Analyzing the mechanisms that facilitate the subtype-specific assembly of \( \gamma \)-aminobutyric acid type A receptors

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Impaired inhibitory signaling underlies the pathophysiology of many neuropsychiatric and neurodevelopmental disorders including autism spectrum disorders and epilepsy. Neuronal inhibition is regulated by synaptic and extrasynaptic \( \gamma \)-aminobutyric acid type A receptors (GABA\(_{A}\)Rs), which mediate phasic and tonic inhibition, respectively. These two GABA\(_{A}\)R subtypes differ in their function, ligand sensitivity, and physiological properties. Importantly, they contain different \( \alpha \) subunit isoforms: synaptic GABA\(_{A}\)Rs contain the \( \alpha_{1-3} \) subunits whereas extrasynaptic GABA\(_{A}\)Rs contain the \( \alpha_{4-6} \) subunits. While the subunit composition is critical for the distinct roles of synaptic and extrasynaptic GABA\(_{A}\)R subtypes in inhibition, the molecular mechanism of the subtype-specific assembly has not been elucidated. To address this issue, we purified endogenous \( \alpha_{1} \) and \( \alpha_{4} \)-containing GABA\(_{A}\)Rs from adult murine forebrains and examined their subunit composition and interacting proteins using liquid chromatography coupled with tandem mass spectrometry (LC-MS/MS) and quantitative analysis. We found that the \( \alpha_{1} \) and \( \alpha_{4} \) subunits form separate populations of GABA\(_{A}\)Rs and interact with distinct sets of binding proteins. We also discovered that the \( \beta_{3} \) subunit, which co-purifies with both the \( \alpha_{1} \) and \( \alpha_{4} \) subunits, has different levels of phosphorylation on serines 408 and 409 (S408/9) between the two receptor subtypes. To understand the role S408/9 plays in the assembly of \( \alpha_{1} \) and \( \alpha_{4} \)-containing GABA\(_{A}\)Rs, we examined the effects of S408/9A (alanine) knock-in mutation on the subunit composition of the two receptor subtypes using LC-MS/MS and quantitative analysis. We discovered that the S408/9A mutation results in the formation of novel \( \alpha_{1}\alpha_{4} \)-containing GABA\(_{A}\)Rs. Moreover, in S408/9A mutants, the plasma membrane expression of the \( \alpha_{4} \) subunit is increased whereas its retention in the endoplasmic
GABA<sub>A</sub>-aminobutyric acid type A receptors (GABA<sub>A</sub>Rs) are ligand-gated ion channels that mediate neuronal inhibition in the adult brain (Farrant and Nusser, 2005). As the primary regulators of neuronal inhibition, they are also the therapeutic sites of action for benzodiazepines, barbiturates, intravenous anesthetics, and neurosteroids (Rudolph and Möhler, 2006). GABA<sub>A</sub>Rs are heteropentamers that can be assembled from the α<sub>1–6</sub>, β<sub>1–3</sub>, γ<sub>1–3</sub>, δ, ε, θ, π, and ρ<sub>1–3</sub> subunits (Rudolph and Möhler, 2006). This subunit diversity provides the structural basis for the generation of distinct GABA<sub>A</sub> R subtypes endowed with distinct physiological and pharmacological properties. While the identity of the endogenous GABA<sub>A</sub>R subtypes that are assembled in neurons remains to be explored, consensus opinion suggests that phasic inhibition is principally mediated by receptor subtypes composed of the α<sub>1–3</sub>, β<sub>1–3</sub>, and γ<sub>2</sub> subunits (Farrant and Nusser, 2005). Tonic inhibition is believed to be mediated by specialized populations of extrasynaptic GABA<sub>A</sub>Rs, which are composed of the α<sub>4–6</sub> and β<sub>2–3</sub> subunits with or without the γ<sub>0</sub> or δ subunit (Farrant and Nusser, 2005).

Individual neurons such as dentate gyrus granule cells (DGGCs) or CA1 principal neurons exhibit both robust phasic and tonic currents. Consistent with this, immunohistochemical studies have shown that single neurons can express the α<sub>1–5</sub>, β<sub>1–3</sub>, γ<sub>1–3</sub>, and δ subunits (Fritschy and Möhler, 1995; Spérk et al., 1997; Pirker et al., 2000). Significantly, there is accumulating experimental evidence showing that the efficacy of phasic and tonic inhibition can be dynamically modulated by reproductive hormones and neurosteroids (Adams et al., 2015; Modgil et al., 2017, 2019; Mukherjee et al., 2017; Parakala et al., 2019). Moreover, deficits in the "equilibrium" between phasic and tonic inhibition are widely believed to play central roles in the pathophysiology of anxiety, depression, epilepsy, and autism spectrum disorders (Farrant and Nusser, 2005; Benham et al., 2014; Tang et al., 2021).

To date, recombinant expression has been used to understand how hetero-oligomeric GABA<sub>A</sub>Rs are constructed from their component subunits. Such experiments have revealed that individual GABA<sub>A</sub>R subunits are assembled into heteropentamers in the endoplasmic reticulum (ER) (Jacob et al., 2008). GABA<sub>A</sub>Rs that reach 'conformation maturity' and form αβγ-, αβδ-, or αβδ-containing receptors exit the ER for transport to and insertion into the plasma membrane (PM) (Connolly et al., 1996; Jacob et al., 2008; Vithlani et al., 2011). Unassembled subunits and transport incompetent combinations are targeted for proteasomal degradation (Jacob et al., 2008). Both synaptic and extrasynaptic GABA<sub>A</sub>Rs enter the PM at extrasynaptic locations and can access the synaptic space (Thomas et al., 2005; Bogdanov et al., 2006). However, only the receptors that contain the α<sub>1–3</sub> subunits are selectively stabilized at synapses due to their affinity for proteins within the postsynaptic scaffold such as gephyrin (Moss and Smart, 2001; Jacob et al., 2008). Thus, the subunit composition of a GABA<sub>A</sub>R, and specifically the α subunit isoform, contributes to the receptor's localization within the PM which is critical for its function. However, how neurons orchestrate the segregation of the α subunit variants into distinct receptor subtypes that mediate phasic and tonic inhibition remains to be defined.

To address this issue, we isolated native GABA<sub>A</sub>Rs subtypes containing the α and α4 subunits from the murine forebrain and used quantitative mass spectrometry to examine their subunit composition. We found that the α1 and α4 subunits formed discrete populations of GABA<sub>A</sub>Rs that associate with distinct sets of binding proteins, but both contained the β3 subunit. Interestingly, our analysis did not detect the δ subunit co-purifying with the α4 subunit. Additionally, the β3 subunit in α1-containing GABA<sub>A</sub>Rs exhibited higher levels of phosphorylation on serines 408 and 409 (S408/9) than the β3 subunit in the α4 subtype. Previous studies have shown that these residues are critical sites for phospho-dependent regulation of GABA<sub>A</sub>R membrane trafficking, which impacts the efficacy of GABAergic inhibition (Kittler et al., 2005; Vien et al., 2015). Mutation of S408/9 compromised the segregation of the α1 and α4 subunits into biochemically distinct receptor subtypes. Mechanistically this mutation acted to reduce the ER retention of the α4 subunit and to increase its accumulation in the PM. Thus, our results suggest a critical role for the β3 subunit and its phosphorylation on S408/9 in orchestrating the subtype-specific assembly of GABA<sub>A</sub>Rs.
Results

Purification and analysis of native GABA$_A$Rs that are assembled from the $\alpha$1 subunit

While it is widely accepted that GABA$_A$Rs containing the $\alpha$1 subunit are major mediators of phasic inhibition (Farrant and Nusser, 2005), quantitative analysis of their subunit composition in the neuronal PM has not been reported. To address this issue, adult murine forebrain tissues were collected and homogenized in isotonic, detergent-free buffers and subjected to serial centrifugation for isolation of highly purified PM fractions (Suski et al., 2014; Smalley et al., 2020). Purified PM was solubilized in 0.5% Triton X-100, a detergent that efficiently extracts GABA$_A$Rs from their native environment without compromising their structure (Chen et al., 2012). To assess the molecular mass of native $\alpha$1-containing GABA$_A$Rs in PM, we subjected the purified material to Blue Native PAGE (BN-PAGE) and immunoblotted with the $\alpha$1 subunit antibody. Two major bands were identified - one at around 250 kDa, which is consistent with the pentameric structure of GABA$_A$Rs, and the other at around 720 kDa (Figure 1A). Next, the solubilized PM fraction was exposed to a monoclonal antibody against the $\alpha$1 subunit or non-immune mouse IgG immobilized on magnetic beads and eluted in a buffer containing 2% Tween and 0.01% SDS, a combination sufficient to elute the material from the beads without disturbing stable protein complexes (Antrobus and Borner, 2011). As measured using immunoblotting, high levels of the $\alpha$1 subunit were retained on the $\alpha$1 subunit antibody compared to flow through (Supplementary Figure 1A). The eluted material was resolved on BN-PAGE and visualized using immunoblotting. Both 250 and 720 kDa bands were detected in the eluted material with the $\alpha$1 subunit antibody but not in material purified on IgG control (Figure 1A).

