extrasynaptic GABA_ARs that mediate tonic inhibition are also expressed by neurons. The efficacy of phasic inhibition and thus neuronal excitability is critically dependent on the accumulation of specific GABA_AR subtypes at inhibitory synapses. Here we evaluate how neurons control the number of GABA_ARs on the neuronal plasma membrane together with their selective stabilization at synaptic sites. We then go on to examine the impact that these processes have on the strength of synaptic inhibition and their possible impact on behavior and the etiology of neuropsychiatric disorders.

I. Introduction

γ -aminobutyric acid (GABA) is the major inhibitory neurotransmitter in the adult mammalian central nervous system (CNS). It is estimated that at least one third of all CNS neurons utilize GABA as their primary neurotransmitter. Most of these GABAergic neurons are interneurons and therefore uniquely able to alter the excitability of local circuits within a given brain region. GABA exerts its powerful inhibitory influence by acting on two distinct classes of receptors based on their electrophysiological and pharmacological properties. Ionotropic GABA_ARs are fast acting ligand-gated chloride channels (Sieghart 2006) while metabotropic GABA_B receptors are coupled indirectly via G proteins to either calcium or potassium channels to produce slow and prolonged inhibitory responses (Bettler and Tiao 2006). GABA_A receptors are clinically relevant drug targets for anti-convulsant, anxiolytic and sedative-hypnotic agents including benzodiazepines, barbiturates, neurosteroids, certain classes of general anesthetics and alcohol. These allosteric compounds bind to discrete sites

distinct from the agonist binding site to either positively or negatively modulate receptor function (Sieghart et al., 2006).

The classic GABA_AR-mediated hyperpolarization of the membrane potential is attributed to the direct activation of an integral ion channel and the resultant influx of chloride along its electrochemical gradient. The concentration-response curve exhibits positive co-operativity consistent with the presence of at least two GABA binding sites on the receptor complex (Baumann et al., 2003). Following brief exposure to high (millimolar) concentrations of GABA (released from presynaptic vesicles) activation of GABA_ARs located at synaptic sites transiently moves the membrane potential away from the spike threshold required for action potential generation underlying what is known as 'phasic' inhibition. In contrast, low (submicromolar) concentrations of ambient GABA in the extracellular space can persistently activate spatially and temporally less restricted extrasynaptic receptors to generate a persistent or 'tonic' inhibitory state (Farrant and Nusser 2005). Given the critical role of GABA_ARs in controlling neuronal excitability and animal behavior, it is of fundamental importance to understand the mechanisms used by neurons to regulate their function.

Emerging evidence indicates that activity-dependent changes in the number of postsynaptic GABA_ARs represent one of the most powerful mechanisms underlying the functional plasticity of GABAergic synapses. Moreover, deficits in the functional expression of GABA_ARs have been implicated in the pathogenesis of a wide range of neurological and psychiatric diseases including epilepsy, anxiety, depression, schizophrenia and substance abuse. A significant amount of research has thus focused on detailing the cellular and molecular mechanisms that regulate the stability of GABA_ARs on the cell surface and their accumulation at the inhibitory postsynapse.

II. GABA_AR structure

GABA_ARs belong to the superfamily of Cys-loop ligand-gated ion channels that comprises nicotinic acetylcholine (nACh) receptors, strychnine-sensitive glycine receptors and 5-hydroxytryptamine type-3 (5-HT₃) receptors. Members of this family are heteropentameric glycoproteins composed of homologous subunits that specifically recognize one another and assemble around an intrinsic ion channel, which in the case of the GABA_AR is permeable to chloride and, to a lesser extent, bicarbonate anions (Unwin, 1993). Each subunit is predicted to share a common membrane topology encompassing a large extracellular N-terminal ligand-binding domain and four transmembrane (TM)1–4 domains. A major cytoplasmic loop lies between TM3 and TM4 that is the most divergent part of the sequence among the GABA_AR subfamily. This intracellular domain is the target for a number of post-translational modifications that can directly affect channel function and/or mediate the interaction with cytosolic proteins important for regulating receptor localization and/or membrane trafficking.

Comparative modeling of the GABA_AR based on the 4-Å resolution model of the *Torpedo* nACh receptor for the first time provided an insight into the 3D organization of this ligand-gated ion channel (Ernst et al., 2005). The extracellular domain of each receptor subunit comprises a variable N-terminal domain and two β-folded sheets that form a twisted

sandwich connected by a signature disulphide bridge. Each subunit has a principal (+) and a complementary (–) side. The GABA binding sites are located in solvent-accessible pockets at the two $\beta+\alpha-$ subunit interfaces and that of benzodiazepines at the $\alpha+\gamma-$ interface (Ernst et al., 2003). The TM domain of the receptor is made up of four loosely-packed helical bundles resulting in a considerable amount of solvent-accessible space within subunits and at the subunit interfaces. It has been proposed that the intersubunit pockets form a continuous groove with their extracellular counterparts, suggesting that they may not only play a role in the conformational mobility of the receptor but may also provide drug-binding sites.

The intrasubunit pockets contain a number of amino acid residues that have been shown to be of key importance in the binding and/or efficacy of a number of modulatory drugs. For example, a pocket within the $\alpha 1$ subunit, defined by the presence of S270 in TM2 and A291 in TM3, is thought to correspond to the site of action of volatile anesthetics (Nishikawa et al., 2002). Similarly, the presence of a homologous serine residue (S265) in the intrasubunit pocket of $\beta 1$, which is replaced by an aspartic residue in the $\beta 2$ and $\beta 3$ subunit isoforms, is believed to account for the subtype-selectivity of etomidate and other related substances (Belelli et al., 1997). Future experiments leading to the improvement in the accuracy of the models will not only continue to provide a new perspective on existing data but also pave the way for structure-based drug design.

III. Subunit composition of synaptic and extrasynaptic GABA_A receptors

To date, 21 GABA_AR subunits have been cloned and sequenced from the mammalian CNS. These have been divided into eight classes on the basis of sequence identity (Whiting et al., 1999): $\alpha(1-6)$, $\beta(1-3)$, $\gamma(1-3)$, δ , $\epsilon(1-3)$, π , θ and $\rho(1-3)$. Subunit isoforms within a single class share approximately 70% sequence identity but between classes this falls to 30–40%. Alternatively spliced variants of several of these subunits have been reported, generating further subunit diversity and the potential for extensive molecular heterogeneity (Sieghart et al., 1999). For example, the $\gamma 2$ subunit exists in short ($\gamma 2S$) and long ($\gamma 2L$) forms, which differ in an eight amino acid insert in the intracellular domain of the $\gamma 2L$ subunit (Whiting et al., 1990; Kofuji et al., 1991). There is a plethora of GABA_AR subunit isoforms; however, due to the selective oligomerization mediated by receptor assembly, only a limited number of subunit combinations are in fact expressed on the neuronal plasma membrane (Sieghart and Sperk, 2002). The majority of GABA_AR subtypes in the brain are composed of $\alpha 1\beta 2\gamma 2$, followed by $\alpha 2\beta 3\gamma 2$ and $\alpha 3\beta 3\gamma 2$ with a likely stoichiometry of $2\alpha:2\beta:1\gamma$, similar to that found for nACh receptors (Chang et al., 1996; Farrar et al., 1999; Knight et al., 2000; Tretter et al., 1997). To a lesser extent, δ , ϵ , π subunits replace the γ subunit to form benzodiazepine-insensitive receptor subtypes, whereas the θ subunit has been shown to replace the β subunit (Sieghart and Sperk, 2002). Conversely, ρ subunits (which are predominantly expressed in the retina) rarely co-assemble with other GABA_AR subunits. Instead they homo- and hetero-oligomerize with other ρ subunits to form a family of pharmacologically distinct GABA-gated chloride channels that have been termed GABA_C receptors (Bormann 2000).

