General Anesthetic Binding Sites in Human $\alpha_{4}\beta_{3}\delta$
\gamma-Aminobutyric Acid Type A Receptors (GABA$\alpha$Rs)*

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Extrasynaptic $\gamma$-aminobutyric acid type A receptors (GABA$\alpha$Rs), which contribute generalized inhibitory tone to the mammalian brain, are major targets for general anesthetics. To identify anesthetic binding sites in an extrasynaptic GABA$\alpha$Rs, we photolabeled human $\alpha_{4}\beta_{3}\delta$ GABA$\alpha$Rs purified in detergent with $[3^H]$azietomidate and a barbiturate, $[3^H]R$-m-tTFD-MPAB, photoactive anesthetics that bind with high selectivity to distinct but homologous intersubunit binding sites in the transmembrane domain of synaptic $\alpha_{1}\beta_{3}\gamma_{2}$ GABA$\alpha$Rs. Based upon $^3$H incorporation into receptor subunits resolved by SDS-PAGE, there was etomidate-inhibitable labeling by $[3^H]$azietomidate in the $\alpha$ and $\beta$ subunits and barbiturate-inhibitable labeling by $[3^H]R$-m-tTFD-MPAB in the $\beta$ subunit. These sites did not bind the anesthetic steroid alphaxalone, which enhanced photolabeling, or DS-2, a $\delta$ subunit-selective positive allosteric modulator, which neither enhanced nor inhibited photolabeling. The amino acids labeled by $[3^H]$azietomidate or $[3^H]R$-m-tTFD-MPAB were identified by N-terminal sequencing of fragments isolated by HPLC fractionation of enzymatically digested subunits. No evidence was found for a $\delta$ subunit contribution to an anesthetic binding site. $[3^H]$azietomidate photolabeling of $\beta$3Met-286 in $\beta$M3 and $\alpha$4Met-269 in $\alpha$M1 that was inhibited by etomidate but not by R-m-tTFD-MPAB established that etomidate binds to a site at the $\beta^{+}$-$\alpha^{-}$ interface equivalent to its site in $\alpha_{1}\beta_{3}\gamma_{2}$ GABA$\alpha$Rs. $[3^H]$Azietomidate and $[3^H]R$-m-tTFD-MPAB photolabeling of $\beta$3Met-227 in $\beta$M1 established that these anesthetics also bind to a homologous site, most likely at the $\beta^{+}$-$\beta^{-}$ interface, which suggests a subunit arrangement of $\beta_{3}\alpha_{4}\beta_{3}\delta$.

$\gamma$-Aminobutyric acid type A receptors (GABA$\alpha$Rs)$^2$ are the major inhibitory neurotransmitter receptors in the mammalian brain. They are members of the pentameric ligand-gated ion channel superfamily that consists of five homologous subunits, each of which has a large extracellular domain, a transmembrane domain of four transmembrane helices (M1–M4), and an intracellular domain connecting the third and fourth transmembrane helices. GABA$\alpha$Rs, which are the target of many drugs, among them benzodiazepines and general anesthetics, are heteropentamers, and drug action often depends on the subunit composition. For example, at synaptic receptors, which commonly have a subunit composition of $(\alpha)2(\beta)2\gamma$, arranged $\beta\alpha\gamma\beta$ counterclockwise when viewed from the synaptic or extracellular side of the receptor, benzodiazepines act in the extracellular domain between $\alpha^{+}$-$\gamma^{-}$ subunits at a site homologous to the GABA binding sites at the two $\beta^{+}$-$\alpha^{-}$ subunit interfaces (Fig. 1) (1–3).

General anesthetics have long been known to bind to sites in the transmembrane domains of pentameric ligand-gated ion channels (reviewed in Refs. 4–7). Photolabeling of endogenous and heterologous GABA$\alpha$Rs by $[3^H]$azietomidate located the etomidate binding site in the two $\beta^{+}$-$\alpha^{-}$ subunit interfaces (8, 9), 50 Å from the GABA site and at a position later shown to overlap with the five ivermectin sites in the crystal structure of the homopentameric glutamate-gated chloride channel (GluCl) (10). More recently, a photoactive, anesthetic barbiturate, $R$-m-tTFD-MPAB, has been shown to bind to sites in the $\gamma^{+}$-$\beta^{-}$ and $\alpha^{+}$-$\beta^{-}$ subunit interfaces homologous to the etomidate binding sites, introducing the concept of subtype-dependent action of general anesthetics (11). Whereas etomidate and $R$-m-tTFD-MPAB bind with high selectivity to their sites, propofol, pentobarbital, and other barbiturates bind with much less selectivity to these two classes of sites.

The in vivo mechanism of action of etomidate has been firmly linked to the GABA$\alpha$R. Heterologously expressed GABA$\alpha$Rs that have an N256M mutation on the M2 helix of the $\beta$3 subunit ($\beta^{+}$ surface of the interface) are relatively insensitive to etomidate (12), and sleep times in knock-in mice bearing the same mutation are much shorter than in wild-type mice (13).
Aziotomidate causes normal anesthesia in wild-type mice with the same potency as etomidate, and its action is similarly attenuated in the knock-in mouse (14). R-mTFD-MPAB also causes general anesthesia in mice and is equally potent in wild-type and N256M knock-in mice (15), consistent with the location of its binding sites at the $\beta^-$ subunit interfaces.

The contrasting subunit-selective actions of these two agents raise questions about the mechanism of general anesthesia itself, because there are 19 known GABA subunits, and which of the possible combinations occur in $\textit{vivo}$ is not yet fully defined. The state of anesthesia involves many behavioral components (16), so subunit-selective general anesthetics might be associated with specific subsets of the behavioral impairments experienced during anesthesia (17). Of particular interest are the relative contributions of phasic (synaptic) and tonic (extrasynaptic) inhibition actions (18, 19). The focus of this study is on the possible combinations occur in $\textit{vivo}$ is not yet fully defined. The state of anesthesia involves many behavioral components (16), so subunit-selective general anesthetics might be associated with specific subsets of the behavioral impairments experienced during anesthesia (17). Of particular interest are the relative contributions of phasic (synaptic) and tonic (extrasynaptic) inhibition actions (18, 19). The focus of this study is on the extrasynaptic $\alpha\beta\delta$ GABA$_A$Rs that are sensitive to endogenous neurosteroids and general anesthetics at concentrations lower than necessary to potentiate inhibitory postsynaptic currents (20–24). Expression studies in fibroblasts and oocytes establish that multiple combinations of $\alpha$, $\beta$, and $\delta$ subunits can combine to form functional receptors, which results in alternative subunit interfaces (25–31).

In this work, we photolabeled detergent-solubilized, purified heterologous $\alpha\beta\delta$ GABA$_A$Rs with $[^3\text{H}]$azietomidate and $[^3\text{H}]R$-mTFD-MPAB. Two distinct high affinity anesthetic sites were identified: 1) $[^3\text{H}]$azietomidate photolabeling established that azietomidate and etomidate bind to a $\beta^+\cdot \alpha 4^{-}$ interface site that does not bind $R$-mTFD-MPAB with high affinity; and 2) $[^3\text{H}]$azietomidate and $[^3\text{H}]R$-mTFD-MPAB share a common binding site with etomidate at a $\beta^-$ subunit interface. DS2, a positive allosteric modulator selective for GABA$_A$Rs containing a $\delta$ subunit (32), did not bind to these sites.