To interrogate the subunit composition of GABA$_A$Rs containing the $\alpha$1 subunit, the gel area containing the lower molecular weight species at around 250 kDa was excised and its contents were analyzed with liquid chromatography coupled with tandem mass spectrometry (LC-MS/MS). The LC-MS/MS data was analyzed with label-free, globally normalized quantification. Briefly, the spectral index of each protein was visualized using immunoblotting, which revealed the 250 kDa material purified with the $\alpha$1 antibody against the $\alpha$1 subunit $\alpha$1–3, $\beta$1–3, $\gamma$2–3, and $\delta$ subunits are significantly enriched in major populations of $\alpha$1-containing GABA$_A$Rs, compared to the IgG control (Figure 1B and Supplementary Table 1A). The most abundant subunits were the $\alpha$1 subunit (SI$_{GI}$ of 0.0087 ± 0.0009, $p < 0.0001$, $n = 5$, Welch’s $t$-test), followed by the $\beta$3 (0.0035 ± 0.0008, $p = 0.0026$, $n = 5$, Welch’s $t$-test), $\gamma$2 (0.0024 ± 0.0003, $p < 0.0001$, $n = 5$, Welch’s $t$-test), $\beta$2 (0.0014 ± 0.0003, $p = 0.0027$, $n = 5$, Welch’s $t$-test), $\alpha$3 (0.0007 ± 0.0001, $p = 0.0009$, $n = 5$, Welch’s $t$-test), $\beta$1 (0.0006 ± 0.0002, $p = 0.0069$, $n = 5$, Welch’s $t$-test), $\delta$ (0.0004 ± 0.0009, $p = 0.0222$, $n = 5$, Welch’s $t$-test), $\alpha$2 (0.0003 ± 0.0009, $p = 0.0094$, $n = 5$, Welch’s $t$-test), and $\gamma$3 (0.0001 ± 0.00003, $p = 0.0050$, $n = 5$, Welch’s $t$-test) subunits (Figure 1B and Supplementary Table 1A). Based on the relative abundance, $\alpha$1$\beta$3$\gamma$2 is likely to be the most common subunit composition of $\alpha$1-containing GABA$_A$Rs. The $\alpha$4 (0.0002 ± 0.0008, $p = 0.1194$, $n = 5$, Welch’s $t$-test), $\alpha$5 (0.0004 ± 0.0002, $p = 0.0723$, $n = 5$, Welch’s $t$-test), and $\gamma$1 (0.00003 ± 0.00003, $p = 0.2035$, $n = 5$, Welch’s $t$-test) subunits were detected but not significantly enriched when compared to IgG control.

Purification and analysis of native GABA$_A$Rs that are assembled from the $\alpha$4 subunit

We also analyzed the subunit composition of GABA$_A$Rs that contain the $\alpha$4 subunit, which are the principal mediators of tonic current in the adult forebrain (Chandra et al., 2006). To quantitatively analyze the subunit composition of $\alpha$4-containing GABA$_A$Rs in the neuronal PM, we first subjected purified PM fractions from murine forebrain tissues to BN-PAGE and immunoblotted with the $\alpha$4 subunit antibody. A major band at around 250 kDa and a weaker band at around 720 kDa were observed (Figure 1C). The PM fraction was then subjected to immunoprecipitation with an immobilized monoclonal antibody against the $\alpha$4 subunit or non-immune mouse IgG and eluted under non-denaturing conditions as described above. The effectiveness of immunoprecipitation with the $\alpha$4 subunit antibody was verified via immunoblotting (Supplementary Figure 1B). The eluted material was resolved by BN-PAGE and visualized using immunoblotting, which revealed the 250 kDa band in the eluted material purified with the $\alpha$4 subunit antibody (Figure 1C). No immunoreactivity was observed with IgG control (Figure 1C).

LC-MS/MS was performed on the major band at 250 kDa and the data were analyzed with label-free normalized quantification and Welch’s $t$-test. We found that only the $\alpha$4 (0.0046 ± 0.0011, $p = 0.0049$, $n = 5$, Welch’s $t$-test) and $\beta$3 (0.0010 ± 0.0002, $p = 0.0013$, $n = 5$, Welch’s $t$-test) subunits are significantly enriched in major populations of $\alpha$4-containing GABA$_A$Rs, when compared to IgG control (Figure 1D and Supplementary Table 1B). The $\alpha$5, $\gamma$1, and $\gamma$3 subunits were
At steady state, the endogenous α1 and α4 subunits are segregated into distinct GABA_A subtypes. **FIGURE 1**

(A) Purified plasma membrane fraction (PM), PM exposed to non-immune IgG, and PM exposed to the α1 subunit antibody for purification of GABA_A Rs that contain the α1 subunit (α1 IP) were resolved on Blue Native PAGE (BN-PAGE) gels and probed for the α1 subunit. Major bands at around 250 and 720 kDa, which represent heteropentameric GABA_A Rs in isolation and heteropentameric GABA_A Rs with binding partners, respectively, were observed.

(B) Gel regions containing the 250 kDa band were subjected to liquid chromatography coupled with tandem mass spectrometry (LC-MS/MS). Analysis of the 250 kDa band with label-free quantification and Welch’s t-test revealed that in α1-containing GABA_A Rs, the α1–3, β1–3, γ2–3, and δ subunits are significantly enriched when compared to non-immune IgG control. Detected but non-significant subunits are not shown (**p < 0.01, ***p < 0.001, ****p < 0.0001, n = 5 replicates).

(C) Purified PM, PM exposed to non-immune IgG, and PM exposed to the α4 subunit antibody (α4 IP) were resolved on BN-PAGE and probed with the α4 subunit antibody. A band at 250 kDa and a weaker band at 720 kDa were observed with immunoblotting.

(D) The contents of the 250 kDa band were analyzed with LC-MS/MS. Only the α4 and β3 subunits are significantly enriched in α4-containing GABA_A Rs when compared to non-immune IgG (**p < 0.01, n = 5 replicates).

(E) The amount of target subunit recovery in 250 kDa bands is higher for the α1 subunit than the α4 subunit (*p < 0.05, n = 5 replicates).

**not detected at all, and others – the α1 (0.0001 ± 0.00007, p = 0.0967, n = 5, Welch’s t-test), α2 (0.00003 ± 0.00003, p = 0.4858, n = 5, Welch’s t-test), α3 (0.00002 ± 0.00002, p = 0.2035, n = 5, Welch’s t-test), α6 (0.0002 ± 0.0002, p = 0.0994, n = 5, Welch’s t-test), β1 (0.00005 ± 0.00002, p = 0.0973, n = 5, Welch’s t-test), β2 (0.0001 ± 0.00008, p = 0.1226, n = 5, Welch’s t-test), γ2 (0.000006 ± 0.000006, p = 0.2035, n = 5, Welch’s t-test), and δ (0.000008 ± 0.000005, p = 0.0983, n = 5, Welch’s t-test) subunits – were detected in low amounts but not significantly enriched when compared to IgG control.**

While the α1 and α4 subunits are mutually exclusive, the β3 subunit is significantly enriched in both α1- and α4-containing populations of GABA_A Rs (Figures 1B,D and Supplementary Tables 1A,B).

A significant limitation in comparing data sets between α1- and α4-containing GABA_A Rs is controlling for the recovery of
the target receptor subtype between purifications. To control for this variable, we directly compared the amount of each target α subunit isolated in the respective purifications. We found that there is an approximately twofold higher level of recovery of the α1 subunit compared to the α4 subunit (Figure 1E; α1 IP: 0.0087 ± 0.0009, α4 IP: 0.0046 ± 0.0011, p = 0.0211, n = 5, unpaired t-test). The lower recovery of the α4 subunit is likely to reflect that the α1 subunit is expressed at about 15-fold higher levels than the α4 subunit in the brain, as quantified by mass spectrometry (Chen et al., 2012). Collectively, our experiments strongly suggest that neurons have the capacity to segregate the α1 and α4 subunits into distinct receptor subtypes with distinct subunit composition.

**Analysis of the proteins that co-purify with α1-containing GABA_ARs**

In addition to GABA_ARs subunits that co-purify with the α1 subunit, we examined which proteins associate with α1-containing GABA_ARs. To do so, we used LC-MS/MS and quantitative analysis to reveal the contents of the gel region containing the 720 kDa bands, which were evident in purifications of α1-containing GABA_ARs. Welch’s t-test was performed against non-immune mouse IgG to identify significantly enriched proteins, and only those that were detected in all replicates were included for subsequent analysis. For the α1 subunit, 93 proteins were identified as significantly enriched proteins (Supplementary Table 2A). Across all replicates, the most abundantly found proteins are Sptan1 (0.2476 ± 0.0426, p = 0.0008, n = 5, Welch’s t-test), Sptbn1 (0.1078 ± 0.0165, p = 0.0005, n = 5, Welch’s t-test), and Sptbn2 (0.0292 ± 0.0031, p < 0.0001, n = 5, Welch’s t-test), which are different isoforms of the spectrin family (Figure 2A and Supplementary Table 2A). Together these isoforms of spectrin constitute over 70% of all the proteins co-purified with the α1 subunit – 47% Sptan1, 20% Sptbn1, and 6% Sptbn2 (Figure 2B). Each of the remaining 90 proteins makes up 5% or less of the total amount of proteins detected with α1-containing GABA_ARs (Figure 2B). These suggest that the detection of the high molecular weight species of α1-containing GABA_ARs at around 720 kDa is mainly driven by the presence of spectrin isoforms. Other notable proteins that are significantly enriched with the α1 subunit are known inhibitory scaffolding proteins such as Cyfip2 (0.0008 ± 0.0002, p = 0.0074, n = 5, Welch’s t-test), and Gphn (gephyrin; 0.0005 ± 0.0001, p = 0.0017, n = 5, Welch’s t-test) (Kneussel et al., 1999; Davenport et al., 2019) (Supplementary Table 2A).