In addition to molecular determination of GABA_AR assembly, subunit composition is further restricted by the spatial and temporal pattern of subunit expression. *In situ* hybridization and immunohistochemical studies have demonstrated that each one of the subunits has a distinct regional and cellular distribution within the brain. Different subunits and/or subunit combinations further dictate the subcellular localization of these ligand-gated chloride channels and determine their biophysical and pharmacological properties. Detailed knowledge of the molecular composition and the exact anatomical expression of different GABA_AR subtypes are therefore crucial in understanding the physiological actions of GABA within the brain and for developing potentially clinically useful subtype-selective drugs that are devoid of the side effects associated with the classical benzodiazepines.

In the cerebellum the $\gamma 2$ subunit has been shown to be a component of all postsynaptic GABA_ARs, whereas the δ subunit is found almost exclusively at extrasynaptic sites (Nusser et al., 1998b). GABA_ARs incorporating a $\gamma 2$ subunit together with $\alpha 1-3$ subunits ($\alpha 1-3\beta 2/3\gamma 2$) are thus the predominant receptor subtypes responsible for mediating phasic inhibition. It is important to note, however, that these receptor subtypes are also abundant at perisynaptic and extrasynaptic sites. This is consistent with recent evidence for the dynamic mobility and rapid exchange of $\gamma 2$ subunit-containing receptors between extrasynaptic and synaptic receptor pools (Jacob et al., 2005; Thomas et al., 2005). The $\alpha 5$ subunit is unique in that, despite its coassembly with a $\gamma 2$ subunit ($\alpha 5\beta 3\gamma 2$), it is enriched at extrasynaptic sites (Brunig et al., 2002). In accordance with this finding, deletion of the $\alpha 5$ subunit eliminates tonic conductance in cultured hippocampal neurons (Caraiscos et al., 2004). However, there is a pool of $\alpha 5$ subunit-containing GABA_ARs that concentrates at GABAergic synapses on CA1 pyramidal cell dendrites (Serwanski et al., 2006). In a recent study, diazepam-sensitive IPSCs elicited by dendrite preferring interneurons in the rat neocortex were blocked by the $\alpha 5$ subtype-selective inverse agonist ($\alpha 5$ IA), suggesting that $\alpha 5\beta \gamma 2$ receptor subtypes may also play a role in phasic inhibition (Ali and Thomson, 2008). $\alpha 4$ and $\alpha 6$ subunits form assembled channel complexes with δ subunits ($\alpha 4\beta \delta$ and $\alpha 6\beta \delta$) that are exclusively extrasynaptic, accounting for tonic conductance in the thalamus and cerebellum respectively (Nusser et al., 1998b). On the other hand, the majority of phasic signaling in the cerebellum is due largely to synaptic $\alpha 6$ subunit-containing receptors of the $\alpha 6\beta 2\gamma 2$ combination (Nusser et al., 1998b).

IV. Functional significance of GABA_AR structural heterogeneity

While the precise subcellular localization of different GABA_AR subtypes undoubtedly contributes to their participation in phasic and tonic forms of signaling, this distinction alone is not sufficient to account for their differential activation. Subunit composition is a major determinant of the binding and gating properties of the ion channels, and hence the magnitude of the response following exposure to ligand. Studies using recombinant GABA_ARs have revealed that sensitivity to GABA is defined by the type of α subunit present. Extrasynaptic $\alpha 6\beta 3\delta$ and $\alpha 4\beta 3\delta$ subunit compositions display the highest affinities for GABA and synaptic $\alpha 3\beta 3\gamma 2$ subtypes the lowest (Bohme et al., 2004). For both $\alpha \beta \gamma$ and $\alpha \beta \delta$ assemblies, the identity of the α subunit also affects the rates of activation, deactivation and desensitization (Bianchi et al., 2002; Caraiscos et al., 2004; Gingrich et al., 1995; Lavoie et al., 1997; McClellan and Twyman, 1999; Tia et al., 1996). Conversely, in $\alpha \beta 3\gamma 2$

subunit-containing receptors replacing the $\gamma 2$ subunit with a δ subunit results in a dramatic reduction in single channel conductance independent of the type of α subunit present (Fisher and Macdonald, 1995). This supports the notion that GABA has a high affinity but low efficacy at δ subunit-containing extrasynaptic receptors. The presence of a $\gamma 2$ or δ subunit also influences channel kinetics: $\alpha\beta\delta$ receptors desensitize more slowly and less extensively than $\alpha\beta\gamma$ receptors (Bianchi and Macdonald, 2002; Haas and Macdonald, 1999; Saxena and Macdonald, 1996). Together the distinct biophysical properties of γ and δ subunit-containing receptors are wholly consistent with the involvement of these receptor subtypes in phasic and tonic signaling respectively.

Differences in subunit composition between synaptic and extrasynaptic receptors are reflected in a differential modulation of phasic and tonic signaling by a number of compounds of therapeutic importance. The most frequently cited example of this is the role of the α subunit in defining receptor affinity for benzodiazepines. GABA_ARs incorporating either an $\alpha 4$ or $\alpha 6$ subunit renders receptors insensitivity to functional modulation by benzodiazepines (Benson et al., 1998) as does elimination or substitution of the $\gamma 2$ subunit. This difference can be attributed solely to the presence of a conserved arginine residue in $\alpha 4$ and $\alpha 6$ subunits, which in $\alpha 1-3$ and $\alpha 5$ subunits is a histidine (Wieland et al., 1992). Thus benzodiazepine site ligands selective for $\alpha 1-3$ subunits largely influence phasic signaling, whereas those selective for $\alpha 5$ subunits are capable of primarily modulating tonic conductance.

The diverse CNS depressant effects of benzodiazepines have been attributed to specific α subunit types of GABA_ARs (Rudolph and Möhler, 2004). A combined molecular genetic and pharmacologic approach has revealed that $\alpha 1$ subunit-containing receptors mediate the sedative, amnesic and, in part, anticonvulsant actions of diazepam, whereas $\alpha 2$ subunits contribute to its anxiolytic and muscle-relaxant effects. Strong pharmacological evidence for the involvement of the $\alpha 3$ subunit in anxiety comes from a study by Atack and colleagues (2005) who showed that the $\alpha 3$ subtype-selective inverse agonist ($\alpha 3$ IA) induces an anxiogenic response in rats. The $\alpha 5$ subunit has been implicated in learning and memory following the observation that a single point mutation (H105R), which prevents the interaction of this receptor subtype with diazepam, abolishes its memory impairing effects (Crestani et al., 2002). The development of tolerance to the sedative actions of chronic diazepam is also associated with a downregulation of $\alpha 5$ subunit-containing GABA_ARs to which $\alpha 5$ -H105R mice are resistant (Rijnsoever et al., 2004).

The diverse functions of GABA in the CNS are matched not just by the heterogeneity of GABA_ARs but also by the complex trafficking and clustering mechanisms that generate and maintain surface receptor populations accessible to the neurotransmitter at inhibitory postsynaptic specializations or extrasynaptic sites, depending on their subunit composition. The regulation of these mechanisms by GABA_Areceptor-associated proteins is discussed below.