### Results

**Biochemical Characterization of the $\alpha\beta\delta$ GABA$_A$R**—Comparison of $[^3\text{H}]$muscimol binding to $\alpha\beta\delta$ GABA$_A$R in membranes and after purification in asolectin/CHAPS established that positive allosteric modulation was retained by etomidate and by DS2, a positive allosteric modulator selective for GABA$_A$Rs containing the $\delta$ subunit (32) (Table 1). In contrast to $\alpha1\beta3\gamma2$ GABA$_A$Rs, which bound $[^3\text{H}]$muscimol with similar affinity in membrane-bound ($K_{\text{eq}} = 50$ nM) and purified ($K_{\text{eq}} = 80$ nM) states (33), $[^3\text{H}]$muscimol bound to $\alpha\beta\delta$ GABA$_A$Rs in membranes ($K_{\text{eq}} = 13$ nM) with higher affinity than after purification in CHAPS/asolectin ($K_{\text{eq}} = 90$ nM). After purification, etomidate (10 $\mu$M) and DS2 (30 $\mu$M) increased the specific binding of 2 nM $[^3\text{H}]$muscimol by ~30%.

When samples of purified human $\alpha\beta\delta$ GABA$_A$R were fractionated by SDS-PAGE and visualized by Coomassie Blue stain, bands were readily visualized at 78 and 58 kDa, along with...
fainter bands at 72, 62, and 54 kDa (Fig. 2, lane 1). When extracted materials from in-gel tryptic digests of these bands were characterized by LC/MS/MS (Table 2), fragments of the GABA<sub>R</sub> γ4 subunit were most enriched in the 72 kDa band, consistent with the expected mobility of the mature subunit (58 kDa + 3 N-linked glycosylations). Fragments from the β3 subunit were concentrated in the 62 and 58 kDa bands, as found for β3 subunit from expressed α1β3γ2 GABA<sub>R</sub>γ4Rs (11). Fragments from the δ subunit were broadly distributed in the 62, 58, and 54 kDa bands, with α4 subunit fragments also recovered from the 54 kDa band. However, in contrast to the recovery of α4 subunit fragments from the 72 kDa gel band, for the 54 kDa band, no fragments were recovered from the α4 cytoplasmic domain beginning about 30 amino acids after the end of the M3 helix (data not shown). This result suggests that the 54 kDa band contains an N-terminal fragment of the α4 subunit containing the M1–M3 helices that was probably produced by proteolytic cleavage during receptor purification. The major component in the 78 kDa band was identified as the chaperone heat shock 70-kDa protein 1A (HSP70-1).

When material eluted from the 72 kDa band was characterized by Edman degradation, the primary sequence identified (XXLNXPGQNQXXXXXXX . . . ) matched a region near the predicted N-terminus of the human α4 GABA<sub>R</sub> subunit (VCNESPGQNQKEEKL . . . ). Multiple amino acids were detected at similar levels at each cycle of Edman degradation of the 62- and 58-kDa samples, which precluded de novo identification of the subunits present. Sequence analysis of material from the 54 kDa band identified a primary sequence (XNDLXXXKYCD . . . ) matching the N terminus region of the β-subunit of the GABA<sub>R</sub> δ GABA<sub>R</sub> subunit sequence (MNIDIGDYK3DDK . . . , with the underline denoting the FLAG peptide sequence). The N termini of the α4 and δ subunits identified by Edman degradation are those predicted to be the N termini of the mature subunits by the signal sequence cleavage site prediction program P-signal (34). No N-terminal sequence was detected from the 78-kDa material, consistent with the fact that the N-terminal alanine of 70-kDa heat shock protein is acetylated (35), preventing Edman degradation.

Table 2: LC/MS/MS identification of major peptides in αβ3δ GABA<sub>R</sub> SDS-polyacrylamide gel bands

| Gel band | Protein (gene name) | Peptides detected | MS/MS scans | Average intensity | Coverage |
|----------|---------------------|-------------------|-------------|------------------|----------|
| 78 kDa   | HSPA1A              | 68                | 911         | 1,450,000        | 70       |
|          | PRMT5               | 40                | 166         | 597,000          | 48       |
|          | GABRA4              | 27                | 127         | 393,000          | 46       |
| 72 kDa   | GABRB3              | 57                | 1,005       | 756,000          | 55       |
|          | LMNB1               | 29                | 65          | 253,000          | 48       |
| 62 kDa   | GABRB3              | 33                | 282         | 723,000          | 41       |
|          | LMNB1               | 32                | 127         | 500,000          | 45       |
| 58 kDa   | GABRD               | 29                | 487         | 924,000          | 38       |
|          | GABRB3              | 33                | 282         | 723,000          | 41       |
|          | LMNB1               | 32                | 127         | 500,000          | 45       |
| 54 kDa   | GABRD               | 35                | 304         | 1,970,000        | 40       |
|          | GABRB3              | 27                | 421         | 1,400,000        | 39       |
|          | GABRA4              | 28                | 115         | 712,000          | 42       |
|          | TUBB2A              | 47                | 476         | 655,000          | 59       |
|          | GABRA4              | 29                | 393         | 892,000          | 40       |
|          | GABRB3              | 15                | 52          | 639,000          | 30       |
|          | TUBA1A              | 13                | 31          | 176,000          | 35       |

α4β3δ GABA<sub>R</sub> General Anesthetic Binding Sites

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Labeling Human αβ3δ GABA<sub>R</sub> with Photoreactive Anesthetics—αβ3δ GABA<sub>R</sub> Rs were photolabeled at anesthetic concentrations with [3H]Methylazolidate or [3H]R-mTFD-MPAB in the presence or absence of etomidate at 1 mM or non-radioactive R-mTFD-MPAB at 60 μM, concentrations at which they each bind selectively to the β<sup>3</sup> or β<sup>1</sup> intersubunit sites in αβ3γ2 GABA<sub>R</sub>γ4Rs (11). When 3H incorporation was determined by fluorography after SDS-PAGE (Fig. 2, lanes 2–7), 3H incorporation was highest in the 58/62 kDa gel region for both photoreactive anesthetics. The 72 kDa band (α4) was labeled prominently only by [3H]Methylazolidate, and that photolabeling was inhibitable by etomidate but not by R-mTFD-MPAB. [3H]Methylazolidate photolabeling in the 58/58/62 kDa gel bands was inhibited to a greater extent by etomidate than by R-mTFD-MPAB, and, conversely, [3H]R-mTFD-MPAB photolabeling of 58/62 kDa gel bands was inhibited to a greater extent by R-mTFD-MPAB than by etomidate. These findings suggested that 1) there is an etomidate/methylazolidate binding site associated with the α4 subunit that does not bind R-mTFD-MPAB with high affinity; 2) methylazolidate, etomidate, and R-mTFD-MPAB share a common binding site associated with the 58/62 kDa gel band; and 3) there may be an R-mTFD-MPAB binding site associated with the 58/62 kDa gel band that does not bind etomidate.