To confirm the association of the proteins identified by LC-MS/MS with the α1 subunit, we resolved purified α1-containing GABA_ARs on BN-PAGE and immunoblotted with antibodies against some of the proteins detected in high- and low-abundance. For high-abundance proteins, we probed for Sptan1, Sptbn1, and Sptbn2 and observed their immunoreactivity at around 720 kDa (Figure 2C). We also assessed the presence of low-abundance interactors – Ercc2, Kcc2 (Slc12a5), and gephrin. For Ercc2 and Kcc2, their immunoreactivity was observed slightly below 720 kDa, with Kcc2 showing an additional band between 242 and 480 kDa (Figure 2C). Gephyrin was observed higher than 720 kDa, at approximately 1.048 kDa (Figure 2C). For these high- and low-abundance interactors of the α1 subunit, the immunoreactivity was specific to purified α1-containing GABA_ARs (Figure 2C), which provided confidence in the veracity of our approach.

Next, we performed the network analysis on the significantly enriched proteins of the α1 subunit to gain insights into the known relationships between the proteins. For the analysis, we included proteins that are (1) significantly enriched, (2) detected in all replicates, and (3) have average SlG values greater than 0.0005. We used STRINGdb database to impute known interactions between proteins (Szklarczyk et al., 2019) and scaled each node to the average SlG value of the protein. Lastly, we overlaid the highest scoring Gene Ontology (GO) Biological Process term for each protein to provide information on the protein function (Szklarczyk et al., 2019). The network of the α1 subunit shows interactions between the spectrin isoforms (Sptan1, Sptbn1, Sptbn2, and Sptb) and ankyrin isoforms (Ank2 and Ank3), as well as interactions between gephrin (Gphn) and Kcc2 (Slc12a5) (Figure 2D). Many of the associated proteins of the α1 subunit are involved in cytoskeleton organization, ion transport, or signal transduction (Figure 2D). These results show that the α1 subunit interacts with scaffolding and cytoskeletal proteins that stabilize α1-containing GABA_ARs in the PM.

**Analysis of the proteins that co-purify with α4-containing GABA_ARs**

We also analyzed the proteins associated with the α4 subunit as described above. The α4 subunit has 19 significantly enriched proteins, none of which are regarded as inhibitory scaffolding proteins (Supplementary Table 2B). Proteins that interact with the α4 subunit in the order of abundance are Ttn (titin; 0.0039 ± 0.0004, p = 0.0464, n = 5, Welch’s t-test), Myo5a (0.0026 ± 0.0010, p = 0.0416, n = 5, Welch’s t-test), and Dsp (desmoplakin; 0.0017 ± 0.0005, p = 0.0271, n = 5, Welch’s t-test) (Figure 3A and Supplementary Table 2B). Only two proteins were detected with both α1- and α4-containing GABA_ARs: Myo5a and Vdac2 (Figures 2A, 3A). Unlike in α1-containing GABA_ARs, there is not a single protein or a family of proteins that dominates the makeup of the proteins purified with α4-containing GABA_ARs – 20% titin, 13% Myo5a, 9% desmoplakin, followed by 8% or less of each of the remaining 16 proteins (Figure 3B). The inhibitory scaffolding protein gephrin was not detected with the α4 subunit. Consistent
The endogenous α1 subunit interacts with inhibitory scaffolding proteins and cytoskeletal proteins. (A) The 720 kDa band of α1-containing GABA\(_A\)Rs purified from PM was subjected to LC-MS/MS. Label-free normalized quantification and Welch’s t-test against non-immune control IgG revealed 93 significantly enriched binding proteins of the α1 subunit. The heatmap shows the top 10 binding proteins of α1-containing GABA\(_A\)Rs by abundance (SI GI value) across all replicates. The most abundant proteins detected with the α1 subunit are Sptan1, Sptbn1, and Sptbn2 (\(n = 5\) replicates). (B) Pie chart shows the relative abundance of the proteins that are significantly enriched with α1-containing GABA\(_A\)Rs. Sptan1, Sptbn1, and Sptbn2 together constitute more than 70% of the total amount of proteins. Proteins that make up less than 2% of the total amount of proteins (86 proteins) were grouped together as ‘Other.’ (C) Immunopurified α1-containing GABA\(_A\)Rs were resolved on BN-PAGE and probed for the α1 subunit, high-abundance proteins (Sptan1, Sptbn1, and Sptbn2), and low-abundance proteins (Erc2, Kcc2, and gephyrin). Sptan1, Sptbn1, and Sptbn2 are observed at around 720 kDa with α1-containing GABA\(_A\)Rs. Erc2 and Kcc2 are present slightly below 720 kDa and gephyrin is observed higher than 720 kDa. The immunoreactivity of these proteins is specific to purified α1-containing GABA\(_A\)Rs. (D) The network analysis shows known inhibitory scaffolding proteins such as Actn1, Cypfl2, and Gphn (gephyrin), and a subnetwork of spectrins (Sptan1, Sptbn1, Sptbn2, and Sptb) and ankyrins (Ank2 and Ank3). GO analysis reveals that most proteins are involved in cytoskeletal organization, ion transport, or signal transduction.
FIGURE 3
The endogenous α4 subunit does not interact with inhibitory scaffolding proteins. (A) Analysis of the 720 kDa band of α4-containing GABA_A Rs with LC-MS/MS and quantitative analysis identified 19 binding proteins. The heatmap shows the top 10 binding proteins by abundance, which include Ttn (titin), Myo5a, and Dsp (desmoplakin). Myo5a and Vdac2 were detected with both the α1 and α4 subtypes (n = 5 replicates). (B) The relative abundance of binding proteins of the α4 subunit is illustrated as a pie chart. Besides 20% by titin and 13% by Myo5a, most proteins constitute less than 10% of the total amount of proteins detected with α4-containing GABA_A Rs. Proteins that make up less than 2% each (6 proteins) were grouped together as ‘Other.’ (C) BN-PAGE blots of purified α4-containing GABA_A Rs show the absence of gephyrin in the α4 subtype. (D) The network analysis reveals no known interactions among the proteins detected with the α4 subunit. (E) PCA plot of the significantly enriched binding proteins of the α1 and α4 subunits on the 720 kDa band reveals high reproducibility of datasets between replicates. It also highlights the difference in the binding proteins of the two receptor subtypes. (F) Target subunit recovery in 720 kDa bands between subtype purifications is comparable (ns ≥ 0.05, n = 5 replicates).
with this, when purified α4-containing GABA₄Rs were resolved on BN-PAGE and immunoblotted with the gephyrin antibody, no immunoreactivity was observed (Figure 3C). The network analysis revealed that there are no known interactions among the proteins detected with the α4 subunit (Figure 3D). These results suggest that the α4 subunit does not interact with components of the sub-synaptic cytoskeleton.

Comparing the multiprotein complexes of α1- and α4-containing GABA₄Rs

To visualize the similarity or difference among the associated proteins of α1- and α4-containing GABA₄Rs and to assess the reproducibility of our purifications, we performed the principal component analysis (PCA). The replicates of the α1 subunit were tightly grouped together, suggesting high reproducibility of the α1 subunit purifications (Figure 3E). It also suggests that there is a core group of proteins that is commonly identified with the α1 subunit across all replicates. The replicates of the α4 subunit, on the other hand, were more spread out on the PCA plot (Figure 3E), suggesting the lack of common components across all replicates. Importantly, the two types of purifications were clearly separated (Figure 3E), further suggesting that the α1 and α4 subunits associate with distinct sets of binding proteins. The first principal component (PC1) accounted for 51.78% of the variance and was positively correlated with Sptan1 and Sptbn1, the two most abundantly detected proteins of the α1 subunit (Figure 2). The second principal component (PC2) accounted for 18.71% of the variance. Together, our findings show that the α1 and α4 subunits form separate receptor populations with distinct subunit composition and binding proteins.

Finally, we compared the amount of target subunit detected in the 720 kDa bands between purifications of the two GABA₄R populations. We found that there was no significant difference between the amounts of the α1 and α4 subunits in each purification (Figure 3F; α1 IP: 0.0008 ± 0.0002, α4 IP: 0.0005 ± 0.0002, p = 0.2205, n = 5, unpaired t-test). This suggests that the input amounts of α1- and α4-containing GABA₄Rs in our experiments were comparable.

