Efficient Expression of Functional (α6β2)2β3 AChRs in Xenopus Oocytes from Free Subunits Using Slightly Modified α6 Subunits

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Abstract

Human (α6β2)(α4β2)β3 nicotinic acetylcholine receptors (AChRs) are essential for addiction to nicotine and a target for drug development for smoking cessation. Expressing this complex AChR is difficult, but has been achieved using subunit concatamers. In order to determine what limits expression of α6 AChRs and to efficiently express α6 AChRs using free subunits, we investigated expression of the simpler (α6β2)β3 AChR. The concatameric form of this AChR assembles well, but is transported to the cell surface inefficiently. Various chimeras of α6 with the closely related α3 subunit increased expression efficiency with free subunits and produced pharmacologically equivalent functional AChRs. A chimera in which the large cytoplasmic domain of α6 was replaced with that of α3 increased assembly with β2 subunits and transport of AChRs to the oocyte surface. Another chimera replacing the unique methionine 211 of α6 with leucine found at this position in transmembrane domain 1 of α3 and other α subunits increased assembly of mature subunits containing β3 subunits within oocytes. Combining both α3 sequences in an α6 chimera increased expression of functional (α6β2)β3 AChRs to 12-fold more than with concatamers. This is pragmatically useful, and provides insights on features of α6 subunit structure that limit its expression in transfected cells.

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Introduction

Human α6β2β3* nicotinic acetylcholine receptors (AChRs) on dopaminergic neurons are important targets for drugs to treat nicotine addiction and Parkinson’s disease [1–5]. These AChRs are critical brain subtypes in processes involving movement, memory, reward, and learning [6,7]. Self-administration of nicotine is inhibited by knockout of α6, β2, or α4 subunits, implying that (α6β2)(α4β2)β3 AChRs are necessary for presynaptically modulating dopamine release associated with reward and induction of nicotine dependence [8]. Antagonists of α6β2β3* AChRs inhibit nicotine self-administration by reducing nicotine-induced release of dopamine [2]. Smoking-relevant concentrations of nicotine sustain smoldering activation of dopaminergic neurons through (α6β2)(α4β2)β3 AChRs [5,9,10]. α6β2β3* AChRs are more abundant in dopaminergic neurons in primates than in rodents, thus probably even more important in humans than in rodents [11]. Therefore, it is critical to be able to express functional human α6β2β3* AChRs to develop better drugs.

However, efficiently expressing the complex (α6β2)(α4β2)β3 subtype or the simpler (α6β2)β3 subtype has been difficult [12–15]. Mixtures of free α6 and β2 subunits efficiently assemble (α6β2) binding sites, but not mature AChRs [12]. β3 subunits promote expression and nicotine-induced upregulation of human α6* AChRs expressed in transfected cell lines, but the amount of AChRs expressed is too small for detecting AChR function [13]. Use of subunit concatamers enabled expression in Xenopus oocytes of both human α6β2β3* AChR subtypes [16]. Linking subunits in pentameric concatamers results in efficient assembly of α6β2β3* AChRs [16]. A pentameric concatamer of (α6β2)(α4β2)β3 AChRs is efficiently transported to the surface [16]. However, transport of concatameric (α6β2)β3 AChRs to the surface membrane is inefficient. It is important to determine what limits expression of these subtypes and to express them efficiently.

Chimeras reveal α6 sequences that limit expression of transfected α6* AChRs. Chimeras with the extracellular domain of α6 and the remainder of α3 or β4 expressed in combination with β2 are efficiently transported to the surface [12]. The large cytoplasmic domain of some AChR subunits promotes assembly and transport of AChRs [17]. This suggests that the large cytoplasmic domain of α3 or β4 might provide efficient transport to the surface that α6 does not. The cytoplasmic domain of α3 is smaller than that of β4 and more closely resembles the sequence of α6.

Analysis of chimeras of α6 and α3 identified α3 sequences that permitted expression of α6* AChRs and sequences of α6 that inhibited expression of α3 AChRs [16]. A region of α6 in the first half of transmembrane domain 1 inhibited expression of α3 AChRs [16]. This region contains a unique methionine at position...
211 which is occupied by leucine in α3 and other α subunits. M211 is in a sequence that in α2 subunits governs stability, assembly, and transport to the cell surface [18]. This sequence is on the side of the α1 subunit that assembles with the accessory subunit (e.g. β1 in α1* AChRs or β3 in α6* AChRs) [19]. This suggests that the α6 sequence containing M211 might be important for associating with β3 accessory subunits. α6β2β3* AChRs are usually expressed in aminergic neurons at presynaptic locations [1,20,21]. These neurons may express a chaperone for Xenopus oocytes or human embryonic kidney (HEK) cell lines. These may express a chaperone for assembly of β3 subunits with α6 that is missing in other cell types such as Xenopus oocytes or human embryonic kidney (HEK) cell lines. These efficiently assemble β3 with other α subunits to form AChRs such as (α4β2)β3 [22].