To further characterize the pharmacological specificity of [3H]Methylazolidate and [3H]R-mTFD-MPAB incorporation at the subunit level, photolabelings were performed on an analytical scale in the presence of various concentrations of etomidate, R-mTFD-MPAB, the neuroactive steroid alaphaxalone, or DS2, with 3H incorporation into the gel bands quantified by liquid scintillation counting (Fig. 3). Etomidate inhibited [3H]Methylazolidate photoincorporation into the 72 kDa (α4) and 58/62 kDa bands with IC<sub>50</sub> values of ~15 μM, with high con-
centrations producing maximal inhibition of subunit photolabeling by 80 and 60%, respectively. Etomidate also inhibited \(^{3}\text{H}\)R-m-tfFd-MPAB photolabeling in the 58/62 kDa bands with similar potency. R-m-tfFd-MPAB inhibited \(^{3}\text{H}\)R-m-tfFd-MPAB photolabeling in the 58/62 kDa bands with an IC\(_{50}\) of 2 \(\mu\)M and a maximal inhibition of 75%. R-m-tfFd-MPAB also inhibited \(^{3}\text{H}\)azietomidate photolabeling in the 58/62 kDa bands with similar potency, but it inhibited \(^{3}\text{H}\)azietomidate labeling in the 72 kDa (\(\alpha 4\)) band only at the highest concentration tested (60 \(\mu\)M, \(<30\%\) inhibition).

As seen for \(^{3}\text{H}\)azietomidate and \(^{3}\text{H}\)R-m-tfFd-MPAB photolabeling of \(\alpha 1/\beta 3/\gamma 2\) GABA\(_R\) subunits (11), in the absence of GABA, the neuroactive steroid alfaxalone at concentrations up to 30 \(\mu\)M potentiated photoincorporation into the GABA\(_R\) subunit bands, maximally by \(<50\%\). This result establishes that alfaxalone does not bind to the sites in the purified \(\alpha 4/\beta 3/\delta\) GABA\(_R\) photolabeled by \(^{3}\text{H}\)azietomidate or \(^{3}\text{H}\)R-m-tfFd-MPAB but that there is positive allosteric linkage between alfaxalone and azietomidate/R-m-tfFd-MPAB binding. At concentrations up to 30 \(\mu\)M, DS2 had little or no effect on photolabeling by \(^{3}\text{H}\)azietomidate or \(^{3}\text{H}\)R-m-tfFd-MPAB in the presence of GABA.

Localization of \(\alpha 4/\beta 3/\delta\) GABA\(_R\) Residues Photolabeled by \(^{3}\text{H}\)Azietomidate—To provide an initial characterization of the locations of photolabeled amino acids, we fractionated by reversed phase HPLC (rpHPLC) EndoLys-C digests of subunit bands isolated from \(\alpha 4/\beta 3/\delta\) GABA\(_R\)s photolabeled with \(^{3}\text{H}\)azietomidate or \(^{3}\text{H}\)R-m-tfFd-MPAB (Fig. 4). For both photoreactive anesthetics, the digests of the 58/62 kDa band (\(\beta 3\) and \(\delta\) subunits) contained peaks of \(^{3}\text{H}\) in hydrophobic fractions (\(<55\%\) and 70% organic solvent) where fragments beginning at the N termini of the \(\beta 3\) and \(\beta 1\) helices are known to elute (36). For the 72 kDa band (\(\alpha 4\)) labeled by \(^{3}\text{H}\)azietomidate, the \(^{3}\text{H}\) eluted in two peaks, a broad hydrophobic peak (55% organic solvent), which for digests of \(\alpha 1\) subunits contains fragments beginning at the N termini of the M1 and M3 helices, and a peak at 40% organic solvent, where fragments from the \(\alpha 1\) subunit extracellular domain elute (36). For the 54 kDa band, for each drug, there were peaks of \(^{3}\text{H}\) at 40, 55, and 70% organic solvent, corresponding to the peaks seen in either of the higher molecular weight gel bands.

Etomidate Inhibits \(^{3}\text{H}\)Azietomidate Photolabeling of \(\alpha 4\)Met-269 (\(\alpha 4\)M1), \(\beta 3\)Met-227 (\(\beta 3\)M1), and \(\beta 3\)Met-286 (\(\beta 3\)M3)—Aliquots were sequenced of unfractionated EndoLys-C digests from the 72 and 58/62 kDa gel bands from GABA\(_R\)s photolabeled with \(^{3}\text{H}\)azietomidate in the absence and presence of non-radioactive etomidate (Fig. 5). For the 72 kDa band, there was a major peak of etomidate-inhibitable \(^{3}\text{H}\) release in cycle 14 (Fig. 5A). For the digest from the 58/62 kDa band (Fig. 5B), there were peaks of etomidate-inhibitable \(^{3}\text{H}\) release in cycles 7 and 12 (photoreologically specific photolabeling) and peaks of \(^{3}\text{H}\) release in cycles 3 and 19 that were not inhibited by etomidate (nonspecific labeling). The 72 kDa gel band digest will contain all possible \(\alpha 4\) subunit proteolytic fragments, including
These residues, receptors were photolabeled in the absence or the presence of 20 µM R-mTfD-MPAB, a concentration sufficient to occupy ~90% of its high affinity binding sites based upon the inhibition of photolabeling at the subunit level (Fig. 3). When fractions from the 58/62 kDa band were sequenced that contained the fragment beginning at β3Arg-216 at 15 pmol was present, along with fragments beginning at α4Val-313 (α4M3, ~2.6 pmol) and α4Ser-235 (~1 pmol). The peak of ″H release in cycle 14 was consistent with labeling of α4Met-269 (230 cpm/pmol) unaffected by R-mTfD-MPAB.

R-mTfD-MPAB Does Not Inhibit [3H]Aziotimidate Photolabeling of β3Met-286 — To determine whether [3H]aziotimidate was photolabeling amino acids in βM3 and/or δM3, we sequenced rpHPLC fractions from EndoLys-C digests of the 58/62 kDa gel band enriched in βM3 and from the 54 kDa gel band enriched in δM3. When fractions from the 58/62 kDa gel band were sequenced containing fragments beginning at β3Ala-280 (~6 pmol) and at δAla-309 (4 pmol) (Fig. 7A), there was a single peak of 3H release in cycle 14, consistent with photolabeling of α4Met-269 in α4M1 at 230 cpm/pmol in the absence or presence of R-mTfD-MPAB.

R-mTfD-MPAB Inhibits [3H]Aziotimidate Photolabeling of β3Met-227 but Not α4Met-269 — To confirm that [3H]aziotimidate photolabeled α4Met-269 and β3Met-227, samples were sequenced after rpHPLC fractionation of EndoLys-C digests of material from the 72, 58/62, and 54 kDa gel bands. To determine whether R-mTfD-MPAB also inhibited photolabeling of these residues, receptors were photolabeled in the absence or the presence of 20 µM R-mTfD-MPAB, a concentration sufficient to occupy ~90% of its high affinity binding sites based upon the inhibition of photolabeling at the subunit level (Fig. 3). When fractions from the 58/62 kDa band were sequenced that contained the fragment beginning at β3Arg-216 at ~15 pmol (Fig. 6A), the peak of 3H release in cycle 12 confirmed labeling of β3Met-227 (220 cpm/pmol), and R-mTfD-MPAB inhibited that labeling by ~80%. When fractions from the 72 kDa (α4) gel band were sequenced (Fig. 6B) that contained the fragment beginning at α4Met-256 (1 pmol), there was a single peak of 3H release in cycle 14, consistent with photolabeling of α4Met-269 in α4M1 at 230 cpm/pmol in the absence or presence of R-mTfD-MPAB.