Examining the levels of the β3 subunit phosphorylation in GABA₄Rs assembled from the α1 and α4 subunits

The β3 subunit is a common component of both α1- and α4-containing GABA₄Rs (Figures 1B,D and Supplementary Tables 1A,B). It has been reported that simultaneous phosphorylation of this subunit on both serine 408 and serine 409 (S408/9) acts to regulate the membrane trafficking of GABA₄Rs by decreasing their binding to adaptins, which are critical for the retrograde protein transport in the endocytic pathway (Kittler et al., 2005; Vien et al., 2015). To examine whether phosphorylation of S408/9 plays a role in the segregation of the α1 and α4 subunits into distinct receptor subtypes, we purified the two GABA₄R populations from the PM and measured the ratio of phosphorylated S408/9 (p-S408/9) to total β3 subunit using a phospho-specific antibody that only recognizes the β3 subunit when both S408 and S409 are phosphorylated (Brandon et al., 2002b, 2003; Jovanovic et al., 2004; Saliba et al., 2012; Vien et al., 2015; Parakala et al., 2019). This approach revealed that the percentage of p-S408/9 was significantly higher in α1-containing GABA₄Rs than in the α4 subtype (Figures 4A,B; α1 IP: 47.48 ± 15.85%, α4 IP: 6.76 ± 3.74%, p = 0.0465, n = 4, unpaired t-test).

To evaluate whether there are other phosphorylation sites within the β3 subunit, we compared global β3 subunit phosphorylation between the two receptor subtypes using LC-MS/MS. Spectral searches were performed to identify phosphorylated serine (S), threonine (T), or tyrosine (Y) residues on the intracellular loop domain of the β3 subunit, which is the site of post-translational modifications that regulate receptor trafficking and PM expression of GABA₄Rs (Moss et al., 1992; Brandon et al., 2002a; Nymann-Andersen et al., 2002; O’Toole and Jenkins, 2011). The levels of phosphorylation on these residues were then estimated by comparing the number of phosphorylated and dephosphorylated peptides. This approach is valid as our purifications do not employ a phospho-enrichment step. In receptors that contain the α1 subunit, in addition to peptides phosphorylated on S408 or S409, LC-MS/MS analysis detected phosphorylated S383 (Figure 4C), which is a previously identified phosphorylation site that is modulated by calcium/calmodulin-dependent protein kinase (Saliba et al., 2012). Seven additional novel sites of phosphorylation – T322, S330, S332, Y391, S396, T402, and T419 – were also evident with mean phosphopeptide ratios of 0.2 – 0.7 (Figure 4C). Similar levels of phosphorylation were found at these residues in α4-containing GABA₄Rs, except at S332 where phosphorylation was not detected (Figure 4C). These results demonstrate that α1β3- and α4β3-containing GABA₄Rs are phosphorylated at multiple sites within the intracellular loop domain of the β3 subunit (Figure 4D).

Mutation of S408/9 does not change the total expression levels of the α1 or α4 subunit

To determine whether S408/9 contribute to the subtype-specific assembly of GABA₄Rs, we utilized mice in which S408/9 have been mutated to alanines (S408/9A). These animals are viable but exhibit increased seizure sensitivity, increased autism-like behaviors, and deficits in GABAergic inhibition
The β3 subunit is more highly phosphorylated at S408 and S409 in α1-containing GABA_ARs than in the α4 subtype. (A) Immunoblots of purified α1- and α4-containing GABA_ARs were resolved on SDS-PAGE and probed for the target subunit (α1 or α4 subunit), β3 subunit, and phosphorylated S408/9 (p-S408/9). (B) The percentage of p-S408/9 in respect to total β3 subunit is significantly higher in the α1 subtype than in the α4 subtype (*p < 0.05, n = 4 replicates). (C) Spectra searches were performed on LC-MS/MS data obtained from purified α1- and α4-containing GABA_ARs to identify phosphorylated serine (S), threonine (T), or tyrosine (Y) residues within the β3 subunit. Three known (S383, S408, and S409) and seven novel (T322, S330, S332, Y391, S396, T402, and T419) phosphorylation sites were detected. S332 was detected in the α1 subtype only (n = 5 replicates). (D) Illustration of a GABA_A R β3 subunit with phosphorylation sites detected by LC-MS/MS is shown. Known and novel sites are in pink and green, respectively.

(Vien et al., 2015). Thus, we investigated whether the ability of neurons to segregate the α1 and α4 subunits into distinct receptor subtypes is disrupted in animals carrying the S408/9A mutation. We first examined if the mutation alters the total expression levels of the α1 or α4 subunit in the forebrain. Levels of the α1 and α4 subunits were measured in the forebrain tissues collected from WT and S408/9A animals and normalized to GAPDH as the loading control. We found that there was no change in the level of the α1 (S408/9A: 0.9113 ± 0.0570, p = 0.7312, n = 3, unpaired t-test) or α4 subunit (S408/9A: 0.8097 ± 0.0293, p = 0.3114, n = 3, unpaired t-test) (Figure 5).

**Mutation of S408/9 leads to the formation of α1α4-containing GABA_ARs**

Next, we examined the effects of the S408/9A mutation on the subunit composition of receptors assembled from the α1 subunit. To do so, α1-containing GABA_ARs were purified from PM fractions collected from S408/9A mice, subjected to BN-PAGE, and visualized with the α1 subunit antibody. Consistent with our experiments in WT, bands at around 250 and 720 kDa were observed (Figure 6A). The major band at
around 250 kDa was then examined using LC-MS/MS and label-free quantitative analysis as detailed above. In the α1 subtype, the α1, α3–5, β1–3, γ2, and δ subunits are significantly enriched when compared to non-immune IgG control (Figure 6B and Supplementary Table 1C). The most abundantly detected subunit was the α1 subunit (0.0067 ± 0.0022, p = 0.0155, n = 5, Welch’s t-test), followed by the γ2 (0.0034 ± 0.0006, p = 0.0006, n = 5, Welch’s t-test), β2 (0.0029 ± 0.0004, p = 0.0001, n = 5, Welch’s t-test), β3 (0.0026 ± 0.0003, p < 0.0001, n = 5, Welch’s t-test), α3 (0.0010 ± 0.0002, p = 0.0038, n = 5, Welch’s t-test), β1 (0.0007 ± 0.0002, p = 0.0081, n = 5, Welch’s t-test), α4 (0.0007 ± 0.0002, p = 0.0352, n = 5, Welch’s t-test), α5 (0.0006 ± 0.0003, p = 0.0430, n = 5, Welch’s t-test), and δ (0.0002 ± 0.00009, p = 0.0367, n = 5, Welch’s t-test) subunits (Figure 6B and Supplementary Table 1C). Subunits that were detected but not found to be significant are the α2 (0.0004 ± 0.0002, p = 0.0733, n = 5, Welch’s t-test), γ1 (0.000002 ± 0.000002, p = 0.2035, n = 5, Welch’s t-test), and γ3 (0.0001 ± 0.000007, p = 0.0793, n = 5, Welch’s t-test) subunits.

From this, the most common composition of α1-containing GABA_ARs in S408/9A mice can be estimated to be α1β2/3γ2/3.

In WT, the α1–3 subunits are the only α subunit isoforms that are significantly enriched in α1-containing GABA_ARs (Figure 1B and Supplementary Table 1A). In S408/9A mice, however, in addition to the α1 and α3 subunits, the α4 and α5 subunits are significantly enriched in α1-containing GABA_ARs (Figure 6B and Supplementary Table 1C). To understand how the presence of these additional α subunit variants affects the stoichiometry of total α subunit isoforms, we assessed the contribution of each α subunit isofrom to the total amount of α subunits detected in our purifications from WT and S408/9A animals. In WT, the α1 subunit makes up 88.28% of all α subunits, followed by 3.82% of the α2 subunit and 7.90% of the α3 subunit (Figure 6C). With the S408/9A mutation, however, the contribution of the α1 subunit to the total α subunits decreases to 70.05%, while that of the α3 subunit increases to 15.62% (Figure 6C). The α4 and α5 subunits constitute 7.81% and 6.52% of all α subunits detected, respectively (Figure 6C).

Considering that the conventional stoichiometry of a GABA_AR is 2α2β3γ1 extra subunit (Wisden et al., 1992; Baumann et al., 2003), the presence of the α4 and α5 subunits in α1-containing GABA_ARs implies significant changes to the receptor population in S408/9A animals. Taken together, our experimental results demonstrate that receptors that contain both the α1 and α4 subunits are formed in S408/9A mice, suggesting a critical role of S408/9 in the segregation of the two α subunits into distinct receptor subtypes.

We also examined the contents of the 720 kDa band detected with the α1 subunit antibody with LC-MS/MS and quantitative analysis to reveal the significantly enriched proteins of the α1 subunit in S408/9A mice. Compared to 93 in WT, only 38 proteins were significantly enriched with α1-containing GABA_ARs in S408/9A mice (Supplementary Table 2C). The most abundantly found proteins with the α1 subunit in S408/9A mice are still Sptan1, Sptbn1, and Sptbn2 (Supplementary Figures 2A,B and Supplementary Table 2C). In addition, gephyrin was detected with the α1 subunit (Supplementary Table 2C). Lastly, the mutation did not alter the efficacy of the purification of α1-containing GABA_ARs (Supplementary Figure 2C). Together, these suggest that the S408/9A mutation does not alter the high-abundance binding partners of α1-containing GABA_ARs or disrupt binding to core cytoskeletal proteins that regulate the accumulation of this receptor subtype at synapses.