V. Membrane trafficking of GABA_ARs

GABA_ARs are not static entities on the neuronal cell surface but are believed to cycle continuously between the plasma membrane and intracellular compartments. The relative rates of receptor exo- and endocytosis are therefore key determinants in controlling the size of the postsynaptic pool accessible to GABA and GABAergic compounds and thus the strength of synaptic inhibition. Importantly, GABA_ARs have recently been reported to be inserted into and removed from the plasma membrane exclusively at extrasynaptic sites (Bogdanov et al., 2006; Thomas et al., 2005), highlighting the importance of lateral diffusion for their postsynaptic specialization.

VI. Exocytosis of GABA_ARs to the plasma membrane

GABA_ARs can be delivered to the cell surface either as newly assembled channel complexes via a *de novo* secretory pathway or reinserted following internalization. The oligomerization of GABA_AR subunits into channel complexes is believed to occur in the endoplasmic reticulum (ER). Evidence suggests that this assembly process plays a critical role in determining the diversity of receptor subtypes expressed on the neuronal plasma membrane. Proteins cannot exit the ER until they have achieved their correctly folded conformation, and misfolded or unassembled proteins are retrotranslocated from this organelle for degradation in the proteasome, restricting the number of subunit combinations that can access the cell surface (Kittler et al., 2002). Following correct assembly, GABA_ARs are trafficked to the Golgi apparatus and segregated into vesicles for transport to and insertion into the plasma membrane facilitated by a number of receptor-associated proteins.

Yeast two-hybrid screens using the $\gamma 2$ subunit ICD as bait isolated the first known GABA_AR associated protein (GABARAP) (Wang et al., 1999), a 17 kDa cytosolic protein belonging to the family of membrane associated proteins (MAPs) involved in membrane trafficking. This includes the estrogen-induced protein first isolated in guinea-pig endometrial cells (GEC)-1 (or GABRAP like-1), Golgi-associated transport enhancer of 16 kDa (GATE-16, or GABARAP like-2), Apg8L and light chain (LC)-3 subunits of MAP1. Interestingly, GABARAP knockout mice do not show differences in punctate staining of $\gamma 2$ subunit-containing GABA_ARs and lack an overt behavioral phenotype (O'Sullivan et al., 2005). GABARAP may thus be functionally redundant. Nevertheless, the notion that GABARAP is involved in the trafficking of GABA_ARs to the plasma membrane is supported by the finding that overexpression of GABARAP in heterologous expression systems and in cultured hippocampal neurons results in an increase in the cell surface expression of $\gamma 2$ subunit-containing GABA_ARs (Leil et al, 2004). In *Xenopus* oocytes expressing $\alpha 1\beta 2\gamma 2$ subunit-containing GABA_ARs this was accompanied by an increase in GABA_AR-mediated synaptic inhibition, an effect requiring the $\gamma 2$ subunit- and microtubule-binding motifs as well as intact polymerized microtubules (Chen et al., 2005). More recently it has been demonstrated that GABARAP-mediated exocytosis of GABA_ARs is necessary for potentiation of inhibitory transmission by NMDA receptor activation, suggesting that GABARAP might have a role in the regulated delivery of GABA_ARs to the plasma membrane after activity rather than in the maintenance of basal surface receptor levels (Marsden et al., 2007).

Evidence in support of a function of GABARAP as a trafficking factor includes the identification of phospholipase C (PLC)-related catalytically inactive protein (PRIP)-1, a 130 kDa protein that is believed to competitively inhibit the binding of the $\gamma 2$ subunit of GABA_ARs to GABARAP (Kanematsu et al., 2002). PRIP1 knockout mice show impairments in GABA_AR modulation by benzodiazepines and zinc-sensitivity, indicating reduced activation of $\gamma 2$ subunit-containing receptors (Kanematsu et al., 2002). These findings suggest that PRIP1 may play a role in the regulation of GABA_AR trafficking by GABARAP, ensuring that only mature $\alpha\beta\gamma$ receptor complexes are delivered to the plasma membrane. In addition, PRIP1 has been shown to directly bind to the ICD of GABA_AR $\beta 1-3$ subunits, serving as an adaptor protein for the protein phosphatase (PP)-1 α , and as such has been implicated in the phospho-dependent modulation of GABA_AR functional expression (Terunuma et al., 2004). Recently, a second PRIP isoform (PRIP2) has been identified that, like PRIP1, binds both GABARAP and PP1 α , suggesting a central role for all PRIP isoforms in modulating GABA_AR functional expression (Uji et al., 2002).

GABA_ARs also interact with the protein that links the integrin-associated protein with the cytoskeleton (Plic)-1. Plic1 is a 67 kDa protein with a ubiquitin-like (UBL) N-terminal domain and a ubiquitin-associated (UBA) C-terminal domain (Kleijnen et al., 2000). It is able to bind ubiquitin ligases and components of the proteasome. As such it is thought to interfere with ubiquitin-dependent proteolysis of proteins (Kleijnen et al., 2000). Yeast two-hybrid screens and glutathione S-transferase (GST) affinity purification assays have shown that Plic1 interacts with the ICD of $\alpha 1-3,6$ and $\beta 1-3$ (but not $\gamma 2$ or δ) subunits of GABA_ARs (Bedford et al., 2001), indicating that Plic-1 function might be relevant for the majority of receptor subtypes expressed in the brain. This interaction has been demonstrated to be of significance in mediating the functional expression of GABA_ARs in human embryonic kidney (HEK-293) cells and in hippocampal slices as revealed using dominant negative peptides (Bedford et al., 2001). In a recent study by Saliba and colleagues (2008), the authors showed that Plic1 increases the accumulation of GABA_AR $\beta 3$ subunits on the cell surface in a manner independent of their rates of internalization. These findings suggest that Plic1 selectively modulates the secretory pathway. In accordance with this, Plic1 was found to significantly increase the half-life of polyubiquitinated GABA_AR $\beta 3$ subunits in the ER and facilitate their insertion into the plasma membrane (Saliba et al., 2008). By increasing the residence times of unassembled subunits in the ER, Plic1 may also increase subunit maturation and production of heteromeric receptors. Plic1 regulation of the ubiquitin-dependent proteasomal degradation of GABA_ARs may thus provide a dynamic mechanism for regulating the efficacy of inhibitory synaptic transmission.

The 190 kDa brefeldin A-inhibited guanine nucleotide exchange factor (BIG)-2 was also identified as a GABA_AR β subunit interacting protein in a yeast two-hybrid screen (Charych et al., 2004). BIG2 is concentrated mainly in the trans-Golgi network but is also found in vesicle-like structures along dendrites and at the postsynaptic plasma membrane (Charych et al., 2004). Interestingly, coexpression of BIG2 with the GABA_AR $\beta 3$ subunit results in an increase in $\beta 3$ exit from the ER, suggesting that BIG2 is involved in the post-Golgi vesicular trafficking of GABA_ARs (Charych et al., 2004). It is currently proposed that BIG2 may play a role in the transport of newly assembled GABA_ARs to the postsynaptic plasma membrane and also be involved in receptor recycling (Shen et al., 2006; Shin et al., 2004). Other

GABA_AR-associated proteins implicated in the forward trafficking of GABA_ARs include the GABA_AR interacting factor (GRIF)-1 (Beck et al., 2002) and the multifunctional protein gC1qR (Schaerer et al., 2001). However, their functional significance remains unclear.