Here we report efficient expression of (α6β2)β3 AChRs in Xenopus oocytes using free subunits with only small changes in α6 subunits, while not altering AChR pharmacology or channel structure. We explored the effects of incorporating M211L and α3 cytoplasmic domain alone, and together, into chimeras with α6. M211L increased assembly with β3, and α3 cytoplasmic domain increased assembly with β2 and transport to the surface. Together, these modifications synergistically permitted expression of high levels of (α6β2)β3 AChRs from free subunits. These AChRs exhibited the same pharmacological properties as concatameric AChRs, but were expressed on the oocyte surface in much greater amounts.

Materials and Methods

Ethics Statement

The anesthesia (1 g/1 L Tricane) used and treatment of Xenopus laevis frogs in our research experiments was performed in accordance with our approved IACUC protocol. The cRNA encoding desired subunits was synthesized from 1 µg of linearized cDNA templates in the pSP64 vectors using SP6 RNA polymerase from the mMessage mMachine kit (Ambion Inc, Austin, TX). Subunit cRNAs were mixed at a 1:1:1 ratio of α6:β2:β3 for constructs of free subunits. Dimeric concatamer was mixed with α6 chimera and β2 subunits at a 1:1:1 ratio. Xenopus laevis oocytes were injected with 100 ng of cRNA mix per oocyte, then incubated in 50% L-15 (Invitrogen, San Diego, CA), 10 mM HEPES, pH 7.5, 10 units/ml penicillin, 10 µg/ml streptomycin, and 50 µg/ml gentamycin [24]. This medium was refreshed daily, and replaced with gentamycin-free medium the day before recording.

Surface Expression of AChR

Surface expression was determined by binding of 125I mono-clonal antibody (mAb) 295 performed on the same day as electrophysiological recording of responses to 30 µM ACh. Groups of 8 oocytes were placed in an Eppendorf tube with 525 µL of L-15 media containing 10% horse serum and 5 nM 125I mAb 295 [25] at room temperature for 3 hours. Unbound 125I mAb was removed by 3 washes with 1 ml of L-15, then 125I bound to individual oocytes was determined in a γ-counter. Nonspecific binding was determined using non-injected oocytes.

3H Epibatidine Binding

Groups of 8 oocytes were homogenized in 1 ml of buffer A (50 mM NaCl, 50 mM sodium phosphate buffer, pH 7.3, 5 mM EDTA, 5 mM EGTA, 5 mM benazamide, 5 mM iodoacetamide, 2 mM phenylmethylsulfonyl fluoride). Then a crude membrane fraction was pelleted by centrifugation for 15 minutes at 13,400 rpm. Membrane proteins were resuspended by pipetting and solubilized in 150 µL of buffer A containing 2% Triton X-100 for 1 hour at room temperature. Debris was removed by centrifugation at 13,400 g for 15 min. Next, mAb 295 coated wells were loaded with aliquots of detergent extracts with 2 nM 3H epibatidine (PerkinElmer Life Sciences, Emeryville, CA) in a total volume of 100 µL in phosphate-buffered saline (PBS) buffer.
containing 0.5% Triton X-100 and 10 mM NaN₃ [12]. These plates were left overnight on a shaker at 4°C. Wells were then washed three times with 0.5% Triton X-100 in PBS. ³H epibatidine bound to AChRs bound to the wells through mAb 295 or mAb 210 was eluted with 30 μl of 0.1 M NaOH and quantitated by liquid scintillation counting [22]. Nonspecific binding was determined using non-injected oocytes.

Sucrose Gradient Sedimentation

Triton X-100-solubilized AChRs from oocytes were prepared as described above from groups of 30–50 oocytes [16]. Aliquots (150 μl) of the extracts, mixed with 2 μl of 1 μg/ml of Torpedo californica electric organ AChR, were loaded onto 11.3 ml sucrose gradients [linear 5–20% sucrose (w/v) in 10 mM sodium phosphate buffer, pH 7.5, that contained 100 mM NaCl, 1 mM NaN₃, 5 mM EDTA, 5 mM EGTA, and 0.5% Triton X-100] [16,26]. Gradients were centrifuged for 16 hours at 40,000 rpm in a SW-41 rotor (Beckman Coulter, Fullerton, CA) at 4°C. Fractions were collected at 15 drops per well from the bottom of the tubes. Fifty microliters of each fraction were transferred to mAb 295-coated wells to isolate β2-containing AChRs for measurement of ³H epibatidine binding, and 20 μl of each fraction were transferred to mAb 210-coated wells to isolate Torpedo californica AChRs used as internal molecular weight standards [26]. Fractions in mAb 295-coated wells were incubated with 2 nM ³H epibatidine at 4°C overnight. Fractions in mAb 210-coated wells were incubated with 1 nM ¹²⁵I α-bungarotoxin at 4°C overnight. Wells were then washed three times with PBS and 0.5% Triton X-100. Bound ³H epibatidine was determined by liquid scintillation counting.