R-mTfD-MPAB Does Not Inhibit [3H]Aziotimidate Photolabeling of β3Met-227 but Not α4Met-269 — To confirm that [3H]aziotimidate photolabeled α4Met-269 and β3Met-227, samples were sequenced after rpHPLC fractionation of EndoLys-C digests of material from the 72, 58/62, and 54 kDa gel bands. To determine whether R-mTfD-MPAB also inhibited photolabeling of these residues, receptors were photolabeled in the absence or the presence of 20 µM R-mTfD-MPAB, a concentration sufficient to occupy ~90% of its high affinity binding sites based upon the inhibition of photolabeling at the subunit level (Fig. 3). When fractions from the 58/62 kDa band were sequenced that contained the fragment beginning at β3Arg-216 at ~15 pmol (Fig. 6A), the peak of 3H release in cycle 12 confirmed labeling of β3Met-227 (220 cpm/pmol), and R-mTfD-MPAB inhibited that labeling by ~80%. When fractions from the 72 kDa (α4) gel band were sequenced (Fig. 6B) that contained the fragment beginning at α4Met-256 (1 pmol), there was a single peak of 3H release in cycle 14, consistent with photolabeling of α4Met-269 in α4M1 at 230 cpm/pmol in the absence or presence of R-mTfD-MPAB.

R-mTfD-MPAB Does Not Inhibit [3H]Aziotimidate Photolabeling of β3Met-286 — To determine whether [3H]aziotimidate was photolabeling amino acids in βM3 and/or δM3, we sequenced rpHPLC fractions from EndoLys-C digests of the 58/62 kDa gel band enriched in βM3 and from the 54 kDa gel band enriched in δM3. When fractions from the 58/62 kDa gel band were sequenced containing fragments beginning at β3Ala-280 (~6 pmol) and at δAla-309 (4 pmol) (Fig. 7A), there was a peak of 3H release in cycle 7 (280 cpm), which R-mTfD-MPAB inhibited by <15%. In contrast, when fractions from the 54 kDa gel band were sequenced containing fragments beginning at β3Ala-280 (1 pmol) and at δAla-309 (2 pmol), the peak of 3H release in cycle 7 was 25 cpm (Fig. 7B). Because the 3H release in cycle 7 was 25 cpm (Fig. 7B). Because the 3H release in cycle 7 was 25 cpm (Fig. 7B). Because the 3H release in cycle 7 was 25 cpm (Fig. 7B). Because the 3H release in cycle 7 was 25 cpm (Fig. 7B). Because the 3H release in cycle 7 was 25 cpm (Fig. 7B). Because the 3H release in cycle 7 was 25 cpm (Fig. 7B). Because the 3H release in cycle 7 was 25 cpm (Fig. 7B). Because the 3H release in cycle 7 was 25 cpm (Fig. 7B).
0.3H release in cycles 7 (50 cpm), 9 (90 cpm), and 10 (75 cpm) that contained the M3 fragments, there were prominent peaks in rpHPLC fractions from the 58/62 kDa digest were sequenced and labeled by [3H]azetomidate, the residues common to both β3M3 and δM3 (○) were not used for the repetitive yield fits. The peak of 3H release in cycle 7, if originating from β3M3, indicated photolabeling of δM3 (A, 300 cpm; B, 25 cpm) correlate well with the amounts of the β3Ala-280 fragment (A, 6 pmol; B, 1 pmol) but not with those of the δAla-309 fragment (A, 4 pmol; B, 2 pmol).

Releases in cycle 7 correlated well with the amount of the β3Ala-280 fragment but not with the amount of the δM3 fragment, the peak of 3H release in cycle 7 indicated photolabeling of β3Met-286 (130 cpm/pmol) rather than δTrp-315. That [3H]azetomidate labeling of β3Met-286 (βM3) and α4Met-269 (αM1) was inhibited by etomidate (Fig. 5), but not by R-mTFD-MPAB, indicates that etomidate and azetomidate bind to a site at the β3"-α4" interface that does not bind R-mTFD-MPAB with high affinity and is homologous to their binding site at the β3-α1 interface.

[3H]R-mTFD-MPAB Photolabels β3Met-227, β3Met-286, and β3Phe-289 in human α4β3δ GABA_{A}Rs (50 pmol of muscimol sites per condition) photolabeled with 0.5 μM [3H]R-mTFD-MPAB in the absence ( ), residues unique to δM3; l_0 = 8/4 pmol, without/with R-mTFD-MPAB) and/or presence (○) of 60 μM R-mTFD-MPAB, the detected pmol of residues common to β3M3 and δM3 (○) were not used for the repetitive yield fits. The 3H release in cycles 7 and 10, if originating from β3M3, indicated labeling of δAla-309 (δM3, residues unique to δM3; l_0 = 2/3 pmol, without/with R-mTFD-MPAB) with the secondary sequence beginning at δAla-309 ( ), residues unique to δM3; l_0 = 4 pmol, 380 fragment (B, 4 pmol; C, 2 pmol).

Fig. 8. [3H]R-mTFD-MPAB specifically photolabels β3Met-227, β3Met-286, and β3Phe-289 in human α4β3δ GABA_{A}Rs. A-C. H ( ), ○ and pmol of PTH-derivatives ( ), residues unique to δM3; l_0 = 4 pmol; residues unique to δM3. The detected pmol of residues common to both β3M3 and δM3 (○) were not used for the repetitive yield fits. The peak of 3H release in cycle 7, if originating from β3M3, indicated photolabeling of δM3 (A, 300 cpm; B, 25 cpm) correlate well with the amounts of the β3Ala-280 fragment (A, 6 pmol; B, 1 pmol) but not with those of the δAla-309 fragment (A, 4 pmol; B, 2 pmol). The peaks of 3H release in cycle 7, if originating from β3M3, indicated photolabeling of δM3 (A, 300 cpm; B, 25 cpm) correlate well with the amounts of the β3Ala-280 fragment (A, 6 pmol; B, 1 pmol) but not with those of the δAla-309 fragment (A, 4 pmol; B, 2 pmol).
Discussion

In this report, we provide a first characterization of the locations of anesthetic binding sites in a GABA<sub>R</sub> subtype expressed extrasynaptically in the CNS. We photolabeled purified human α4β3δ GABA<sub>R</sub>Rs with [3H]azietomidate and [3H]R-mTDF-MPAB, photoreactive anesthetics that have been used previously to identify two homologous but pharmacologically distinct classes of anesthetic binding sites in α1β3γ2 GABA<sub>R</sub>Rs (11). Based upon the identification of photolabeled amino acids and the results of competition photolabeling assays carried out at the level of intact subunits, we demonstrate that etomidate, but not R-mTDF-MPAB, binds with high affinity to a site at the β<sup>+</sup>-α<sup>-</sup> subunit interface in α4β3δ GABA<sub>R</sub>Rs that is equivalent to its binding site in α1β3γ2 GABA<sub>R</sub>Rs. In contrast to α1β3γ2 GABA<sub>R</sub>Rs, which bind R-mTDF-MPAB, but not etomidate, with high affinity to sites at the α<sup>+</sup> / γ<sup>-</sup> - β<sup>-</sup> interfaces in proximity to β3Met-227 in βM1, we find that etomidate as well as R-mTDF-MPAB bind with high affinity to a site in α4β3δ GABA<sub>R</sub>Rs containing β3Met-227. As discussed below, this site is most likely to be at a β<sup>-</sup> - β<sup>-</sup> subunit interface. The sites identified by photolabeling with [3H]azietomidate and [3H]R-mTDF-MPAB are distinct from the binding sites for alfaxalone, an anesthetic steroid, or DS-2, a δ subunit-selective positive allosteric modulator (32), because neither drug inhibited photolabeling.