The subunit composition of α4-containing GABA_ARs is not affected by the S408/9A mutation

We also analyzed the subunit composition of α4-containing GABA_ARs in S408/9A mice. When purified GABA_ARs that
Mutation of S408/9 leads to the formation of α1α4-containing GABA\(_A\)Rs. (A) Purified PM from S408/9A mice were exposed to non-immune control IgG or the α1 subunit antibody and probed for the α1 subunit. Major bands at 250 and 720 kDa were observed. (B) Quantitative analysis of the contents of the 250 kDa band revealed that the α4 and α5 subunits are significantly enriched in α1-containing GABA\(_A\)Rs in S408/9A mice, along with the α1, α3, β1–3, γ2, and δ subunits (*\(p < 0.05\), **\(p < 0.01\), ***\(p < 0.001\), ****\(p < 0.0001\), \(n = 5\) replicates). (C) Contribution of each α subunit variant to the total amount of α subunits is compared between α1-containing GABA\(_A\)Rs from WT and S408/9A mice. With the S408/9A mutation, the percentages of the α1 and α2 subunits decrease whereas those of the α3, α4, and α5 subunits increase. (D) The α4 subunit antibody recognized a major band at 250 kDa in purified α4-containing GABA\(_A\)Rs from S408/9A mice. (E) In α4-containing GABA\(_A\)Rs from S408/9A mice, only the α4 and β3 subunits are significantly enriched (**\(p < 0.01\), \(n = 5\) replicates). (F) The S408/9A mutation does not affect the efficacy of the purification of either α1- or α4-containing GABA\(_A\)Rs (ns ≥ 0.05, \(n = 5\) replicates).

To ensure that the S408/9A mutation does not interfere with the efficacy of the purification of GABA\(_A\)Rs that contain either the α1 or α4 subunit, we compared the target subunit recovery in each purification between the two genotypes. We found that there is no significant difference in the amount
of the α1 (WT: 0.0087 ± 0.0009, S408/9A: 0.0067 ± 0.0022, p = 0.4237, n = 5, unpaired t-test) or α4 (WT: 0.0046 ± 0.0011, S408/9A: 0.0063 ± 0.0020, p = 0.3158, n = 5, unpaired t-test) subunit detected in the 250 kDa band between WT and S408/9A animals (Figure 6F). Collectively, these results suggest that the S408/9A mutation does not alter the subunit composition of α4-containing GABA_ARs.

Analysis of the 720 kDa band revealed that α4-containing GABA_ARs in S408/9A mice interact with only 9 proteins, none of which were known inhibitory scaffolding proteins (Supplementary Figures 2D,E and Supplementary Table 2D). In addition, there was no overlap between binding proteins of the α4 subtype between WT and S408/9A mice (Figure 3, Supplementary Figures 2D,E, and Supplementary Tables 2B,D). The mutation had no effect on the efficacy of α4-containing GABA_AR purification (Supplementary Figure 2F). Together, these results suggest the lack of proteins that specifically anchor the α4 subunit in the membrane.

The S408/9A mutation does not induce gross changes in the basal phosphorylation of the β3 subunit

Next, we assessed if mutation of S408/9 induced aberrant phosphorylation on other residues within the β3 subunit in either receptor subtype using LC-MS/MS. We found that in α1-containing GABA_ARs, the same phosphorylation sites that are present in WT were also detected in S408/9A mice: T322, S330, S332, S383, Y391, S396, T402, and T419 (Figure 7A). Two sites were more significantly phosphorylated in α1-containing GABA_ARs in S408/9A mice than in WT: T322 (WT: 0.54 ± 0.01, S408/9A: 0.63 ± 0.04, p = 0.0464, n = 5, unpaired t-test) and T419 (WT: 0.40 ± 0.03, S408/9A: 0.05 ± 0.01, p = 0.0380, n = 5, unpaired t-test) (Figure 7A). The same eight phosphorylation sites were detected in the α4 subtype in S408/9A mice as well (Figure 7B). Together, these results demonstrate that the mutation of S408/9 does not induce gross changes in the sites of basal phosphorylation within the β3 subunit in either the α1 or α4 receptor subtype. These results also suggest that the presence of α1α4-containing GABA_ARs in S408/9A mice is specifically due to the S408/9A mutation.

The endoplasmic reticulum retention of the α4 subunit is decreased in S408/9A mice

Our results have demonstrated that the S408/9A mutation leads to altered subunit composition of α1-containing GABA_ARs in the forebrain (Figures 6B,C). The assembly of GABA_ARs from their component subunits occurs within the ER, a process that acts to limit the structural diversity of receptor subtypes that accumulate in the PM (Jacob et al., 2008). To understand the effects of the S408/9A mutation further, we investigated the expression of GABA_AR subunits between subcellular compartments. We first subjected homogenized murine forebrain tissues to

![Figure 7](https://example.com/figure7.jpg)

**Figure 7**

The S408/9A mutation does not lead to gross changes in global β3 subunit phosphorylation. (A) Phosphorylated residues within the β3 subunit of α1-containing GABA_ARs are compared between WT and S408/9A mice. The mutation does not induce aberrant phosphorylation on the β3 subunit, as indicated by the detection of the same phosphorylation sites in WT and S408/9A mice. Two sites – T322 and T419 – are more highly phosphorylated in S408/9A than in WT mice (*p < 0.05, n = 5 replicates). (B) Phosphorylated residues within the β3 subunit of α4-containing GABA_ARs from WT and S408/9A mice are shown. Phosphorylated S332 was only detected in the α4 subtype from S408/9A mice (n = 5 replicates).
serial centrifugation from which cytoplasm, mitochondria, ER, and PM fractions were obtained (Suski et al., 2014; Smalley et al., 2020). These fractions were immunoblotted for marker proteins specific for each subcellular compartment. Enrichment of hsp90 in cytoplasmic fraction, hsp60 in mitochondria, calreticulin in ER, and n-cadherin in the PM (Supplementary Figure 3) validated our method of subcellular fractionation.

We then subjected purified total lysate (TL), PM, and ER fractions to SDS-PAGE and probed for n-cadherin and the α1, α4, and β3 subunits. The distribution of each GABA A R subunit between the PM and ER was calculated as a percentage of total expression in TL, PM, and ER. In WT mice, the α1 subunit immunoreactivity was significantly higher in the PM than in the ER (Figures 8A,B; PM: 46.57 ± 3.63%, ER: 17.43 ± 3.26%, p = 0.0040, n = 3, unpaired t-test). In contrast, similar amounts of the α4 subunit were found in the PM and the ER (Figures 8A,C; PM: 36.81 ± 3.94%, ER: 32.36 ± 5.90%, p = 0.5641, n = 3, unpaired t-test). Similarly to the α1 subunit, the majority of the β3 subunit immunoreactivity was found in the PM (Figures 8A,D; PM: 57.16 ± 1.94%, ER: 22.79 ± 4.40%, p = 0.0020, n = 3, unpaired t-test). In subcellular fractions prepared from S408/9A mice, all three subunits were more significantly expressed in the PM than in the ER: α1 (PM: 62.18 ± 0.82%, ER: 11.11 ± 1.81%, p < 0.0001, n = 3, unpaired t-test), α4 (PM: 60.30 ± 0.78%, ER: 23.56 ± 2.82%, p = 0.0002, n = 3, unpaired t-test), and β3 subunit (PM: 40.51 ± 5.39, ER: 18.93 ± 1.80, p = 0.0191, n = 3, unpaired t-test) (Figures 8E–H).

To examine the effects of the S408/9A mutation on ER accumulation of GABA A Rs subunits further, we prepared primary cultured mixed cortical and hippocampal neurons from WT and S408/9A mice. We used immunocytochemistry to visualize the α4 subunit with calreticulin around the soma (Figure 9A). We found that there is no significant difference in the total area of the α4 subunit puncta (WT: 25.83 ± 3.67 pixel², S408/9A: 25.02 ± 2.59 pixel², p = 0.8577, n = 18, unpaired t-test) or calreticulin alone (WT: 24.74 ± 3.17 pixel², S408/9A: 22.93 ± 2.63 pixel², p = 0.6630, n = 18, unpaired t-test) between the two genotypes (Figures 9B,C). However, the total area of the α4 subunit and calreticulin co-localization is significantly reduced in S408/9A cells (Figure 9D; WT: 40.57 ± 5.56 pixel², S408/9A: 24.84 ± 4.54 pixel², p = 0.0352, n = 18, unpaired t-test). This is consistent with the results from the immunoblot of the PM and ER fractions (Figure 8) and suggests that there is less accumulation of the α4 subunit in the ER in S408/9A mice than in WT mice.

Discussion

In the forebrain, phasic inhibition is mediated by GABA A Rs that contain the α1 subunit whereas tonic inhibition is dependent upon those assembled from the α4 subunit, and these populations of GABA A Rs are often co-expressed in neurons (Fritschy et al., 1998; Pirker et al., 2000). Here, we have begun to evaluate how the endogenous α1 and α4 subunits are segregated into distinct receptor subtypes. To do so, we have developed methods to isolate native α1- and α4-containing GABA A Rs from the PM and evaluated their subunit composition and binding proteins using BN-PAGE coupled with quantitative mass spectrometry.