Electrophysiology

Electrophysiological recording was performed with a twomicroelectrode voltage clamp amplifier (Oocyte Clamp OG-725; Warner Instruments, Hamden, CT) on Xenopus oocytes with a constant flow of ND-96 solution (96 mM NaCl, 1.8 mM CaCl₂, 1 mM μgCl₂, and 5 mM HEPES, pH 7.5) containing 0.5 mM atropine [16,27]. Whole-cell membrane currents were recorded in response to application of agonists 5–7 days after RNA injection at a clamp potential of −70 mV. Currents were measured in response to application of various concentrations of agonists for 4 seconds. Between agonist applications, the recording chamber was washed with ND-96 buffer containing 0.5 μM atropine for 3 minutes. Responses were normalized to the maximum response induced by acetylcholine (ACh)(30 μM). The mean value of at least five oocytes was used for graphing concentration/response curves. Values are expressed ± standard error.

Data were analyzed with KaleidaGraph version 4.1 software for common statistical determinants (Synergy Software, Reading, PA). EC₅₀, efficacy and Hill co-efficiency were obtained from the Hill equation as described [24].

Results

Design of (α6β[2]α3)β3 AChR Constructs

Subunit concatamers and chimeras of α6 and α3 subunits were incorporated into a series of constructs for expressing (α6β[2]α3)β3 AChRs as shown in Figure 1. Concatameric pentamer (construct 1) linked through (AGS)ₙ linkers (Figure 1B and 1D) enabled successful expression of functional (α6β[2]β3) AChRs in oocytes [16]. Free native α6, β2, and β3 subunits (construct 2) did not permit assembly of functional AChRs [12,16]. To investigate the effect of exchanging methionine 211 and/or cytoplasmic domain of α6 subunit to that of α3, three chimeras were made to build constructs 3, 4 and 5 (Figure 1C and 1D). Chimeras α6α3cyt-C; and construct 6 were made to investigate the effect of the short C-terminus following the last transmembrane domain of the α6 subunit. To increase β3 incorporation, a QAP linker was used to link β3 with α6. Combining β3-QAP-α6 concatamer with various chimeras shown in Figure 1C, constructs 7, 8 and 9 were made with one or two chimeric α6 subunits. Numbering and nomenclature of the constructs described above is used in the subsequent data figures.

Expression Efficiency in Oocytes

To evaluate efficiency of expression of the nine constructs in Figure 1, Xenopus oocytes were injected with mRNAs and six days later tested for total binding site assembly, surface protein level and responses to ACh. ACh binding sites were measured by binding of ³H epibatidine to immunosolated detergent solubilized components containing β2 subunits. Expression in the surface membrane was assayed by binding to oocytes of ¹²⁵I α-bungarotoxin to β2 subunits. Biophysical properties were determined by examining the currents induced by 30 μM ACh. Results are shown in Figure 2.

Various constructs produced very different amounts of ACh binding sites (Figure 2A). The amounts of AChRs on the oocyte surface were not proportionate to the total amount of ACh binding sites. For example, constructs 2, 4 and 5 all yielded more than 90 fmol of ³H epibatidine bound per oocyte but expressed very different amounts of AChRs on the surface, as low as 0.168±0.070 fmol for construct 2 or as high as 8.31±4.97 fmol for construct 5. This indicates that some constructs result in incompletely assembled AChRs or properly assembled AChRs that were not transported to the cell surface, as shown previously [12,16]. AChR function and AChR expression on the oocyte surface were closely correlated (Figure 2B), indicating that AChRs on the surface produced by most constructs had similar functional properties. The pentameric concatamer 1 (β3−α6−β2−α6−β2) exhibited the expected pharmacological properties and is considered the positive control [16]. Mature concatameric AChRs were assembled, but were inefficiently transported to the oocyte surface. Free native construct 2 (α6+β2+β3) subunits are the negative control. Free native subunits assembled large numbers of ACh binding sites. These α6β2α3 complexes assemble into amorphous aggregates, but virtually no mature AChRs [12]. Construct 3 (α6α3β2+β3) did not increase surface expression or function, and decreased assembly of ACh binding sites. Construct 4 (α6α3α3+β2+β3) greatly increased assembly with β2, surface expression, and function compared to construct 3. Construct 5 (α6α3α3α3+β2+β3) combines the α3 components of constructs 3 and 4 in one α6 chimera. Construct 5 increased expression on the surface and function 40–80 fold compared to free subunits and 10–13 fold compared to concatamer 1.