VII. Endocytosis of GABA_ARs from the plasma membrane

GABA_ARs have been shown to be localized in clathrin-coated pits, suggesting that they undergo clathrin-mediated endocytosis, a process that is further dependent on dynamin for endocytic vesicle formation. The clathrin adaptor protein (AP)-2 is a central component in the formation of these vesicles, forging a link between membrane proteins and clathrin, which forms the outer layer of the coat. AP2 is a heterotetrameric complex composed of two large (~100 kDa) α and β 2 subunits, a medium (50 kDa) μ 2 subunit, and a small (19 kDa) σ 2 subunit, commonly referred to as adaptins. The α adaptin is responsible for targeting the protein to the plasma membrane, where the β 2 adaptin interacts with clathrin to trigger clathrin assembly, forming coated pits. This in turn leads to the activation of μ 2 adaptin phosphorylation, inducing a conformational change in the subunit that allows the complex to directly bind to endocytic motifs in cell surface receptors, clustering the protein cargo into the assembling coated pit. Cargo is bound by the β 2 and μ 2 adaptins that mostly recognize a dileucine motif or the canonical tyrosine-based YXX Φ motif (where X denotes any amino acid and Φ a bulky hydrophobic residue) (Clague, 1998; Le Borgne and Hoflack, 1998). The σ 2 adaptin is responsible for stabilizing the complex at the core by mediating subunit interactions. The AP2 complex also interacts with several accessory proteins that form essential components of the endocytic machinery (Slepnev and De Camilli, 2000).

GABA_ARs are intimately associated with AP2 in the brain through a direct binding of the β 1–3 and γ 2 GABA_AR subunits (Kittler et al., 2000). In the GABA_AR β 2 subunit a dileucine AP2- β 2 adaptin-binding motif (L³⁴³L³⁴⁴) has been identified. This motif is important for clathrin-mediated endocytosis in HEK-293 cells and in cortical slices (Herring et al., 2003, 2005). It is also present in the ICDs of receptor β 1 and β 3 subunits, but evidence suggests that it is not involved in the interaction of these subunits with the AP2 complex (Kittler et al., 2005). In addition, an atypical AP2-binding motif conserved within the ICD of all GABA_AR β subunit isoforms has been identified (KTHLRRRSSQLK in the β 3 subunit) (Kittler et al., 2005). This motif, which is enriched in lysine and arginine residues, also incorporates the major sites of phosphorylation for PKA and PKC within this class of receptor subunits: S409 in β 1, S410 in β 2 and S408/9 in β 3 (Moss et al., 1992). *In vitro* binding experiments have shown that phosphorylation and/or mutation of these residues confers a reduction in binding of the GABA_AR β subunits to the μ 2 adaptin of the AP2 complex (Jacob et al., 2009; Kittler et al., 2005). Moreover, neurons expressing GABA_ARs incorporating fluorescent β 3^{S408/9A} subunits exhibit reduced endocytosis and enhanced functional expression at both synaptic and extrasynaptic sites (Jacob et al., 2009). Intriguingly, while analyzing the cell surface distributions of wild-type and S408/9A mutant β 3 subunit-containing GABA_ARs, the authors noted an apparent increase in long, filopodia-like spines on neurons expressing the latter. This deficit in spine maturity was reversed by pharmacological blockade of GABA_ARs. Regulating the efficacy of synaptic inhibition by modulating GABA_AR membrane trafficking may thus play a critical role in regulating spinogenesis and synaptic plasticity (Jacob et al., 2009).

More recently a tyrosine based AP2- μ 2 adaptin-binding motif in the GABA_AR γ 2 subunit (Y³⁶⁵GY³⁶⁷ECL) has been identified that is also conserved in the γ 1 and γ 3 subunits (Kittler et al., 2008). These tyrosine residues are the principal sites for phosphorylation by Fyn and Src kinase (Jurd et al; 2010; Brandon et al., 2001; Moss et al., 1995). Utilizing nonphosphorylated and phosphorylated peptides corresponding to Y365 and Y367 the authors showed that this high affinity interaction is phospho-dependent (Kittler et al., 2008). Introduction of the nonphosphorylated γ 2 peptide into neurons induced a large increase in the mIPSC amplitude that was accompanied by an increase in the number of receptors on the cell surface (Kittler et al., 2008).

The physiological significance of GABA_AR endocytosis has been further evaluated by creating a knock-in mouse in which Y365 and Y367 in the γ 2 subunit have been mutated to phenylalanine mutations, significantly increasing the affinity for μ 2-AP2. Homozygotes for these mutations die *in utero*. Heterozygotes are viable and do not exhibit any gross behavioral deficits. Consistent with the roles of Y365/7F in mediating high affinity binding to μ 2-AP2 Y365/7F knock-in mice have elevated total and cell surface expression levels. In the hippocampus this results in an increase in the size of inhibitory synapses and efficacy of synaptic inhibition that is specific to CA3. This increase in inhibition correlates with a specific deficit in spatial memory in Y365/7F mice, a behavioral paradigm dependent upon CA3. Modifying GABA_AR endocytosis clearly has behavioral consequences.

VIII. Post-endocytic GABA_AR sorting

The fate of internalized receptors is another determinant of surface receptor levels. Following internalization GABA_ARs are either rapidly recycled back to the neuronal plasma membrane or, over longer time frames, are targeted for lysosomal degradation, an endocytic sorting decision that is regulated by the Huntingtin-associated protein (HAP)-1. Yeast two-hybrid screens have revealed that HAP1 interacts with the GABA_AR β 1 subunit but not the α 1, γ 2, or δ subunits (Kittler et al., 2004). Overexpression of HAP1 in cultured neurons has been shown to increase the number of receptors on the cell surface at steady-state, an increase that correlates with a dramatic increase in the mean amplitude of mIPSCs but that does not affect the frequency or kinetics of these events (Kittler et al., 2004). However, whether HAP1 promotes GABA_AR recycling or prevents receptor lysosomal degradation is an unresolved issue. Nevertheless, the importance of HAP1-dependent regulation of GABA_AR trafficking is clear, as evidenced by a study in which selective suppression of hypothalamic HAP1 by siRNA induced a decrease in feeding behavior in mice that was attributed to reduced surface expression and activity of GABA_ARs (Sheng et al., 2006). More recently it was established that HAP1 plays a role in regulating the targeting of GABA_ARs to inhibitory synapses dependent on the molecular motor-KIF5 (Twelvetrees et al., 2010).

IX. Postsynaptic targeting and clustering of GABA_ARs

The highly selective subcellular localization of GABA_AR subtypes implies that subunit composition plays a major role in the postsynaptic targeting and clustering of these receptors. While the exact molecular mechanisms that govern the accumulation of

GABA_ARs at inhibitory synapses are not yet fully understood, it involves a number of receptor-associated proteins and cytoskeletal elements that are concentrated at postsynaptic densities (PSDs).