Our approach revealed that affinity purified GABA A Rs containing the α1 subunit exhibited a native molecular mass of 250 kDa, which is consistent with the pentameric structure of GABA A Rs. Quantitative analysis of the 250 kDa species showed that α1-containing receptors contain the α1–3, β1–3, γ2–3, and δ subunits but not the α4 or α5 subunit. Based on the relative abundance, the subunit composition of endogenous α1-containing GABA A Rs was determined to be α1β3γ2α, which has previously been found to be the most commonly encountered subunit composition in the brain (Olsen and Sieghart, 2009). Out of the α subunit isoforms, only the α2 and α3 subunits were significantly enriched with the α1 subunit when compared to control. This is consistent with previous studies that suggest the α1–3 subunits as synaptic α subunit variants (Brüning et al., 2002). Compared to the α1 subunit, lower levels of the α2 and α3 subunits were detected. Thus, the majority of synaptic GABA A Rs are likely to contain one or more copies of the α1 subunit. Such preferential assembly may result from the α1 subunit being expressed at 5–20-fold higher levels than other α subunit variants in the rodent forebrain (Chen et al., 2012). Notably, the δ subunit was detected as a significant binding partner of the α1 subunit in our purifications. While this result was unexpected, this is consistent with earlier studies suggesting that the δ subunit co-purifies with the α1, α3, β2–3, and γ2 subunits (Mertens et al., 1993). Moreover, receptors containing both the α1 and δ subunits have been suggested to be principal mediators of tonic inhibition in hippocampal interneurons (Fritschy and Mohler, 1995; Glykys et al., 2007).

The subunit composition of GABA A Rs assembled from the α4 subunit was explored using a similar approach involving affinity purification with the α4 subunit antibody and BN-PAGE. These receptors also exhibited a native molecular mass of 250 kDa. Quantitative analysis revealed the significant presence of only the α4 and β3 subunits in this 250 kDa species. No other α, β, or γ subunit variants or the δ subunit was significantly enriched. While a major limitation of our experimental approach is that we purify GABA A Rs en masse and thus low-abundance species may be overlooked, GABA A Rs composed of the α and β subunits only have been shown to mediate tonic inhibition in hippocampal pyramidal neurons (Mortensen and Smart, 2006). In fact, it is estimated that up to 10% of extrasynaptic GABA A Rs in hippocampal pyramidal neurons are αβ-containing...
The S408/9A mutation selectively affects the distribution of the α4 subunit across subcellular compartments. (A) Immunoblots of total forebrain lysate (TL), PM, and ER fractions from WT animals were probed for N-cadh, α1, α4, and β3 subunits. (B) The distribution of each GABA_A R subunit between PM and ER was calculated as a percentage of total expression in TL, PM, and ER. Higher percentage of the α1 subunit is expressed in the PM than in the ER (**p < 0.01, n = 3). (C) Similar levels of the α4 subunit are observed in the PM and ER (ns ≥ 0.05, n = 3). (D) The β3 subunit is more significantly enriched in the PM than in the ER (***p < 0.001, n = 3). (E) Levels of N-Cadh, α1, α4, and β3 subunits were assessed in immunoblots of TL, PM, and ER fractions from S408/9A animals. (F–H) In S408/9A animals, higher levels of the α1, α4, and β3 subunits are expressed in the PM than in the ER (*p < 0.05, ***p < 0.001, ****p < 0.0001, n = 3).

FIGURE 8

The S408/9A mutation selectively affects the distribution of the α4 subunit across subcellular compartments. (A) Immunoblots of total forebrain lysate (TL), PM, and ER fractions from WT animals were probed for N-cadh, α1, α4, and β3 subunits. (B) The distribution of each GABA_A R subunit between PM and ER was calculated as a percentage of total expression in TL, PM, and ER. Higher percentage of the α1 subunit is expressed in the PM than in the ER (**p < 0.01, n = 3). (C) Similar levels of the α4 subunit are observed in the PM and ER (ns ≥ 0.05, n = 3). (D) The β3 subunit is more significantly enriched in the PM than in the ER (***p < 0.001, n = 3). (E) Levels of N-Cadh, α1, α4, and β3 subunits were assessed in immunoblots of TL, PM, and ER fractions from S408/9A animals. (F–H) In S408/9A animals, higher levels of the α1, α4, and β3 subunits are expressed in the PM than in the ER (*p < 0.05, ***p < 0.001, ****p < 0.0001, n = 3).

receptors (Mortensen and Smart, 2006). Likewise, studies using immunoprecipitation coupled with immunoblotting suggest that more than 50% of α4-containing GABA_A Rs in the brain are composed of the α and β subunits only (Bencsits et al., 1999).

The current dogma states that α4βδ is the most common subunit combination for GABA_A Rs mediating tonic inhibition (Nusser and Mody, 2002; Wei et al., 2003; Peng et al., 2004; Jia et al., 2005). However, to our surprise, we did not detect a significant enrichment of the δ subunit in our
Colocalization of the α4 subunit with calreticulin is reduced in cultured neurons from S408/9A mice. (A) Representative images of the soma of DIV21 neurons cultured from WT and S408/9A mice stained with the α4 subunit and calreticulin antibodies are shown. Individual channels and areas of colocalization for each cell are shown on the right. (B) Total area of the α4 subunit staining in the soma is not affected by the S408/9A mutation (ns ≥ 0.05, n = 18 cells from three individually prepared cultures). (C) There is no significant difference in calreticulin staining in the soma between WT and S408/9A cultures (ns ≥ 0.05, n = 18 cells from three individually prepared cultures). (D) Colocalized area of the α4 subunit with calreticulin is reduced in S408/9A cultures when compared to WT cultures (*p ≤ 0.05, n = 18 cells from three individually prepared cultures).

Purifications of α4-containing GABA_A Rs. While neurons may assemble this receptor subtype in distinct brain regions such as the thalamus and DGCCs (Sur et al., 1999; Wei et al., 2003), our results suggest that α4β3 is the dominant subunit composition of α4-containing GABA_A Rs in the adult mouse forebrain. This is consistent with a previous study that showed that only 7% of the α4-containing GABA_A Rs contain the δ subunit (Bencsits et al., 1999). Given the sensitivity of δ-containing GABA_A Rs to modulation by neurosteroids (Stell et al., 2003), the expression of α4δ-containing GABA_A Rs may be induced by these endogenous modulators of GABAergic inhibition.

In addition to the 250 kDa species, a larger species of 720 kDa was seen in the purification of α1-containing GABA_A Rs.
from the PM. Analysis of this larger species revealed that the α1 subunit co-purifies with 93 proteins, with spectrin isoforms (Sptan1, Spbn1, and Spbn2) making up more than 70% of the stable multiprotein complex. Spectrins are cytoskeletal integrators that bind to actin, microtubules, and intermediate filaments to provide structural support in cells (Liem, 2016). They are heterodimeric proteins that contain one α and one β isoforms – most spectrin molecules found in the brain are composed of αII-spectrin (encoded by Sptan1) and βII-spectrin (encoded by Spbn1) (Machnicka et al., 2014; Liem, 2016). With each spectrin isoform at around 240 kDa, the high molecular weight species of α1-containing GABA Rs at around 720 kDa is expected to be the pentameric GABA Rs with dimeric spectrins. In neurons, spectrins form structures with ankyrins (Ank2/3), actin (Actn1), and adducin (Add1/3) (Xu et al., 2013; Han et al., 2017), all of which are significantly detected with the α1 subunit in low abundance. Together these proteins anchor cell surface transmembrane proteins to the cytoskeleton (Xu et al., 2013; Han et al., 2017).

Among the low-abundance proteins co-purifying with the α1 subtype is Kcc2 (encoded by Slc12a5), which is a neuronal membrane protein whose expression and function are critical for synaptic inhibitory transmission (Moore et al., 2017). The immunoblot of Kcc2 showed a band between 242 and 480 kDa for synaptic inhibitory transmission (Moore et al., 2017). The expression and function in neurons have not been determined, however. Another binding protein of the α4 subunit is desmoplakin, which interacts with n-cadherin, a cell adhesion protein in the membrane (Tanaka et al., 2012). Interestingly, desmoplakin was found to associate with the γ2 subunit as well (Ge et al., 2018). For the α4 subunit, the relative abundance of each protein is comparable, suggesting that the 720 kDa species would be a summation of pentameric α4-containing GABA Rs with these low-abundance proteins. We also noted that neither gephyrin nor spectrin isoforms co-purified with the α4 subunit. However, two proteins were detected with both the α1 and α4 subunits in our findings: Myo5a and Vdac2. Myo5a has been implicated in subcellular localization of gephyrin (Fuhrmann et al., 2002) and Vdac2 has been found to associate with GABA Rs (Bureau et al., 1992; Darbandi-Tonkabon et al., 2004).