Other constructs did not approach the efficiency of expression obtained with construct 5 (α6α3α3+β2+β3). Construct 6 (α6α3α3+β2+β3) reduced assembly of ACh binding sites, surface expression, and function compared to construct 4 (α6α3α3+β2+β3). Thus, including α3 transmembrane domain 4 and C-terminal domain impaired assembly and transport. Subsequent experiments will show that pharmacological function was also altered. Construct 7 (β3−α6−α6+α3+β2) was intended to test whether the β3−α6 concatamer equaled or exceeded the effect of the α6α311 chimera in promoting assembly with β3 and whether a single α3 cytoplasmic domain was sufficient for enhanced assembly and transport. Construct 7 assembled fewer ACh binding sites than construct 4 with two α3 cytoplasmic domains, and was equally expressed on the surface. This is consistent with the ideas that assembly of β2 with α6 in the concatamers was reduced due
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A
Extracellular
Transmembrane
Cytoplasmic

C

α6
α3
α6_{211L}
α6/α3_{cyt}
α6_{211L}/α3_{cyt}
α6/α3_{cyt-C}

D

1. β3-α6-β2-α6-β2
2. α6 + β2 + β3
3. α6_{211L} + β2 + β3
4. α6_{α3 cyt} + β2 + β3
5. α6_{211L}, α3_{cyt} + β2 + β3
6. α6_{α3 cyt-C} + β2 + β3
7. β3-α6 + α6_{α3 cyt} + β2
8. β3-α6 + α6_{211L}, α3_{cyt} + β2
9. β3-α6_{211L}, α3_{cyt} + β2
to the absence of an α3 cytoplasmic domain in this α6 and that one α3 cytoplasmic domain per AChR is sufficient for transport to the surface. The β3–α6 concatamer provided no benefit. Construct 8 (β3–α6 + α6211L,α3cyt+β2) had no better surface expression or function than 7 and lower assembly of ACh binding sites. Thus, the β3–α6 concatamer with a single or double α6 chimera provided no benefit. Construct 9 (β3–α6211L,α3cyt+α6211L,α3cyt+β2) expressed functional AChRs better than 8. However, the β3–α6211L,α3cyt concatamer substantially impaired assembly relative to that achieved with the α6 double chimeras as free subunits. This is consistent with the idea that this concatamer did not promote assembly of β3 with α6 and prevented M211 from promoting this assembly which does occur with free subunits. The β3–α6 concatamer used a QAP linker rather than the (AGS)n linkers used in construct 1. The change in linker may account for the ineffectiveness of the β3–α6 concatamer, but this linker has proven effective in several other constructs (unpublished).

Most constructs exhibited a similar ratio of current per surface AChR in response to activation by 30 μM ACh (Figure 3). This suggests that they have similar channel opening and/or channel conductance. Construct 6 (α6,α3cyt+C+β2+β3) had larger current per surface AChR. This suggests that α3 transmembrane domain 4 and/or the extracellular C-terminal domain altered channel

Figure 1. Illustration of (α6β2)2β3 AChR constructs. A) Diagrammatic representation of an AChR subunit. B) Diagrammatic representation of two AChR subunits joined by a linker. Direction of the linker is indicated by arrows. C) Representation of α6 and α3 sequences used in the α6/α3 chimeras studied. D) Representation of (α6β2)2β3 AChRs assembled from the various constructs. Agonist binding sites are shown as solid triangles between two subunits. The number and nomenclature for each construct depicted here are used in the following data figures.

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Figure 2. Efficiency of expression of functional (α6β2)2β3 AChRs using various constructs in Xenopus oocytes. The mRNAs of the nine constructs were injected on the same day and the assays were performed 6 days later. Values are the average of results from at least 8 oocytes. A) Comparison of total ACh binding sites versus surface expression for each (α6β2)2β3 AChR construct. Total number of ACh binding sites in partially or completely assembled AChRs inside cells plus in the surface membrane was assayed by binding of 3H epibatidine. B) Comparison of function versus surface expression for each (α6β2)2β3 AChR construct. Electrophysiological function was assayed by the current evoked by 30 μM ACh voltage clamped oocytes. Mature AChRs on the oocyte surface were assayed by binding of 125I labeled mAb 295 to β2 subunits.