The inhibitory synaptic marker gephyrin is a 93 kDa subsynaptic scaffolding protein that was originally implicated in regulating the postsynaptic clustering of glycine receptors in the spinal cord by directly binding to the receptor β subunit (Meyer et al., 1995). More recently, data derived from gephyrin knockout mice and knockdown experiments using antisense oligonucleotides or shRNAi have revealed that reducing gephyrin expression also leads to an extensive loss in the punctate staining of GABA_ARs incorporating a $\gamma 2$ subunit together with $\alpha 2$ (but not $\alpha 1$) subunits (Essrich et al., 1998; Jacob et al., 2005; Kneussel et al., 1999; Levi et al., 2004). This suggests the existence of both gephyrin-dependent and -independent GABA_AR clustering mechanisms. Gephyrin clusters are absent in $\gamma 2$ subunit knockout mice. The remaining α and β subunit-containing receptors show diffuse staining (Essrich et al., 1998). Furthermore, transfection of $\gamma 2$ subunit-deficient neurons with chimeric $\alpha 2/\gamma 2$ constructs revealed that the TM4 of the $\gamma 2$ subunit is sufficient for targeting these receptors to postsynaptic sites, but that both the TM4 and ICD of the $\gamma 2$ subunit were necessary for recruiting gephyrin to the synapse and rescuing GABAergic inhibitory synaptic function (Allred et al., 2005). A role for this receptor-associated protein in stabilizing previously clustered GABA_ARs at the cell surface, rather than their specialization at inhibitory synapses, has thus been proposed. Concurrent with this, postsynaptic GABA_ARs have been shown to be three times more mobile in cells where gephyrin expression has been impaired as measured by fluorescence recovery after photobleaching (Jacob et al., 2005). It is emerging that the rate of lateral mobility of GABA_ARs plays a central role in determining the number of these receptors at synaptic sites, a process revealed by the use of single particle tracking. This approach has illustrated that there is rapid exchange between extrasynaptic and synaptic populations of GABA_ARs and that this process is subject to modulation via the activity of protein phosphatase IIB. This Ca⁺-dependent mechanism activated via NMDA receptors leads to an increase in the lateral mobility of GABA_ARs and a reduction in the size of inhibitory synapses, a process that favors neuronal depolarization (Bannai et al., 2009).

Gephyrin is believed to anchor postsynaptic GABA_ARs to the plasma membrane by cross-linking the receptor molecule to the tubulin and actin cytoskeleton. Gephyrin binds with high affinity to tubulin and as such is viewed as a *bona fide* microtubule binding protein MAP (Ramming et al., 2000). Gephyrin also mediates an indirect interaction with the cytoskeleton by binding to LC1 and LC2 of the dynein motor proteins and the microfilament-associated proteins belonging to the Mena/VASP family (Fuhrmann et al., 2002). The membrane-associated protein collybistin II is a guanine-nucleotide exchange factor (GEF) that binds to and so permits the subsynaptic localization of the otherwise intracellular gephyrin, where it directly interacts with and immobilizes specific GABA_AR subtypes on the plasma membrane (Harvey et al., 2004; Kins et al., 2000). In addition, the dystrophin glycoprotein complex (DGC) and neuroligin (NL)-2 bridge the synaptic cleft by interacting with presynaptic β -neurexin, promoting GABAergic synaptogenesis (Tretter and Moss, 2008).

Although gephyrin is found to be concentrated at GABAergic synapses in both hippocampal and cortical neurons (where glycine receptor expression is relatively low), in initial studies using standard biochemical methods it failed to co-purify with any of the GABA_AR subunits. Hence an intermediate linker protein was postulated. Given its ability to directly bind the $\gamma 2$ subunit ICD and gephyrin the prime candidate was GABARAP. However, subsequent analysis revealed that GABARAP is not clustered at gephyrin-rich postsynaptic specializations but instead is mainly localized in intracellular compartments including the ER, Golgi apparatus and, to some extent, in subsynaptic secretory vesicles (Kittler et al., 2001; Wang et al., 1999). This is consistent with a role for this protein in intracellular trafficking rather than postsynaptic clustering.

In a recent study Tretter and colleagues (2008) for the first time provided evidence that gephyrin binds directly to the $\alpha 2$ subunit ICD *in vitro*. This interaction was blocked by low concentrations of detergent, providing a possible explanation as to why previous attempts to identify a direct association between gephyrin and GABA_ARs were unsuccessful. However, under the same conditions only very weak binding to $\beta 3$ and $\gamma 2$ subunits was observed (Tretter et al., 2008), suggesting that α subunits isoforms may play a predominant role in the synaptic accumulation of GABA_ARs via their ability to directly bind gephyrin

The 81 kDa actin-binding protein radixin, a member of the ezrin/radixin/moesin protein family, has recently been shown to directly link the $\alpha 5$ subunit to the cytoskeleton via a radixin-binding motif conserved within the ICD of $\alpha 1-3$ and $\alpha 5$ subunits, differing only by two amino acids in the $\alpha 2$ subunit (Loebrich et al., 2006). In neurons, both depletion of radixin and replacement of the radixin actin-binding motif dramatically decreased $\alpha 5$ subunit-containing GABA_AR clusters without altering total surface expression levels. Radixin also showed limited colocalization with postsynaptic gephyrin, consistent with previous reports that GABA_AR $\alpha 5$ subunits localize mainly at extrasynaptic sites (Brunig et al., 2002; Loebrich et al., 2006). However, the mechanisms responsible for the formation of radixin-dependent extrasynaptic GABA_AR clusters remain to be elucidated.

X. Modulation of GABA_AR function by post-translational modifications

The cell surface stability of GABA_ARs is further regulated by post-translational modifications such as palmitoylation, ubiquitination and phosphorylation. These modifications have been implicated in altering the biophysical and pharmacological properties of these ligand-gated ion channels (Arancibia-Carcamo and Moss, 2009).

A. Palmitoylation

Palmitoylation is the covalent attachment of the saturated fatty acid palmitate to cysteine residues of a given protein by the palmitoyl transferase Golgi-specific DHHC zinc finger domain protein (GODZ). Palmitoylation enhances the hydrophobicity of proteins and contributes to their membrane association. As such it has been shown to be involved in the postsynaptic clustering and subcellular trafficking of a number of proteins, including AMPA receptors (Hayashi et al., 2005) and the neuronal scaffold proteins PSD-95 and glutamate receptor-interacting protein (GRIP)-1 (DeSouza et al., 2002; Smotrys and Linder, 2004). It is unique in that it is the only reversible lipid modification and thus allows the cell to

dynamically regulate the location of specific proteins. In a yeast two-hybrid screen, the 34 kDa protein GODZ was identified as a GABA_AR γ 2 subunit interacting protein that recognizes a 14 amino acid cysteine-rich domain conserved in the ICD of all γ 1–3 subunits, N-terminal to the GABARAP binding site (Rathenberg et al., 2004). Analysis of Cys-Ala mutant γ 2 constructs in transfected COS-7 cells revealed that the γ 2 subunit is palmitoylated at all four cysteines within the GODZ binding domain (Rathenberg et al., 2004). Mutation of these cysteine residues resulted in a loss of GABA_AR clusters at the cell surface, as did drug-induced global inhibition of palmitoylation by Br-palmitate (Rathenberg et al., 2004). Likewise, disrupting GODZ function or expression levels using dominant negative or RNAi approaches resulted in a significant reduction in the amplitude of mIPSCs attributed to a decrease in postsynaptic GABA_AR number (Fang et al., 2006). From these studies it is evident that palmitoylation can dynamically regulate the efficacy of neuronal inhibition by controlling the accumulation of GABA_ARs at the postsynaptic membrane, although the exact mechanism by which this is achieved is unknown.