α4β3δ GABA<sub>R</sub> Composition—Based upon mass spectrometry and Edman degradation, the affinity-purified α4β3δ GABA<sub>R</sub>Rs used in this work contain α4 and β3 subunits as well as the δ subunit, whose presence is assured because the FLAG epitope used for purification is attached near the δ subunit N terminus. However, we do not know whether the preparation is characterized by a single dominant subunit composition. Whereas receptors having a β3α4β3αδ subunit arrangement (counterclockwise when viewed from the extracellular side) with two β3<sup>+</sup>-αδ<sup>-</sup> interfaces containing the agonist sites and a δ subunit replacing the γ subunit have been reported to be strongly favored in transiently transfected HEK cells (27, 28, 37), other studies indicate that subunit stoichiometry can be variable and dependent upon the subunit cDNA transfection ratios (26). Also, studies using concatenated subunits provide evidence that the δ subunit can assume multiple positions in a receptor pentamer and can contribute to a β<sup>-</sup> - δ<sup>-</sup> agonist binding site (25, 27, 30).

In the absence of independent definition of the subunit composition and arrangement in our purified α4β3δ GABA<sub>R</sub>Rs, consideration of our photolabeling results suggests a β3α4β3δδβ3 or β3δβ3α4β3 organization for the stably transfected cell line used in our studies. We favor these stoichiometries because 1) they have a β3-β3 interface required for the shared azietomidate/etomidate/R-mTDF-MPAB binding site, and 2) they have three β3 subunits to every one α4 subunit, consistent with the similar levels of [3H]azietomidate incorporation (cpm/pmol) at the amino acid level in the β3 and α4 subunits (Fig. 6) in the presence of a higher level of [3H]azietomidate photolocation in the β3 gel band than in the α4 band (Fig. 2). However, a β2α4δ4δα4β2 pentameric concatemer, containing a β-β interface, also forms a functional receptor (30).

α4β3δ GABA<sub>R</sub> General Anesthetic Binding Sites

An Etomidate Binding Site at the β3<sup>+</sup>-α4<sup>-</sup> Interface—[3H]Azietomidate photolabeled β3Met-286 in β3M3 (β3<sup>+</sup> side of an interface) and α4Met-269 in α4M1 (α4<sup>-</sup> side), with etomidate inhibiting labeling by >90% and R-mTDF-MPAB by <15%. Because [3H]azietomidate also photolabeled β3Met-286 in α1β3γ2 GABA<sub>R</sub>Rs and α4Met-269 is homologous to α1Met-236 that was also photolabeled (11), the simplest interpretation of these results is that there is an etomidate/azietomidate binding site at a β<sup>+</sup> - α<sup>-</sup> interface homologous to the etomidate site at the β3<sup>-</sup> - α<sup>-</sup> interfaces in α1β3γ2 GABA<sub>R</sub>Rs. This conservation of etomidate binding sites between α4β3δ and α1β3γ2 GABA<sub>R</sub>Rs is not unexpected, in view of the strong conservation of amino acids in the regions of the α2 and α1 subunit M1 and M2 helices that contribute to the α<sup>-</sup> surface of the etomidate binding sites (Fig. 9) and the fact that etomidate produces similar allosteric modulation in α1β3δ and α1β3γ2 GABA<sub>R</sub>Rs (38).

An Etomidate/R-mTDF-MPAB Binding Site at a β3<sup>-</sup> Interface—The most prominently labeled residue in the β3 subunit for both [3H]azietomidate and [3H]R-mTDF-MPAB was β3Met-227. Whereas in α1β3γ2 GABA<sub>R</sub>Rs, etomidate enhanced [3H]R-mTDF-MPAB photolabeling of this residue in α<sup>+</sup> - β<sup>-</sup> and / or γ<sup>-</sup> - β<sup>-</sup> intersubunit sites (11), etomidate inhibited this photolabeling by >90% in the α4β3δ GABA<sub>R</sub>, where β3Met-227 can potentially contribute to anesthetic binding sites at the α4<sup>-</sup> - β3<sup>+</sup>, δ<sup>-</sup> - β3<sup>-</sup>, or β3<sup>-</sup> - β3<sup>-</sup> subunit interfaces.

Several lines of evidence indicate that the β3<sup>-</sup> - β3<sup>-</sup> interface is the most likely interface for the site binding etomidate, azietomidate, and R-mTDF-MPAB with high affinity. 1) Photolabeling studies with expressed α1β3 GABA<sub>R</sub>Rs establish that etomidate, [3H]azietomidate, and [3H]R-mTDF-MPAB all bind with high affinity to the β3<sup>-</sup> - β3<sup>-</sup> interface pocket that is present in α1β3 but not in α1β3γ2 GABA<sub>R</sub>Rs (9, 39). 2) Examination of the amino acid residues that would contribute to the three alternative binding pockets (Fig. 9) identifies non-conservative substitutions contributing to the (+)-surface of the binding pocket that are expected to prevent the high affinity binding of etomidate in an α4<sup>-</sup> - β3<sup>-</sup> or δ<sup>-</sup> - β3<sup>-</sup> intersubunit pocket. In α1β2/3γ2 GABA<sub>R</sub>Rs, β2/3Asn-265 (βM2-15<sup>1</sup>) is known to be a major determinant of etomidate binding affinity, and in vitro and in vivo mutational analyses establish that replacement by Ser (α4M2-15<sup>1</sup>) or Met (δM2-15<sup>1</sup>) reduces etomidate potency by >10-fold (12, 13, 40 – 42). Similarly, substitution of β3Met-286 by Trp (the δ residue in the β3Met-286 position) also inhibits the effects of etomidate (40, 43). Therefore, it is unlikely that etomidate can bind with high affinity at either the α4<sup>-</sup> - β3<sup>-</sup> or δ<sup>-</sup> - β3<sup>-</sup> interface.