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Efficiency of Mature (α6β2)β3 AChR Assembly

Efficiency of assembly of mature AChR was analyzed by sedimentation velocity analysis on sucrose gradients (Figure 5). Confirming previous observations [16], the pentameric concatemer construct 1 exhibited a high proportion of mature AChRs of the expected size, intermediate between monomers and dimers of Torpedo AChR. Construct 2 (α6+β2+β3) resulted in high proportions of partially assembled AChRs and large aggregates, but virtually no mature AChRs. This confirms previous observations that free wild type subunits do not assemble into mature AChRs in Xenopus oocytes [12]. Construct 3 (α6β2+β2+β3) resulted in a substantial proportion of mature AChRs. Thus, replacing the methionine unique to α6 in transmembrane domain 1 with the leucine found in α3 and most other α subunits greatly promoted assembly of mature AChRs. Construct 4 (α6β2α3+β2+β3) resulted in mature AChRs, but also some partially assembled AChRs and a substantial proportion of large aggregates. Figure 2 showed that the absolute amounts of ACh binding sites assembled, AChRs transported to the surface, and functional AChRs were greater with the α6 chimera incorporating only the large cytoplasmic domain of α3 (construct 3) than with the chimera incorporating only leucine 211 of α3 (construct 3). Combining the two chimeras in an α6 subunit in construct 5 (α6β2α3+α3+β2+β3) gave a synergistic effect. The proportion of mature AChRs was large, and partially assembled AChRs and the largest aggregates were eliminated (Figure 5E). In addition, the absolute amount of AChRs assembled and transported to the surface membrane where their function was assayed increased 13–80 fold compared to when the chimeras were expressed individually (Figure 2).

Construct 6 (α6β2α3+β2+β3) produced a high proportion of mature AChRs (Figure 5F). Thus, the α3 transmembrane domain 4 and/or the α3 C-terminal extracellular domain contributed to assembly of an increased proportion of mature AChRs and a decreased proportion of large aggregates and assembly intermediates compared to the α3 cytoplasmic domain alone (Figure 5D). However, the absolute amount of functional AChRs on the cell surface was lower than with only the α3 cytoplasmic domain in construct 4 (Figure 2B). Constructs containing β3–α6 concatemers in constructs 7, 8, and 9 exhibited substantial proportions of mature AChR (Figure S1), although the absolute amounts of AChRs assembled or functional AChRs on the cell surface were much smaller than achieved with construct 5 containing the α6 double chimera, β2, and β3 as free subunits.

Pharmacology of (α6β2)β3 AChR Constructs

Although all of the (α6β2)β3 AChR constructs retain the α6 extracellular domain for agonist binding (Figure 1), altering transmembrane and cytoplasmic domains may change the pharmacology of (α6β2)β3 AChRs. Therefore, acetylcholine and various full and partial agonists of AChRs were used to investigate activation and desensitization of (α6β2)β3 AChR constructs (Figure 6, 7). Most constructs exhibited similar high sensitivities to activation by agonists (Figure 6, Table 1). Constructs 2 and 3 showed very little activation by acetylcholine, with maximum responses less than 20 nA. Thus their pharmacology was not studied in detail. EC50 of acetylcholine for activating all constructs, except 9, is around 1 μM. Kinetics of activation by ACh are displayed in Figure 7 and Figure S2. Constructs exhibited similar kinetics of activation and desensitization by various concentrations of ACh. For example, construct 5 (α6β2α3+β2+β3) responded to 30 μM ACh with 10 fold greater amplitude than construct 1 (β3–α6–β2–α6–β2) but had similar response kinetics. There were noticeable concentration-dependent shifts in peak times in
constructs 5, 6 and 7 (Figure 7 and Figure S2). The response persists after the ACh application period due to design of our electrophysiology testing apparatus, permitting washout to be slower than wash in. This effect might also result from calcium-activated chloride channels that remain open longer than the α6 receptors. Theoretically this effect could be avoided by omitting Ca”” from outside solution or clamping at a lower voltage. However, Ca”” may potentiate activation of AChRs by agonists and α6 AChR evoked peak current is strongly influenced by extracellular Ca”” [28,29]. Considering the overall low functional responses of the nine constructs, experiments were all executed at a holding potential of -70 mV and in a buffer with normal concentration of Ca””. Finally, altered kinetics of agonist responses were only exhibited by constructs that were not especially effective, thus characterizing them in detail is tangential to our goal of effectively expressing α6β2β3 AChRs.

Partial agonists nicotine, cytisine, and varenicline activate AChRs similarly, except construct 6 (Figure 6 and Table 1). Construct 6 (α6α5β1Cβ2β3) is an outlier in the concentration/response curves for nicotine, cytisine, and varenicline (Figure 6) and EC₅₀ values for nicotine and varenicline (Table 1). This suggests that α3 transmembrane domain 4 and/or the C-terminal domain alter sensitivity to activation.