B. Ubiquitination

The regulation of GABA_AR trafficking by the ubiquitin-related protein Plic1 suggests that GABA_ARs may also be direct targets for modification by the polypeptide ubiquitin. The covalent attachment of one or more copies of the 76-amino acid ubiquitin monomer to lysine residues of target proteins is referred to as ubiquitination. Monoubiquitination is reversible and serves as an active signal in diverse intracellular trafficking pathways, including as a trigger for endocytosis. In contrast, polyubiquitination is required for the translocation of proteins from the ER back into the cytosol, where they are degraded by the proteasome. Activity-dependent polyubiquitination of GABA_AR β 3 subunits has been shown to reduce the stability of newly-translated and assembled receptors in the ER via a mechanism dependent on the activity of the proteasome (Saliba et al., 2007).

Coincident with a loss of cell surface expression levels, chronic blockade of neuronal activity by tetrodotoxin (TTX) treatment reduced both the amplitude and frequency of mIPSCs (Saliba et al., 2007). TTX had no effect on the enhanced functional expression of GABA_ARs incorporating β 3 subunits in which all twelve lysine residues within the ICD of this subunit had been mutated to arginines (β 3^{K12R}). These mutations did not alter GABA_AR cell surface half-life or internalization rates but did significantly enhance receptor insertion into the plasma membrane (Saliba et al., 2007).

C. Phosphorylation

Protein phosphorylation is fundamental to the activity of cellular signaling networks. It is achieved through protein kinase catalyzed transfer of a phosphate group to serine, threonine and/or tyrosine residues of a given protein substrate that can be reversed in a dephosphorylation reaction catalyzed by protein phosphatases. GABA_ARs are well-established phosphoproteins. Diverse studies on GABA_AR phosphorylation have implicated this process in altering channel gating, conductance and/or kinetics, sensitivity of the receptors to pharmacological agents, protein-protein interactions and membrane trafficking (Vithlani and Moss, 2009). Hence the coordinated activity of kinases and phosphatases plays a pivotal role in controlling neuronal excitability.

Studies in heterologous expression systems have demonstrated that GABA_AR function, depending on the subtype analyzed, can be differentially modulated by phosphorylation of key residues within the ICDs of receptor β 1–3 and γ 2 subunits by a number of kinases, including cAMP-dependent protein kinase (PKA), calcium/phospholipid-dependent protein kinase (PKC), calcium/calmodulin-dependent kinase II (CaMKII), protein kinase B (PKB - also known as Akt), cGMP-dependent protein kinase (PKG) and tyrosine kinases of the Src family (summarized in Table 1.2). This is best illustrated by the differential modulation of GABA_AR subtypes by PKA, dependent upon the identity of the β subunit. *In vitro* studies using purified bacterially expressed GST fusion proteins combined with site directed mutagenesis revealed that the consensus motif for PKA-induced phosphorylation conserved within the ICDs of GABA_AR β 1–3 subunits is RRRXSQLK, where S is a serine at position 409 in β 1 and β 3 subunits and at position 410 in the β 2 subunit (McDonald and Moss, 1997) and X represents either an alanine residue in β 1 and β 2 subunits or a serine residue at position 408 in the β 3 subunit. However, in contrast to *in vitro* findings, analysis of recombinant GABA_ARs in HEK-293 cells has demonstrated that PKA-induced phosphorylation of the β 3 subunit occurs at both S408 and S409 (rather than at S409 alone), whereas the β 2 subunit is not a substrate for this kinase in heterologous expression systems (McDonald et al., 1998). Interestingly, PKA was shown to depress GABA-activated currents in HEK-293 cells expressing β 1 subunit-containing GABA_ARs, whereas in β 3 subunit-expressing cells a potentiation was observed (McDonald et al., 1998). The latter was attributed to the presence of two juxtaposed serine residues (S408 and S409) in β 3, as selective mutation of S408 to alanine (to structurally resemble the GABA_AR β 1 subunit at this site) converted this potentiation to a depression (McDonald et al., 1998). On the other hand, PKA had no effect on β 2 subunit-expressing cells (McDonald et al., 1998), which can be accounted for by the selective recruitment of PKA to GABA_AR β 1 and β 3, but not β 2, subunits via the A-kinase anchoring protein (AKAP) of 79 (rat)/150 (human) kDa (Brandon et al., 2003).

AKAPs have also been shown to interact directly with PP2B and PKC in addition to PKA (Klauck et al., 1996; Colledge and Scott, 1999); however, the targeting of PP2B and PKC to GABA_ARs by AKAP remains to be investigated. Interestingly, PP2B has been shown to bind directly to the ICD of the γ 2 subunit, thereby dephosphorylating the receptor and inducing long-term depression (LTD) of synaptic inhibition in the CA1 region of the hippocampus, a mechanism that is believed to contribute to learning and memory (Wang et al., 2003). PKC is recruited to the GABA_AR via the anchoring protein receptor for activated C kinase (RACK)-1. Like PKA it has different effects on receptor function depending on subunit composition. In agreement with the findings *in vitro* all of the β subunit isoforms were found to be a substrate for this kinase in HEK-293 cells (Krishek et al., 1994; McDonald et al., 1998; Brandon et al., 2000). Functional studies employing site-directed mutagenesis have demonstrated a downregulation of receptor activity in response to phorbol esters mediated by PKC-induced phosphorylation of β 1 S409, β 2 S410, γ 2S S327 and γ 2L S327/343. However, the use of a constitutively active catalytic domain of PKC (PKM) produced results contradicting those obtained with phorbol esters. In transfected mouse fibroblasts PKM enhanced the response to GABA in a manner that was dependent on the phosphorylation of the β 1 subunit at S409 and the γ 2L subunit at S327 and S343 (Lin et al.,

1996). The reason for this discrepancy remains unresolved but it has been speculated that different PKC isoforms may produce different responses (Song and Messing, 2005). In contrast, brain-derived neurotrophic factor (BDNF)-induced PKC-mediated phosphorylation of the $\beta 3$ subunit has been shown to transiently increase the amplitude of mIPSCs in cultured neurons, an effect paralleled by an increase in GABA_AR cell surface stability (Jovanovic et al., 2004). Furthermore, the initial enhancement observed in receptor activity was followed by a prolonged depression that was attributed to PP2A-mediated dephosphorylation of the $\beta 3$ subunit (Jovanovic et al., 2004).

No adaptor protein is known for CaMKII, although the kinase is capable of coimmunoprecipitating with GABA_ARs from detergent-soluble mouse brain extracts (McAinsh et al., unpublished data). *In vitro* studies have revealed that the serine/threonine kinase CaMKII directly phosphorylates specific residues within the ICDs of GABA_AR β and $\gamma 2$ subunits. These include S384/409 in $\beta 1$, S410 in $\beta 2$ and S383/409 in $\beta 3$, as well as the $\gamma 2$ subunit at S343 ($\gamma 2L$ only), S348 and T350. Interestingly, the intracellular application of purified, active CaMKII failed to modulate the function of GABA_ARs heterologously expressed in HEK-293 cells but significantly potentiated the amplitudes of whole-cell GABA-activated currents recorded from rat cultured cerebellar granule neurons and from recombinant GABA_ARs expressed in neuroblastoma-glioma hybrid (NG108-15) cells. These findings imply the contribution of some essential neuronal factor (Houston and Smart, 2006).