Collectively, our experiments suggest that at steady state, neurons segregate the α1 and α4 subunits into biochemically distinct receptor subtypes. Moreover, analysis of the binding proteins of the two receptor subtypes suggests that GABA Rs containing the α1 subunit are associated with core components of the postsynaptic inhibitory scaffold, which is consistent with their role in mediating phasic inhibition. In contrast, those assembled from the α4 subunit are not associated with inhibitory scaffolding proteins. The significant difference in the components of stable protein complexes between the two GABA R subtypes further strengthens the suggestion that the α1 and α4 subunits are sorted into separate populations.

Interestingly, the β3 subunit is significantly enriched in both α1 and α4 subtypes. Furthermore, the β3 subunit associated with the α1 subunit exhibits higher phosphorylation of S408/9 than that in the α4 subtype. Previous studies have shown that S408/9 phosphorylation acts to modify membrane trafficking of GABA Rs and promotes their residence time in the PM (Kittler et al., 2005; Vien et al., 2015). Moreover, aberrant levels of S408/9 phosphorylation have been observed in mouse models of status epilepticus and fragile X syndrome (FXS) (Terunuma et al., 2008; Vien et al., 2015). We found that disrupting the phospho-regulation of S408/9 compromises the segregation of the α1 and α4 subunits into distinct receptor subtypes, leading to the production of α1α4-containing GABA Rs. In addition to the α4 subunit, the amount of the α5 subunit detected with the α1 subunit was significantly enhanced in the mutants. The presence of the α4 and α5 subunits altered the proportion of receptors occupied by the α1 subunit, suggesting that α1-containing proteins – most spectrin molecules found in the brain are composed of αII-spectrin (encoded by Sptan1) and βII-spectrin (encoded by Spbn1) (Machnicka et al., 2014; Liem, 2016). With each spectrin isoform at around 240 kDa, the high molecular weight species of α1-containing GABA Rs at around 720 kDa is expected to be the pentameric GABA Rs with dimeric spectrins. In neurons, spectrins form structures with ankyrins (Ank2/3), actin (Actn1), and adducin (Add1/3) (Xu et al., 2013; Han et al., 2017), all of which are significantly detected with the α1 subunit in low abundance. Together these proteins anchor cell surface transmembrane proteins to the cytoskeleton (Xu et al., 2013; Han et al., 2017).

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Recently, two proteins - Lhfp4 and Shisa7 - have been reported to bind to GABA Rs and regulate their membrane trafficking (Davenport et al., 2017; Han et al., 2019; Yamasaki et al., 2017). However, we did not detect significant levels of either protein co-purifying with the α1 subunit. Our failure to detect these proteins may reflect that we utilized purified plasma membranes as a starting material while those on Lhfp4 and Shisa7 used brain extracts or synaptosomal fractions. Alternatively, differences between the extraction buffers and/or detergents employed may also be of significance.

Interestingly, we found that purified α4-containing GABA Rs also form a stable multiprotein complex at around 720 kDa and that it contains 19 proteins. Among the proteins that were significantly enriched with the α4 subunit, the most abundant is titin, which is known primarily as a protein that provides elasticity in muscles (Freundt and Linke, 2019). Its function and expression in neurons have not been determined, however. Another binding protein of the α4 subunit is desmoplakin, which interacts with n-cadherin, a cell adhesion protein in the membrane (Tanaka et al., 2012). Interestingly, desmoplakin was found to associate with the γ2 subunit as well (Ge et al., 2018). For the α4 subunit, the relative abundance of each protein is comparable, suggesting that the 720 kDa species would be a summation of pentameric α4-containing GABA Rs with these low-abundance proteins.

Collectively, our experiments suggest that at steady state, neurons segregate the α1 and α4 subunits into biochemically distinct receptor subtypes. Moreover, analysis of the binding proteins of the two receptor subtypes suggests that GABA Rs containing the α1 subunit are associated with core components of the postsynaptic inhibitory scaffold, which is consistent with their role in mediating phasic inhibition. In contrast, those assembled from the α4 subunit are not associated with inhibitory scaffolding proteins. The significant difference in the components of stable protein complexes between the two GABA R subtypes further strengthens the suggestion that the α1 and α4 subunits are sorted into separate populations.

Interestingly, the β3 subunit is significantly enriched in both α1 and α4 subtypes. Furthermore, the β3 subunit associated with the α1 subunit exhibits higher phosphorylation of S408/9 than that in the α4 subtype. Previous studies have shown that S408/9 phosphorylation acts to modify membrane trafficking of GABA Rs and promotes their residence time in the PM (Kittler et al., 2005; Vien et al., 2015). Moreover, aberrant levels of S408/9 phosphorylation have been observed in mouse models of status epilepticus and fragile X syndrome (FXS) (Terunuma et al., 2008; Vien et al., 2015). We found that disrupting the phospho-regulation of S408/9 compromises the segregation of the α1 and α4 subunits into distinct receptor subtypes, leading to the production of α1α4-containing GABA Rs. In addition to the α4 subunit, the amount of the α5 subunit detected with the α1 subunit was significantly enhanced in the mutants. The presence of the α4 and α5 subunits altered the proportion of receptors occupied by the α1 subunit, suggesting that α1-containing
GABA\(_{A}\)Rs in S408/9A mice may only have one copy of the \(\alpha1\) subunit. Based on the relative abundance of subunits detected, \(\alpha1\beta2/3\beta2\) is still likely the subunit composition of \(\alpha1\)-containing GABA\(_{A}\)Rs in the mutant mice. Functional studies have shown that the spontaneous amplitude of inhibitory synaptic currents is increased in S408/9A mice, while tonic currents are decreased (Vien et al., 2015). Thus, the presence of \(\alpha1\alpha4\) containing receptors in S408/9A mice may contribute to these reciprocal modifications in phasic and tonic inhibition.

For the binding proteins, the mutation did not interfere with the interactions between \(\alpha1\)-containing GABA\(_{A}\)Rs and spectrin isoforms (Sptan1, Sptbn1, and Sptbn2), Kcc2, or gephyrin. This suggests that the stabilization and synaptic targeting of the \(\alpha1\) subunit are still intact even with the presence of the \(\alpha4\) and \(\alpha5\) subunits. It also suggests that the novel \(\alpha1\alpha4\)-containing GABA\(_{A}\)Rs may be localized to the synaptic areas. In an animal model of FXS, in which the level of S408/9 phosphorylation is significantly altered compared to healthy mice, the \(\alpha4\) subunit selective agonist Ro15-4513 potentiates synaptic currents (Vien et al., 2015; Modgil et al., 2019). In addition, the \(\alpha4\) subunit is found in synaptic areas (Zhang et al., 2017), although the molecular mechanism for the mis- trafficking has not been revealed. It is possible that aberrant S408/9 phosphorylation leads to the formation of \(\alpha1\alpha4\)-containing GABA\(_{A}\)Rs and as a result, the \(\alpha4\) subunit is mis-trafficked to synapses.

While the S408/9A mutation did not modify the total level of the \(\alpha1\), \(\alpha4\), or \(\beta3\) subunit and did not induce any gross changes in the phosphorylation of the \(\beta3\) subunit, it did reduce the levels of the \(\alpha4\) subunit in the ER and increase its accumulation in the PM. The steady state levels of GABA\(_{A}\)R subunits within the ER are dependent upon oligomerization into transport-competent hetero-oligomers, as well as their rates of retrograde transport and subsequent proteasomal degradation (Jacob et al., 2008; Luscher et al., 2011). Thus, S408/9A are likely to induce the assembly of the \(\alpha4\) subunit into hetero-oligomers, promoting their subsequent transport from the ER to PM and contributing to changes in GABAergic regulation (Vien et al., 2015). Our study also identified several novel phosphorylation sites on the intracellular loop of the \(\beta3\) subunit - T322, S330, S332, Y391, S396, T402, and T419. Phosphorylated S332 was only detected on the \(\beta3\) subunit that is associated with the \(\alpha1\) subunit in WT animals. However, it was detected in both \(\alpha1\) and \(\alpha4\) subtypes in S408/9A animals. Additionally, T322 and T419 were more significantly phosphorylated in the \(\alpha1\) subtype in the mutants than in WT. Further investigation into these novel phosphorylation sites may provide additional insights into mechanisms of GABA\(_{A}\)R regulation.

In summary, our results have provided insights into the mechanisms neurons utilize to orchestrate the assembly of GABA\(_{A}\)R subunits into distinct receptor subtypes, which is required to support phasic and tonic inhibition. They also suggest that receptor assembly is subject to dynamic modulation via S408/9 in the \(\beta3\) subunit, which are key sites for phosphorylation-dependent regulation of surface GABA\(_{A}\)R expression. How S408/9 phosphorylation regulates the segregation of the \(\alpha1\) and \(\alpha4\) subunits into distinct receptor subtypes is an open question, but our results suggest that this process may have an impact on the \(\beta3\) subunit’s half-life in the ER, a key determinant for subunit oligomerization, ER exit, and transport to the PM (Nakamura et al., 2016). Thus, cell signaling molecules that regulate S408/9 phosphorylation, such as neurosteroids (Adams et al., 2015; Modgil et al., 2017, 2019; Parakala et al., 2019), may exert long lasting effects on neuronal activity by regulating GABA\(_{A}\)R assembly.