Contributions of δ Subunit Residues to Etomidate/Barbiturate Binding Sites—In our study, we did not identify any δ subunit amino acids photolabeled in an anesthetic-inhibitable manner by [3H]azietomidate or [3H]R-mTDF-MPAB. Based upon sequence analyses of samples containing variable amounts of βM3 and δM3, any pharmacologically specific photolabeling in δM3 is at <15% the level of βM3. It is possible that [3H]azietomidate does bind in a pocket containing δM3 residues that is homologous to the β<sup>-</sup> - α<sup>-</sup> site without photolabeling any residue in δM3, because the pocket would lack the methionine side chains favored by azietomidate’s photoreac-
sequence analysis. In fact, δM1 may contribute to a barbiturate binding site, because studies with receptors containing α1, β3, and chimeric γ/δ subunits indicated that pentobarbital sensitivity determinants were contained within a fragment containing the amino terminus and the first 3 amino acids of δM1 (44). Further studies will be necessary to clarify whether general anesthetics also bind with high affinity in the pocket at the β+δ site in the cell line used in this study or at the α+δ interface in δβαδ GABAαRs.

Functional Significance of the Identified Binding Sites—Photolabeling studies provided a detailed description of two classes of pharmacologically distinct binding sites for intravenous general anesthetics at subunit interfaces in the αβ3γ2 GABAαR transmembrane domain (8, 11, 36) that overlap with the binding sites for ivertemcin (45). Mutational analyses of the residues identified by photoaffinity labeling as well as neighboring residues in the shared subunit interface pockets have demonstrated their contributions to GABAαR gating and as determinants of anesthetic efficacy (12, 40, 43, 46, 47). In addition, the capacity of anesthetics to protect against modification of substituted cysteines has expanded the definition of residues contributing to anesthetic binding sites (48, 49). Mutational analyses will be necessary to determine, for example, whether the β′-α site, and β′-β sites identified by photoaffinity labeling are equally important for etomidate enhancement of GABA responses in an αβδδ GABAαR. However, in view of the difficulty of expressing αβ3δ GABAαR with defined subunit stoichiometry and subunit arrangement, these studies should be carried out using pentameric concatenated receptors.

Experimental Procedures

Materials—[3H]Muscimol (36 Ci/mmol) was from Perkin-Elmer Life Sciences. The detergents n-dodecyl β-D-maltopyranoside and CHAPS were from Anatrace-Affymetrix (anagrade quality). R-mTFD-MPAB and [3H]R-mTFD-MPAB (38 Ci/mmol) were prepared previously (50), as was [3H]azietomidate (19.3 Ci/mmol) (39). Soy bean asolectin, R- etomidate, and GABA were from Sigma. DS2 and alphaxalone were from Tocris. EndoLys-C was from Roche Applied Sciences.

Purification of αβδδ GABAαR—A detailed description of the expression and affinity purification of αβδδ GABAαR will be presented elsewhere. As described previously for αβ3γ2 GABAαRs (33), a stably transfected, tetracycline-inducible HEK293-TetR cell line expressing human GABAαR subunits α4, β3 (splice variant 2), and δ containing a FLAG tag near its N terminus (between δGly-29 and δAsp-30) was induced and grown for 2–3 days, and then membranes were harvested, flash-frozen in liquid N2, and stored at −80 °C until use. GABAαRs were solubilized with 30 mm n-dodecyl β-D-maltopyranoside and affinity-purified as described (11), using a FLAG M2 antibody column. Columns were washed with buffer supplemented with 200 μM asolectin and 5 mM CHAPS and then eluted with 1.5 mM FLAG peptide in the wash buffer. Aliquots of the eluate fractions were assayed for [3H]muscimol binding, and eluate fractions were flash-frozen in liquid N2 and stored at −80 °C until use. Membranes harvested from 60 15-cm plates contained ~5–10 nmol of [3H]muscimol binding sites (15–20 pmol of sites/mg of membrane protein), and the
eluted with the purifications used for photolabeling contained 50–70 nM [3H]muscimol sites. Based upon [3H]muscimol binding, the receptor was purified at 10–25% yield from the starting membranes. Because the receptor was eluted in the presence of 1.5 mM FLAG peptide, it was not possible to estimate purity in terms of pmol of muscimol binding/mg of protein. Based upon analyses by SDS-PAGE and LC/MS/MS (see “Results”), GABAAR subunits were the dominant polypeptides in the preparation.

Radioiodinated Binding Assays—[3H]Muscimol binding to purified GABAAR was measured by filtration after precipitation with polyethylene glycol (8). The total concentration of sites in eluate fractions was determined at 250 nm [3H]muscimol with 1 µM GABA to determine nonspecific binding. Allosteric modulation of 2 nm [3H]muscimol binding was determined as described (9, 11).

Sequence Numbering—For α4, residue 1 is the predicted signal sequence Met; for β3, residue 1 is the predicted N terminus of the mature protein (splice variant 1, QSNVND ...), with β3Met-286 at the 15th position in the M2 helix (M2–15’); and for δ, the numbering begins with the signal sequence Met and excludes the inserted FLAG sequence (DYKDDDDK). The primary structure locations of transmembrane helices M1–M4 in the figures correspond to the extent of the individual α-helices in the β3 monomeric GABAAR crystal structure (Protein Data Bank code 4COF).

Analysis of the α4β3δ GABAAR Preparation by LC/MS and N-terminal Sequencing—Three aliquots (24 pmol of [3H]muscimol sites each) of α4β3δ GABAAR were separated by SDS-PAGE. Based upon Coomassie Blue staining, bands migrating at 78, 72, 62, 58, and 54 kDa were excised. The bands from one lane were submitted to the Harvard Medical School Taplin Mass Spectrometry Facility for reduction and alkylation, in-gel trypsin digestion, and peptide extraction for microcapillary LC/MS/MS analysis. The material from the equivalent gel bands from the other two lanes was eluted and subjected to N-terminal sequence analysis.

GABAAR Photolabeling—Aliquots of purified α4β3δ GABAAR in elution buffer were photolabeled at analytical or preparative scale (150–200 µl or 1–2 ml of GABAAR per condition, respectively) to characterize photoincorporation at the subunit level or to identify individual photolabeled amino acids by protein microsequencing. Aliquots of [3H]azetomidate or [3H]R-mTDF-MPAB were dried under a gentle argon stream and resuspended with GABAAR solutions for 30 min on ice with gentle vortexing. For preparative photolabeling, non-radioactive drugs were added directly to this resuspension, whereas for analytical photolabeling, drug aliquots were added by the use of a 1-µl syringe (Hamilton 86200) to 10 µl of purified GABAAR, which was then combined with 90–150 µl of GABAAR equilibrated with radioligand. With the exception of studies with alphaxalone, all photolabeling was carried out in the presence of 300 µM GABA. Samples were transferred to 96-well plastic plates or 3.5-cm diameter Petri dishes (Corning catalogue numbers 2797 and 3001) for analytical or preparative scale labeling and irradiated on ice with a 365-nm UV lamp (Spectroline EN-280L) for 30 min at a distance of <1 cm. Samples were then denatured by mixing 2 parts sample with 1 part SDS-PAGE sample buffer, incubated for ~30 min, and fractionated by modified Laemmli SDS-PAGE (11).

Stock solutions of non-radioactive R-mTDF-MPAB (60 mM), etomidate (60 mM), and alphaxalone (8 mM) were prepared in methanol. For these drugs, all samples during photolabeling contained methanol at a final concentration of 0.5% (v/v). DS2 was prepared at 6 mM in 90% methanol, 10% DMSO. For assays with DS2, samples during photolabeling contained methanol/DMSO at final concentrations of 0.45%/0.05% (v/v). To minimize losses of hydrophobic drugs due to adsorption on plastic surfaces, glass syringes, capillaries, pipettes, and vials were used for all material transfers up to the equilibration with the purified GABAAR in detergent/lipid.