Besides difference in activation, construct 6 (α6α5β1Cβ2β3) also exhibits more rapid desensitization than the other constructs (Table 2 and Figure S3). The desensitization rate of construct 6 increased 48% compared to that of construct 1. Construct 5, which has both M211L and the α3 large cytoplasmic domain, desensitized at a rate close to construct 1 (Table 2). Thus, α3 transmembrane domain 4 and/or the α3 C-terminal extracellular domain increase the rate of desensitization.

Discussion

It is hard to study a single AChR subtype in vivo because various subtypes often co-express together in brain. Establishing in vitro model systems expressing α6-containing AChRs is challenging. By changing a single unique amino acid in transfembrane domain 1 of α6 and the cytoplasmic domain of α6 to that of α3, we succeeded in efficiently expressing Xenopus oocytes large amounts of human (α6β2)β3 AChRs. This will be very useful for characterizing functional properties of these AChRs and drugs directed at them.

These AChRs have the pharmacological properties of (α6β2)β3 AChRs formed from wild type subunits linked in a concatamer. As discussed in our previous study, concatameric (α6β2)β3 AChRs showed the pharmacology expected from measuring dopamine release in the brain tissue [16]. EC₅₀ values for nicotine obtained in vivo ranged from 0.1 μM to 1 μM for activation of α6* AChRs not containing α4 [5,21,30]. Such variation may be due to technical variations. The pentameric concatamer and AChRs expressed from the free subunits of our highest expressing free subunit construct 5 have very similar pharmacological properties, (e.g. EC₅₀ for nicotine = 0.239 or 0.203 μM) but most of the constructs exhibit similar properties.

Expression of these modified free subunits is much more efficient than expression from (α6β2)β3 AChRs formed from wild type subunits linked in a concatamer [16]. Concatamers will still be required to ensure efficient expression and subunit order of the more complex (α6β2)α4β2β3 AChR subtype [16]. Dopaminergic neurons which uniquely express α6β2β3* AChRs may have chaperones for efficiently assembling β3 accessory subunits with α6 and ensuring efficient assembly of (α6β2) ACh binding sites in AChRs with α4β2 ACh binding sites [12,13]. In cells expressing free α4, α6, and β2 subunits, α6 and β2 do not effectively assemble to form (α6β2)α4β2β3 AChRs, instead α4β2 AChRs predominate (Kuryatov unpublished).
**Figure 5. Assembly of (α6β2)3 AChRs constructs evaluated by sucrose sedimentation velocity gradient analysis.** After centrifugation, gradient fractions were immunoisolated on microwells coated with mAb 295 to β2 to isolate (α6β2)3 AChRs prior to labeling with 3H epibatidine. Properly assembled mature (α6β2)3 AChRs sediment between the two internal standards, 9S monomer and 13S dimer of Torpedo californica AChRs. Peaks on the left of dimers in the gradient indicate multimers or aggregates of AChRs, while peaks on the right of monomers represent partially assembled AChRs. A) Expression of pentameric concatamer construct 1 (β3α6β2α6β2β2) resulted in a high proportion of mature AChRs, as expected (16). B) Expression of construct 2 (α6α6β2β2β2) resulted in a high proportion of aggregates and partially assembled AChRs and a very low proportion of mature AChRs, as expected (12). C) Expression of construct 3 (α6α3Cy-tβ2β3) resulted in a large proportion of mature AChRs. D) Expression of construct 4 (α6α3Cy-tβ2β3) resulted in mature AChRs but also partially assembled AChRs and both large and very large aggregates. E) Expression of construct 5 (α6α3Cy-tβ2β3) showed mature AChRs and some aggregates. F) Expression of construct 6 (α6α3Cy-Ctβ2β3) showed a high proportion of mature AChRs and few aggregates.

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**Figure 6. Concentration/response curves for constructs that resulted in significant amounts of functional AChRs.** Full agonist (ACh) and partial agonists (cytisine, nicotine, and varenicline) were used on (α6β2)3 AChRs. Each point is the average response of at least 5 oocytes. Arrows indicate construct 5, that behaves like construct 1, and constructs 6 and 9 that are divergent.

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Chimeras of α6 subunits with α3 subunits have revealed sequences of α6 that inhibit expression of α3β2 AChRs and sequences of α3 that permit expression of α6β2 AChRs [12,16]. Studies reported here build on this foundation to achieve efficient expression of (α6β2)2β3 AChRs and further explain structural limits to α6*-AChR expression.

The unique methionine 211 in the first transmembrane domain of α6, when exchanged for the leucine present in most other α subunits, promotes efficient assembly with β3 and formation of a concatamer (α6β2)2β3 AChR, which responds more rapidly with greater currents and more extensive desensitization at higher ACh concentrations. B) Construct 5 (α6211L, α3cyt+β2+β3) response kinetics to ACh are similar, but amplitudes of responses are much larger.