In addition to binding PKC and the GABA_AR β subunits RACK1 also binds the tyrosine kinase Src to facilitate the phosphorylation of the $\gamma 2$ subunit on Y365 and Y367 (Brandon et al., 2001; Kittler and Moss, 2003; Moss et al., 1995). Coexpression of Src with recombinant $\alpha 1\beta 1\gamma 2$ subunit-containing GABA_ARs has been shown to increase whole-cell GABA-activated currents in HEK-293 cells, an effect abolished by site-specific mutagenesis of both of these tyrosine residues to phenylalanines (Moss et al., 1995). Interestingly, mutation of these sites led to increased phosphorylation of the $\beta 1$ subunit at Y384 and Y386, a subunit that exhibits relatively low stoichiometry of phosphorylation in response to Src compared to wild-type control (Moss et al., 1995). In primary neuronal cultures intracellular application of sodium vanadate, a potent tyrosine phosphatase inhibitor, enhanced benzodiazepine-sensitive GABA_AR function, suggesting high endogenous tyrosine kinase and phosphatase activity under basal conditions (Moss et al., 1995). Consistent with this studies using phospho-antibodies have revealed that Y365/7 are principally phosphorylated via the activity of Fyn and Src. Significantly, Fyn is able to bind specifically to phospho-Y367, providing a mechanism for the recruitment of this tyrosine kinases to GABA_ARs (Brandon et al., 2001; Jurd et al., 2010). Studies with phospho-antibodies have revealed significant variations in the stoichiometry of Y367 phosphorylation in the brains of rodents, suggesting input-specific control of phosphorylation; however, the signaling pathways that regulate the phosphorylation of the $\gamma 2$ subunit remain to be defined. In keeping with their roles in mediating high affinity binding to AP2 phosphorylation of Y365/7 increases the cell surface stability of GABA_ARs and enhances the amplitude and frequency of mIPSCs (Kittler et al., 2008; Tretter 2009).

XI. Concluding Remarks

GABA_ARs play a central role in mediating neuronal inhibition and mediating the effects of a broad range of anxiolytic, anticonvulsant, hypnotic and sedative agents. GABA_ARs mediate both phasic and tonic modes of inhibition, phenomena that are mediated via receptor subtypes with distinct molecular structures. Phasic inhibition is mediated largely by benzodiazepine-sensitive receptor subtypes that are highly enriched at synaptic sites and principally assembled from α 1–3, β 1–3 and γ 2 subunits. A critical determinant of receptor number at inhibitory synapses is the steady-state cell levels of these receptors on the neuronal plasma membrane. This is likely determined by the relative rates of receptor exo- and endocytosis, processes that are intimately controlled by covalent modifications and interaction with specific binding partners. While the molecular details of these processes are being evaluated their significance for the efficacy of neuronal inhibition and ultimately for behavior remains to be established. Our comprehension of how GABA_ARs are selectively stabilized at inhibitory synapses is limited; however, a central role for gephyrin is emerging. The direct binding of gephyrin to the GABA_AR subtypes containing α 2 subunits has been recently (Tretter et al., 2008) The significance of this interaction for GABA_AR subtypes containing other α -subunit isoforms warrants further investigation. Gephyrin-independent clustering mechanisms have also been postulated in particular for subtypes containing α 5 subunits The importance of these emerging processes for other receptor subtypes remains to be demonstrated.

The cellular mechanisms that regulate the cell surface accumulation are under active investigation. Future studies to ascertain the significance of these molecular mechanisms in determining the efficacy of neuronal inhibition under both normal and pathological conditions are essential.

Acknowledgements

This work was supported by the National Institutes of Health National Institute of Neurological Disorders and Stroke [Grants R01NS047478, R01NS048045, R01NS051195, R01NS056359 and R01NS054900 to SJM], and by the Wellcome Trust. MT is supported by a National American Heart Association Senior Scientist Award.

Non standard abbreviations

AKAP	A-kinase anchoring protein
GABA_AR	γ -aminobutyric acid type-A receptor
BIG-2	brefeldin A-inhibited guanine nucleotide exchange factor
BDNF	Brain-derived neurotrophic factor
PKA	cAMP-dependent protein kinase
CaMKII	calcium/calmodulin-dependent kinase II
PKC	calcium/phospholipiddependent protein kinase
PKM	constitutively active catalytic domain of PKC
DGC	dystrophin glycoprotein complex

ER	endoplasmic reticulum
GABARAP	GABA _A R associated protein
GEF	guanine-nucleotide exchange factor
GODZ	Golgi-specific DHHC zinc finger domain protein
GST	glutathione S-transferase
ICD	intracellular domain
HEK-293	Human embryonic kidney cell line A293
5-HT₃	5-hydroxytryptamine type-3 receptor
NL	Neuroigin
(nACh)	nicotinic acetylcholine receptor
Plic	phospholipase C-related catalytically inactive protein
PP	protein phosphatase
PSD	postsynaptic densities
PKB/Akt	protein kinase B
Plic-1	PP, protein that links the integrin-associated protein with the cytoskeleton
RACK-1	Receptor for activated C kinase
TTX	Tetrodotoxin
TM	Transmembrane domain
UBL	Ubiquitin-like
UBA	ubiquitin-associated domain

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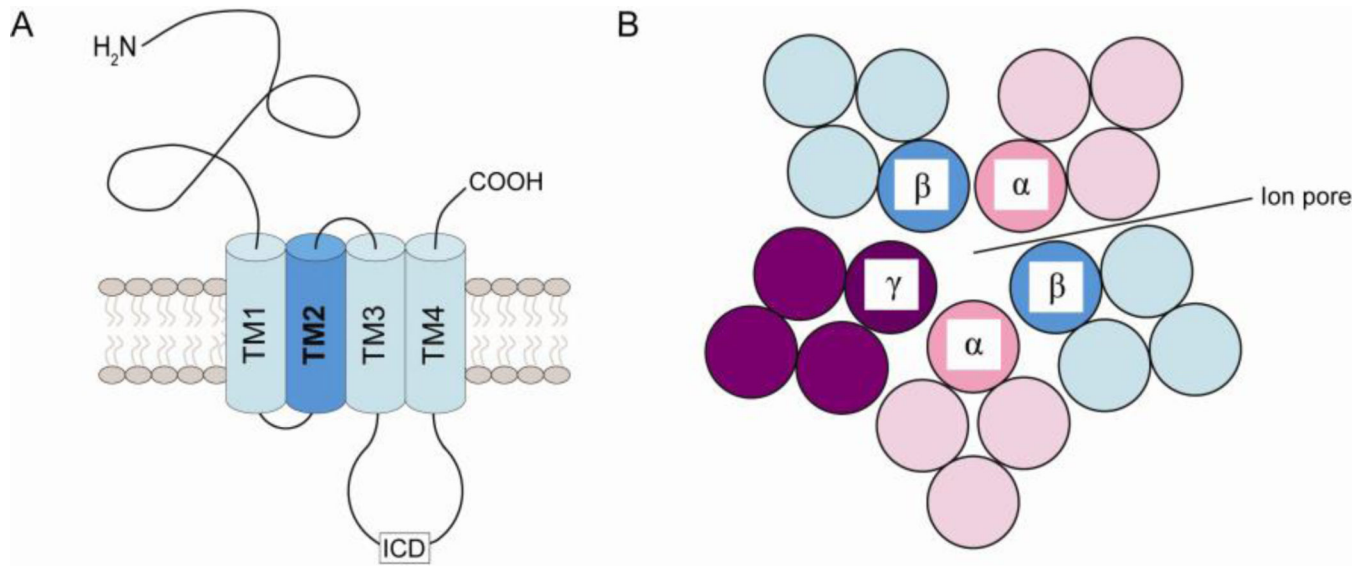


Figure 1. GABA_AR structure

(A) Transmembrane topology of the GABA_AR. Each receptor subunit is composed of a large extracellular ligand-binding N-terminal region that is also the site of action of various drugs followed by four hydrophobic transmembrane (TM)1–4 α -helices and a short, barely extruding C-terminus. Each receptor subunit also contains a large intracellular domain (ICD) between TM3 and TM4 that mediates the majority of protein-protein interactions and is subject to a number of post-translational modifications. (B) Traverse view of the assembly of GABA_AR subunits to form an ion channel. TM2 faces the lumen of the channel ion pore and TM4 is anchored in the lipid membrane. TM1 and TM3 interact with the neighboring subunits.