**Materials and methods**

**Animals**

WT C57BL/6 and S408/9A animals were kept in a temperature-controlled room on a 12-h light/dark cycle and fed ad libitum with 2 cage changes per week. S408/9A animals were generated by gene targeting in murine ES cells as previously described (Vien et al., 2015, 2022). 8–12 weeks old male and female S408/9A homozygous mice and WT littermate controls were used for all biochemistry experiments. For immunocytochemistry experiments, P0 WT and S408/9A homozygous pups were used. All experimental procedures were approved by the Tufts University Institutional Animal Care and Use Committee (IACUC).

**Antibodies**

The following antibodies were used for immunoprecipitation (IP), western blot (WB), or immunocytochemistry (ICC): Gabra1 (mouse, NeuroMab clone N95/35, IP), Gabra1 (rabbit, Abcam ab33299, WB 1:1000, ICC 1:1000), Gabra4 (mouse, NeuroMab clone N398A/34, IP, ICC 1:500), Gabra4 (rabbit, PhosphoSolutions 845A-GA4C, WB 1:1000), Gabrb3 (rabbit, PhosphoSolutions 863A-GB3C, WB 1:1000), p-S408/9 (PhosphoSolutions p1130–4089, WB 1:1000), Sptan1 (rabbit, Cell Signaling 2122S, WB 1:1000), Sptbn1 (rabbit, Abcam ab72239, WB 1:1000), Sptbn2 (rabbit, Proteintech 55107-1-AP, WB 1:1000), Gephyrin (rabbit, Cell Signaling 14304S, WB 1:1000), N-Cadherin (mouse, Cell Signaling 14215S, WB 1:1000), Calreticulin (rabbit, Cell Signaling 12238S, WB 1:1000, ICC 1:500), Hsp60 (rabbit, Cell Signaling 12165S, WB 1:1000), Hsp90 (rabbit, Cell Signaling 4877S, WB 1:1000), GAPDH (mouse, Santa Cruz sc-32233, WB 1:5000), Goat anti-mouse Alexa Fluor 488 (Thermo Fisher A11029, ICC 1:1000), Goat anti-rabbit Alexa Fluor 568 (Thermo Fisher A11011, ICC 1:1000), Donkey anti-mouse conjugated HRP.
Overnight at 4°C. The beads were resuspended in 0.05% PBS-T and incubated three times with 1× PBS with 0.05% Tween-20 (v/v; 0.05% PBS-TBS-T, and incubated with primary antibodies diluted in TBS-T overnight with colloidal Coomassie blue. For liquid chromatography coupled with tandem mass spectrometry (LC-MS/MS), the gels were fixed in 4% paraformaldehyde in PBS (v/v) for 10 min at room temperature. After washing with TBS-T, the membranes were incubated with secondary antibodies diluted in TBS-T for 1 h at room temperature. Protein bands were visualized with SuperSignal West Dura Extended Duration Substrate (Thermo Scientific 34075) and imaged using a ChemiDoc MP (Bio-Rad). Images were cropped at the bottom (below 150 kDa) to remove the dye front, which contains unbound colloidal Coomassie blue. For SDS-PAGE, proteins were transferred to nitrocellulose membranes overnight at 4°C. The membranes were blocked in 5% milk or BSA, incubated with primary and secondary antibodies, and imaged as described above (Smalley et al., 2020; Kontou et al., 2021).

### Primary neuronal culture

P0 pups were anesthetized on ice and immediately euthanized by decapitation. The brains were quickly removed, and cortical and hippocampal tissues were dissected in ice-cold Hank’s buffered salt solution (HBSS; Thermo Fisher 14185052) with 10 mM HEPES. The tissues were trypsinized and triturated in Neurobasal-A media (Thermo Fisher 10888022) containing 2% B27 (v/v; Thermo Fisher 17504044), 1% GlutaMAX (v/v; Thermo Fisher 35050061), and 1% Penicillin/Streptomycin (v/v; Thermo Fisher 15140122). Dissociated neurons were filtered through a 40 μm nylon mesh strainer (Thermo Fisher 22363547) and counted using a hemocytometer. Cells were plated on poly-L-lysine (PLL)-coated 13 mm coverslips in 24-well plates at a density of 100–125 K cells/well in media. At days in vitro (DIV) 21, cells were fixed in 4% paraformaldehyde in PBS (v/v) for 10 min at room temperature.
temperature and placed in PBS at 4°C (Kontou et al., 2021).

**Immunocytochemistry**

Fixed cultures were permeabilized for 1 h in blocking solution [3% BSA (w/v), 10% normal goat serum (v/v), 0.2 M glycine in PBS, 0.1% Triton-X100 (v/v)] (Kontou et al., 2021). Coverslips were incubated with primary antibodies diluted in blocking solution for 1 h at room temperature. After a brief wash with PBS, the coverslips were incubated with fluorophore-conjugated secondary antibodies diluted in blocking solution for 1 h at room temperature. The coverslips were then washed in PBS, briefly dipped in DAPI ready-made solution (Sigma-Aldrich MBD0015), and mounted onto microscope slides with Fluoromount-G (SouthernBiotech 0100-01). The coverslips were imaged using a Nikon A1 confocal microscope (Nikon Instruments, Melville, NY, USA) using a 60× oil immersion objective lens. Laser settings were manually assigned for each fluorescent channel and images were acquired at the scan size of 1,024 × 1,024. Settings were kept the same between imaging sessions and between genotypes.

**Image analysis**

For immunoblots, individual band intensity was quantified using densitometry on ImageJ (Version 1.53) normalized to GAPDH and further normalized to the corresponding control condition where appropriate. For immunocytochemistry, 18 neurons from 3 separate cultures per genotype were analyzed on FIJI (Version 2.3.0). The background was subtracted using the rolling ball radius of 50, and the images were smoothed with the median filter of 1.0. The total areas of individual protein staining were quantified using Analyze Particle with the range of the median filter of 1.0. The total areas of colocalization were calculated using ImageJ (Version 1.53). Laser settings were manually assigned for each fluorescent channel and images were acquired at the scan size of 1,024 × 1,024. Settings were kept the same between imaging sessions and between genotypes.

**LC-MS/MS analysis**

Peptide sequences were determined by matching protein or translated nucleotide databases with the acquired fragmentation pattern using the MSGF+ (Kim and Pevzner, 2014). Raw mzXML files were used to search the UniProt mouse reference proteome (last modified May 4th 2020, containing 21,989 sequences), which contains the Thermo list of common contaminants. For phosphopeptide detection, a variable modification of mass units to serine, threonine, and tyrosine was included in the database searches. For LC-MS/MS analysis, significant proteins were detected by performing the Welch's t-test to identify significantly enriched proteins. For network analysis, the protein lists were compared against the latest version of the STRINGdb database (Szklarczyk et al., 2019). The interaction for each protein with Gabra1 or Gabra4 immunoprecipitation were compared to non-immune IgG purifications by Welch’s t-test to identify significantly enriched proteins. For network analysis, the protein lists were compared against the latest version of the STRINGdb database (Szklarczyk et al., 2019). The interaction for each protein with Gabra1 or Gabra4 was imputed and network diagrams were constructed in R using the igraph package (accessed February 1st, 2019) and the nodes were scaled to the average SI values for each protein. The highest scoring Gene Ontology Biological Process term was extracted for each protein using the mygene package (accessed July 29th, 2020). For PCA, The SI values for proteins contained within each gel band were normalized by z-transformation using the ggfortify package in R. Only the significantly enriched proteins detected in all replicates were included in the analysis.

**Statistics**

For LC-MS/MS data, significant proteins were detected by performing the Welch’s t-test between the samples and non-immune IgG purifications. Significance on immunoblots and
ICC data was determined by performing the unpaired Student’s t-test on GraphPad Prism (Version 9.3.1). Varying levels of significance were denoted as the following: ns (p ≥ 0.05), * (p < 0.05), ** (p < 0.01), *** (p < 0.001), **** (p < 0.0001).

Data availability statement

The datasets presented in this study can be found in online repositories. The names of the repository/repositories and accession number(s) can be found below: http://www.proteomexchange.org/, PXD036295.

Ethics statement

The animal study was reviewed and approved by Tufts University Institutional Animal Care and Use Committee (IACUC).

Author contributions

CC performed the experiments, analyzed the data, and wrote the manuscript. JS, AL, and QR performed the experiments. CB and JD maintained the mouse colony and performed genotyping. SM and JS conceptualized the project and designed the experiments. CC, JS, PD, and SM edited the manuscript. All authors contributed to the article and approved the submitted version.

Funding

This work was supported by the National Institutes of Health (NIH) – National Institute of Neurological Disorders and Stroke Grant NS108378 (SM and PD) and National Institute of Mental Health Grants MH118263 (SM) and MH126542 (CC).

Acknowledgments

We would like to thank the Taplin Mass Spectrometry Facility at Harvard Medical School.

Conflict of interest

Author SM serves as a consultant for AstraZeneca and SAGE Therapeutics, relationships that are regulated by Tufts University. Author SM holds stock in SAGE Therapeutics.

The remaining authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

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Supplementary material

The Supplementary Material for this article can be found online at: https://www.frontiersin.org/articles/10.3389/fnmol.2022.1017404/full#supplementary-material

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