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### Table 1. Pharmacological Properties of Various (α6β2)2β3 AChR Constructs.

| Construct | Agonist 1 456 7 8 9 | EC50 (μM) 1.21 | EC50 (μM) 0.47 | Efficacy (%) 17.0 | EC50 (μM) 1.0 | EC50 (μM) 0.152 | Efficacy (%) 14.9 | EC50 (μM) 1.0 | EC50 (μM) 0.152 | Efficacy (%) 14.9 |
|-----------|----------------------|----------------|----------------|-------------------|----------------|----------------|-------------------|----------------|----------------|-------------------|
| ACh       | 1                    | 1.21 ± 0.16    | 0.33 ± 0.25    | 34.5              | 0.219 ± 0.12    | 0.189 ± 0.05   | 17.0              | 0.219 ± 0.12    | 0.189 ± 0.05   | 17.0              |
| Nicotine  | 2                    | 0.740 ± 0.096  | 0.33 ± 0.25    | 34.5              | 0.219 ± 0.12    | 0.189 ± 0.05   | 17.0              | 0.219 ± 0.12    | 0.189 ± 0.05   | 17.0              |
| Cytisine  | 3                    | 0.300 ± 0.04   | 0.250 ± 0.03   | 20.0              | 0.219 ± 0.12    | 0.189 ± 0.05   | 17.0              | 0.219 ± 0.12    | 0.189 ± 0.05   | 17.0              |
| Varenicline | 4                | 0.200 ± 0.02   | 0.150 ± 0.10   | 12.0              | 0.219 ± 0.12    | 0.189 ± 0.05   | 17.0              | 0.219 ± 0.12    | 0.189 ± 0.05   | 17.0              |

Note the EC50 values for construct 6, which contains the α3 cytoplasmic domain, transmembrane domain 4, and the extracellular C-terminal domain, are significantly different from those of construct 1 for nicotine (p = 0.0087).

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Figure 7. Kinetics of responses to increasing concentrations of ACh by constructs 1 and 5. A) Concatamer 1 (β3−α6−β2−α6−β2) responds more rapidly with greater currents and more extensive desensitization at higher ACh concentrations. B) Construct 5 (α6211L, α3cyt+β2+β3) response kinetics to ACh are similar, but amplitudes of responses are much larger.
Table 2. T1/2 of Various (α6β2)2 AChR Constructs against 30μM ACh.

| AChR Construct | T1/2 (min) | EC50 (μM) |
|----------------|-----------|-----------|
| 1 β6* β2* β3* | 0.57 ± 0.061 | 0.35 ± 0.064 |
| 2 β6* β2* β3* | 0.59 ± 0.038 | 0.105 ± 0.038 |
| 3 β6* β2* β3* | 0.54 ± 0.038 | 0.061 ± 0.061 |
| 4 β6* β2* β3* | 0.54 ± 0.038 | 0.064 ± 0.064 |
| 5 β6* β2* β3* | 0.54 ± 0.038 | 0.061 ± 0.061 |
| 6 β6* β2* β3* | 0.54 ± 0.038 | 0.061 ± 0.061 |
| 7 β6* β2* β3* | 0.54 ± 0.038 | 0.061 ± 0.061 |
| 8 β6* β2* β3* | 0.54 ± 0.038 | 0.061 ± 0.061 |
| 9 β6* β2* β3* | 0.54 ± 0.038 | 0.061 ± 0.061 |

Figure S1 Sucrose sedimentation velocity gradient analysis of the size of 3H epibatidine binding components formed by various constructs. Monomers (9.5S) and high proportion of mature AChRs. This suggests that a unique chaperone in dopaminergic neurons might bind to this region of α6 to promote assembly with β3. No chaperone is needed to get efficient assembly of β3 with α4 subunits that have a leucine at this position [22]. Replacing leucine 211 of α3 with methionine inhibits assembly of α3β2 AChRs [16].

The concatameric construct 1 (β3−α6−β2−α6−β2) efficiently assembles into mature AChRs, but these are not efficiently transported to the cell surface, as are the concatamers β3−α6−β2−α4−β2 or β3−α4−β2−α6−β2 [16]. This suggests that the large cytoplasmic domain of α4 may contribute to transport to the cell surface. The cytoplasmic domain of α4 is uniquely large. Here we show the normal sized cytoplasmic domain of α3, a subunit closely related to α6 in sequence, when substituted for that of α6 can promote both transport to the cell surface and increase assembly with β2 subunits. It is known that large cytoplasmic domain chimeras can increase the assembly and transport of AChRs [17]. Incorporation of shorter sequences of α3 cytoplasmic domain will be required to determine the minimum sequences required to promote assembly with β2 or surface transport. Use of shorter α3 cytoplasmic domain sequences might avoid incorporation of the α3 amphipathic α helix which could alter cation selectivity [17,31]. Identification of these shorter sequences may further suggest their mechanisms of action in transport and assembly and help to explain their collaboration with the single amino acid change M211L to promote expression of human (α6β2)3 AChRs from free subunits in Xenopus oocytes.