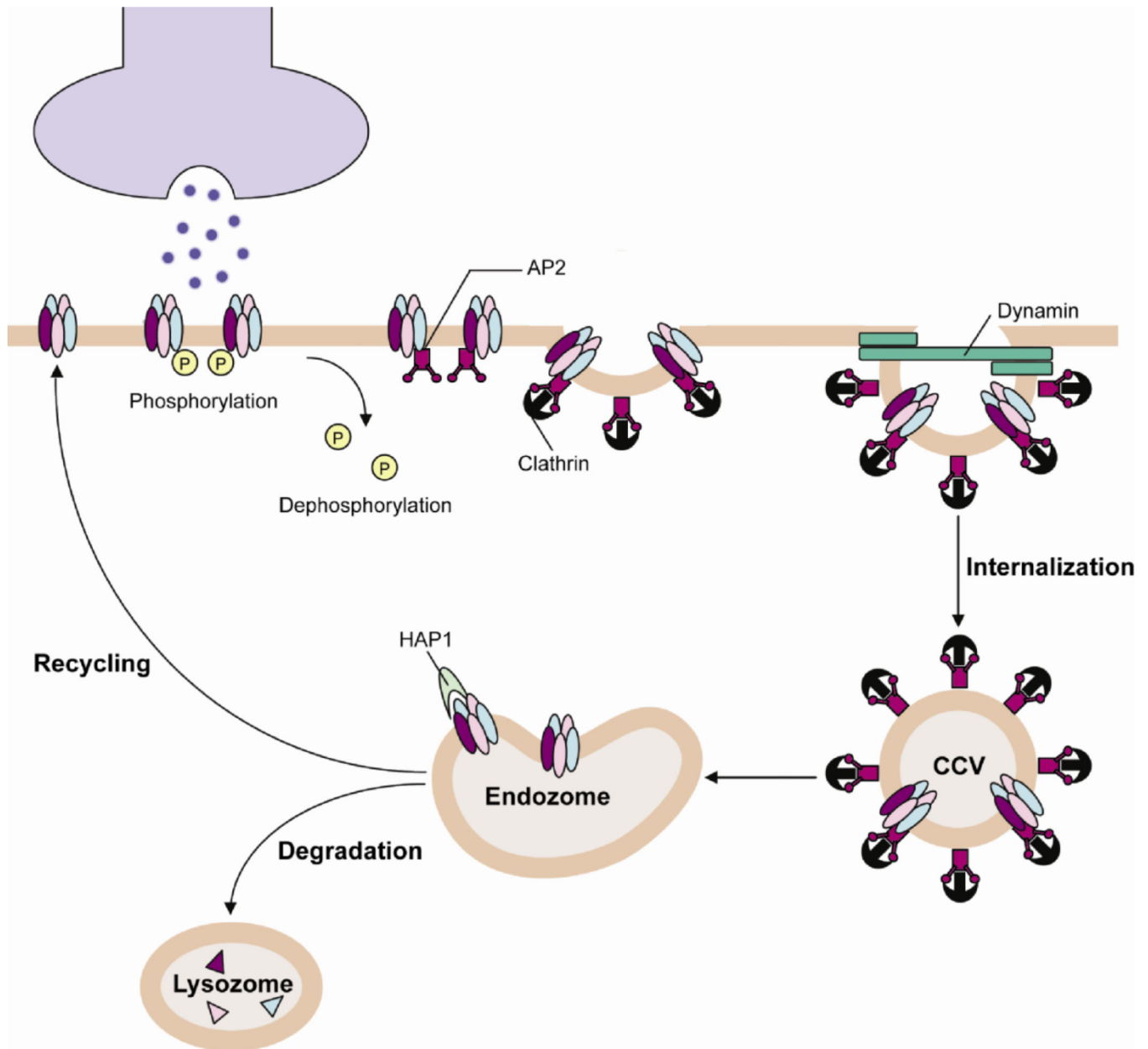


Figure 2. Clathrin-mediated endocytosis

The receptors cluster in specialized sites at the plasma membrane known as clathrin-coated pits, which invaginate and pinch off to form clathrin-coated vesicles (CCVs), a process that is dependent on dynamin. The clathrin adaptor protein (AP)-2 is a central component in the formation of these vesicles, forging a link between membrane proteins and clathrin that forms the outer layer of the coat. The vesicles subsequently lose their coat and fuse together to form an early endosome. Internalized receptors are then either subject to rapid recycling or are targeted for lysosomal degradation, an endocytic sorting decision that is regulated by the Huntingtin-associated protein (HAP)-1.

Table 1GABA_A receptor associated proteins

Protein	Subcellular Localization	Subunit specificity	Proposed function
GABARAP	Golgi	γ	Facilitates exocytosis
PRIP1/2	Synapses	β	Enhances cell surface stability by inhibiting PP1 α mediated dephosphorylation
PLIC1	Intracellular compartments	α/β	Facilitates exocytosis by inhibiting proteasome-dependent degradation
BIG2	Intracellular compartments	β	Promotes ER exit and structural integrity of recycling endosomes
AP2	Clathrin-coated pits	β/γ	Regulates clathrin-mediated endocytosis
HAP1	Endosomes	β	Regulates post-endocytic sorting by inhibiting lysosomal degradation
GODZ	Golgi	γ	Palmitoylation
Gephyrin	Synapses	$\alpha 1$	Facilitates clustering/scaffolding at synaptic sites
Radixin	Extrasynaptic sites	$\alpha 5$	Facilitates clustering at extrasynaptic sites
GRIF1	Intracellular compartments	$\beta 2$	Unknown
gC1qR	Intracellular compartments	$\beta 3$	Unknown

Adapted from Arancibia-Carcamo and Moss, 2006

Table 2GABA_A receptor phosphorylation sites

Subunit	Phosphorylation site	Protein kinase		
		<i>In vitro</i>	Heterologous cell lines	Primary neurons
β1	S384	CaMKII	PKA, PKC	
	S409	PKA, PKC, CaMKII, PKG		
β2	S410	PKA, PKC, Akt, CaMKII, PKG	PKC, Akt	Akt
β3	S383	CaMKII	PKA, PKC PKA, PKC	PKA, PKC PKA, PKC
	S408	PKC		
	S409	PKA, PKC, CaMKII, PKG		
γ2	S327	PKC		
	S343	PKC, CaMKII		
	S348	CaMKII	Src Src	
	T350	CaMKII		
	Y365	Src		
	Y367	Src		

Adapted from Brandon et al., 2002

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