After electrophoresis, gels were stained with Coomassie Brilliant Blue. In analytical scale experiments, [3H] incorporation into subunits was determined either by fluorography or by liquid scintillation counting of excised gel bands as described (11). In preparative scale experiments, material was eluted from the excised stained bands as described (11) and resuspended in gel digestion buffer (15 mM Tris, 0.5 mM EDTA, and 0.1% SDS, pH 8.4) for further analysis. Results from three preparative photoaffinity labelings of purified human α4β3δ GABAARs are presented in this work: 1) GABAAR (145 pmol of muscimol sites per condition) photolabeled with 3 µM [3H]azetomidate in the presence of 300 µM GABA with or without 1 mM etomidate; 2) GABAAR (110 pmol of muscimol sites per condition) photolabeled with 3.5 µM [3H]azetomidate in the presence of 300 µM GABA with or without 1 µM etomidate; and 3) GABAAR (50 pmol of muscimol sites per condition) photolabeled with 0.5 µM [3H]R-mTDF-MPAB in the presence of 300 µM GABA with or without 60 µM R-mTDF-MPAB.

To determine the relative binding affinity for anesthetics at the [3H]azetomidate or [3H]R-mTDF-MPAB binding sites, aliquots of α4β3δ GABAAR were photolabeled in the presence of various concentrations of a drug, and subunit gel slice counts from these aliquots were fit to the equation,

$$B(x) = \frac{B_0 - B_{ns}}{1 + \left(\frac{x}{IC_{50}}\right)} + B_{ns}$$

where $B(x)$ represents the gel slice [3H]cpm at total inhibitor concentration $x$, $B_0$ is the gel slice [3H]cpm in the absence of competitor, $B_{ns}$ is nonspecific [3H]cpm incorporation in the presence of maximal concentration of a competitor, and $IC_{50}$ is the total drug concentration producing 50% inhibition. Data were fit using Sigma Plot version 11.0 (Systat Software, Inc.) with $IC_{50}$ and $B_{ns}$ as adjustable parameters; $B_0$ was fixed at the experimentally observed value. Due to limited quantities of receptor, competition assays were done only once, and the S.E. values given are from the least-squares fits.

Proteolysis, Reversed-phase HPLC, and N-terminal Sequence Analysis—Aliquots of labeled subunits isolated from gel bands were digested (2 weeks, 20 °C, 0.3–1 units/sample) with EndoLys-C (Roche Applied Science). Digests were fractionated by RP-HPLC as described (51), except that the gradient began at 95% aqueous solvent (0.08% TFA) and 5% organic solvent (60% acetonitrile, 40% isopropyl alcohol, 0.05% TFA) and progressed
to 100% organic in 75 min by approximating (in 5-min intervals) the quadratic growth curve, \( f(x) = 5 + 0.017 \times x^2 \), where \( x \) is time in minutes and \( f(x) \) is percentage of organic solvent. The flow rate was 200 \( \mu \)l/min, and fractions were collected every 2.5 min, with 10% assayed for \(^3\text{H}\). Fractions of interest were pooled and drop-loaded onto glass fiber filters for N-terminal sequence analysis on an Applied Biosystems Procise 492 protein sequencer modified so that two-thirds of each cycle were injected for PTH-derivative detection and quantification, whereas one-third was collected for scintillation counting. Some samples were sequenced without rpHPLC separation by loading them onto Applied Biosystems ProSorb\(^{\text{TM}}\) PVDF filters by diluting the samples 10-fold into 0.1% TFA. The pmol of PTH-derivatives detected were calculated by using rpHPLC peak heights at 269 nm compared with a standard injection.

Photolabeling in \( \alpha \Delta \)M1 or \( \Delta \)M3 was determined by sequencing appropriate rpHPLC fractions from digests of the 72 kDa gel band. Labeling in \( \beta \Delta \)M1 and \( \beta \Delta \)M3 was identified by sequencing fractions from the 58/62 kDa gel bands. In preliminary studies, we established that that fragments containing \( \Delta \)M1 and \( \Delta \)M3 were present at the highest level in the fractions containing \( \beta \)M3 from the 58/62 kDa gel band, whereas one-third was collected for scintillation counting. Some of the paper with input from all authors. All authors approved the final version of the manuscript.

References

1. Sigel, E. (2005) The benzodiazepine recognition site on GABA\(_A\) receptors. Med. Chem. Rev. 2, 251–256
2. Miller, P. S., and Smart, T. G. (2010) Binding, activation and modulation of Cys-loop receptors. Trends Pharmacol. Sci. 31, 161–174
3. Sieghart, W. (2015) Allosteric modulation of GABA\(_A\) receptors via multiple drug-binding sites. Adv. Pharmacol. 72, 53–96
4. Hemmings, H. C., Jr., Akabas, M. H., Goldstein, P. A., Trudell, J. R., Orser, B. A., and Harrison, N. L. (2005) Emerging molecular mechanisms of general anesthetic action. Trends Pharmacol. Sci. 26, 503–510
5. Forman, S. A., Chiara, D. C., and Miller, K. W. (2015) Anesthetics target interfacial transmembrane sites in nicotinic acetylcholine receptors. Neurorapharmacology 96, 169–177
6. Sauguet, L., Shashsavar, A., and Delarue, M. (2015) Crystallographic studies of pharmacological sites in pentameric ligand-gated ion channels. Biochim. Biophys. Acta 1850, 511–523
7. Puthekamal, R., Hieckel, M., Simeone, X., Suwattanasophon, C., Fiedlbauer, R. V., Ecker, G. F., and Ernst, M. (2016) Structural studies of GABA-A receptor binding sites: which experimental structure tells us what? Front. Mol. Neurosci. 9, 44
8. Li, G.-D., Chiara, D. C., Sawyer, G. W., Husain, S. S., Olsen, R. W., and Cohen, J. B. (2006) Identification of a GABA\(_A\) receptor anesthetic binding site at subunit interfaces by photolabeling with an etomidate analog. J. Neurosci. 26, 11599–11605
9. Chiara, D. C., Dostalova, Z., Jayakar, S. S., Zhou, X., Miller, K. W., and Cohen, J. B. (2012) Mapping general anesthetic binding site(s) in human \( \alpha \)1B3 \( \gamma \)-aminobutyric acid type A receptors with \(^{[3\text{H}]\text{TDZBz}}\) etomidate, a photoreactive etomidate analog. Biochemistry 51, 836–847
10. Hibbs, R. E., and Gouaux, E. (2011) Principles of activation and permeation in an anion-selective Cys-loop receptor. Nature 474, 54–60
11. Chiara, D. C., Jayakar, S. S., Zhou, X., Zhang, X., Savechenkov, P. Y., Bruzik, K. S., Miller, K. W., and Cohen, J. B. (2013) Specificity of intersubunit general anesthetic-binding sites in the transmembrane domain of the human \( \alpha \)1B3\( \gamma \)-aminobutyric acid type A (GABA\(_A\)) receptor. J. Biol. Chem. 288, 19343–19357
12. Belelli, D., Lambert, J. J., Peters, J. A., Waford, K., and Whiting, P. J. (1997) The interaction of the general anesthetic etomidate with the \( \gamma \)-aminobutyric acid type A receptor is influenced by a single amino acid. Proc. Natl. Acad. Sci. U.S.A. 94, 11031–11036
13. Jurd, R., Arras, M., Lambert, S., Drexlar, B., Siegwart, R., Crestani, F., Zaugg, M., Vogt, K. E., Ledermann, B., Antkowiak, B., and Rudolph, U. (2003) General anesthetic actions in vivo strongly attenuated by a point mutation in the GABA\(_A\) receptor \( \beta \)3 subunit. FASEB J. 17, 250–252
14. Liao, M., Sonner, J. M., Husain, S. S., Miller, K. W., Jurd, R., Rudolph, U., and Eger, E. I. 2nd (2005) \( R \) (+) etomidate and the photoactivatable \( R \) (+) azetomidate have comparable anesthetic activity in wild-type mice and comparably decreased activity in mice with a N265M point mutation in the \( \gamma \)-aminobutyric acid receptor \( \beta \)3 subunit. Anesth. Analg. 101, 131–135, table of contents
15. Amlong, C. A., Perkins, M. G., Houle, T. T., Miller, K. W., and Pearce, R. A. (2016) Contrasting effects of the \( \gamma \)-aminobutyric acid type A receptor \( \beta \)3 subunit N265M mutation on loss of righting reflexes induced by etomidate and the novel anesthetic barbiturate \( R \)-mTFD-MPAB. Anesth. Analg. 123, 1241–1246
16. Patel, P. M., Patel, H. H., and Roth, D. (2011) General anesthetics and therapeutic gases. In Goodman and Gilman's The Pharmacological Basis of Experimental Therapeutics (Brunton, L., Chabner, B., and Knollman, B., eds) pp. 527–564, McGraw-Hill, New York
17. Drexlar, B., Antkowiak, B., Engin, E., and Rudolph, U. (2011) Identification and characterization of anesthetic targets by mouse molecular genetics approaches. Can. J. Anaesth. 58, 178–190
modulation of an α1β3γ2 γ-aminobutyric acid type A (GABA_A) receptor by binding to a site in the transmembrane domain at the γ-2/γ-1 interface. J. Biol. Chem. 290, 23432–23446
37. Herrera, N. P., Betta, J., You, H., Henderson, R. M., Martin, I. L., Dunn, S. M. J., and Edwarsson, J. M. (2008) Atomic force microscopy reveals the stoichiometry and subunit arrangement of the α4β3δ GABA_A receptor. Mol. Pharmacol. 73, 960–967
38. Feng, H. J., Jouaidi, Y., Haburcak, M., Yang, X., and Forman, S. A. (2014) Etomidate produces similar allosteric modulation in α1β3δ and α1β3δ2 GABA(A) receptors. Br. J. Pharmacol. 171, 789–798
39. Jayakar, S. S., Zhou, X., Chiara, D. C., Dostalova, Z., Savechenkov, P. Y., Buznik, K. S., Dailey, W. P., Miller, K. W., Eckenhoff, R. G., and Cohen, I. B. (2014) Multiple propofol-binding sites in a γ-aminobutyric acid type A receptor (GABA_A) identified using a photoactivatable propofol analog. J. Biol. Chem. 289, 27456–27468
40. Siegwart, R., Jurd, R., and Rudolph, U. (2002) Molecular determinants for the action of general anesthetics at recombinant αβ3γ2 γ-aminobutyric acid (A) receptors. J. Neurochem. 80, 140–148
41. Reynolds, D. S., Rosahl, T. W., Cirone, J., O’Meara, G. F., Haythornthwaite, A., Newman, R. J., Myers, J., Sur, C., Howell, O., Rutter, A. R., Atack, J., Macaulay, A. J., Hadingham, K. L., Hutson, P. H., Belelli, D., et al. (2003) Sedation and anesthesia mediated by distinct GABA-A receptor isoforms. J. Neurosci. 23, 8608–8617
42. Stewart, D. S., Pierce, D. W., Hotta, M., Stern, A. T., and Forman, S. A. (2014) Mutations at β2N265 in γ-aminobutyric acid type A receptors alter both binding affinity and efficacy of potent anesthetics. PLoS One 9, e111470
43. Stewart, D., Desai, R., Cheng, Q., Liu, A., and Forman, S. A. (2008) Tryptophan mutations at α2-ε1-motif coexpression sites on α1 or β2 subunits enhance GABA-A receptor gating and reduce etomidate modulation. Mol. Pharmacol. 74, 1687–1695
44. Feng, H. J., and Macdonald, R. L. (2010) Barbiturates require the N termi-nus and first transmembrane domain of the β subunit for enhancement of α1β3δ GABA-A receptor currents. J. Biol. Chem. 285, 23614–23621
45. Estrada-Mondragon, A., and Lynch, J. (2015) Functional characterization of ivermintin binding sites in α1β2γ12 GABA(A) receptors. Front. Mol. Neurosci. 8, 55
46. Krasowski, M. D., Nishikawa, K., Nikolaeva, N., Lin, A., and Harrison, N. L. (2001) Methionine 286 in transmembrane domain 3 of the GABA_A receptor β subunit controls a binding cavity for propofol and other alkylphenol general anesthetics. Neuropharmacology 41, 952–964
47. Maldfiass, M. C., Baur, R., and Sigel, E. (2016) Functional sites involved in modulation of the GABA_A receptor channel by the intravenous anesthetics propofol, etomidate and pentobarbital. Neuropharmacology 105, 207–214
48. Stewart, D. S., Hotta, M., Li, G. D., Desai, R., Chiara, D. C., Olsen, R. W., and Forman, S. A. (2013) Cysteine substitutions define etomidate binding and gating linkages in the α-M1 domain of γ-aminobutyric acid type A (GABA_A) receptors. J. Biol. Chem. 288, 30373–30386
49. Nourmahnad, A., Stern, A. T., Hotta, M., Stewart, D. S., Ziemba, A. M., Szabo, A., and Forman, S. A. (2016) Tryptophan and cysteine mutations in M1 helices of α1β3γ2 γ-aminobutyric acid type A receptors indicate distinct intersubunit sites for four intravenous anesthetics and one orphan site. Anesthesiology 125, 1144–1158
50. Savechenkov, P. Y., Zhang, X., Chiara, D. C., Stewart, D. S., Ge, R., Zhou, X., Raines, D. E., Cohen, J. B., Forman, S. A., Miller, K. W., and Buznik, K. S. (2012) Allyl m-trifluoromethylmethylazide mepobarbital: an unusually pot ent enantioselective and photoactive barbiturate general anesthetic. J. Med. Chem. 55, 6554–6565
51. Ziebell, M. R., Nirthanan, S., Husain, S. S., Miller, K. W., and Cohen, J. B. (2004) Identification of binding sites in the nicotinic acetylcholine receptor for [3H]azetidomide, a photoactivatable general anesthetic. J. Biol. Chem. 279, 17640–17649