The synergism of these two chimeras appears to result from 211L promoting a high proportion (but low amount) of assembly of β3 combined with the cytoplasmic domain of α3 promoting extensive assembly with β2, and to a lesser extent β3, and facilitating transport of assembled AChRs to the oocyte surface. Linking β3 to α6 in a concatamer can force their assembly [16]. However, the dimeric concatamer used here did not exceed the effect of combining 211L and α3 cytoplasmic domain in generating functional (α6β2)3 AChRs.

The α6 chimeras with 211L and the α3 large cytoplasmic domain (construct 5) effectively expressed functional (α6β2)3 AChRs without altering their pharmacological properties. Transfection of HEK 293 cells with a 2:1:10 ratio of a chimera with the extracellular domain of α6 and the rest of α3, β2, and β3 with and a V9S gain of function mutation resulted in functional (α6β2)3 AChRs [32]. Despite these extensive modifications necessary in those studies to assemble β3 and get α6 to the surface of the cell line, some of the pharmacological properties of its AChRs were similar to constructs I and 5, while others were divergent. On the cell line, EC50 values for ACh (0.23 μM), nicotine (0.15 μM), cytisine (0.072 μM), and varenicline (0.022 μM) were lower than we observed by 1.4 to 5.9 fold, as might be expected as a result of their using a hyperactive mutant β3 subunit. Similarly, the efficacy for nicotine they observed was two fold greater than we observed for constructs I and 5.

In conclusion, we simplified and improved expression of (α6β2)3 AChRs in Xenopus oocytes while maintaining their pharmacological properties. The critical structural features that we identified as important for assembly and transport to the surface will be beneficial for understanding what happens in vivo and in developing α6* cell lines for drug screening.

Supporting Information
dimer (13S) of *Torpedo californica* AChR were sedimented as internal standards. A) Expression of construct 7 (β3–α6+2β2* cyt 3cyt) resulted in some very large aggregates, a substantial proportion of mature AChRs, and some partially assembled AChRs. B) Expression of construct 8 (β3–α6+2β2* cyt 3cyt 4* cyt) resulted in aggregate and substantial proportion of mature AChRs, and significant amounts of partially assembled AChRs. C) Expression of construct 9 (β3–α6+2β2* cyt 3cyt 5* cyt 6* cyt) resulted in a high proportion mature AChRs, and a substantial fraction of partially assembled AChRs.

**Figure S2** Kinetics of responses to increasing concentrations of ACh by constructs 4, 6, 7, and 9. A) Construct 4 (β3–α6+2β2* cyt 3cyt) was confirmed by western blot. B) Construct 6 (β3–α6+2β2* cyt 3cyt 4* cyt) was confirmed by western blot. C) Construct 7 (β3–α6+2β2* cyt 3cyt 4* cyt 5* cyt) was confirmed by western blot. D) Construct 8 (β3–α6+2β2* cyt 3cyt 4* cyt 5* cyt 6* cyt) was confirmed by western blot. E) Construct 9 (β3–α6+2β2* cyt 3cyt 4* cyt 5* cyt 6* cyt) was confirmed by western blot.

**Figure S3** Response kinetics to 30 μM ACh by constructs 1, 4–9.

**Figure S4** Stability of concatamer in construct 8 (β3–α6+2β2* cyt 3cyt 4* cyt 5* cyt 6* cyt) was confirmed by western blot. AChR solubilized from 100 oocytes solubilized by Triton X-100 was purified and concentrated by immunofluorescence chromatography using mAb 295 linked to resin before being resolved to subunits by SDS-polyacrylamide gel electrophoresis. After overnight incubation at 4°C, the column was spun at 5,000 rpm for 15 minutes to remove unbound material and washed with PBS, 0.5% Triton solution. 40 μl of LDS sample buffer (Invitrogen) was placed in the column and heated for 30 minutes at 37°C. The eluent was resolved by SDS-polyacrylamide gel electrophoresis and then transferred using a semidyry electroblotting method [16]. Blots were then quenched with 5% Carnation dried nonfat milk for 20 minutes in PBS, 0.5% Triton X-100, 10 mM NaNO3 for one hour. Blots were probed with rat antiserum to α6 (1:500) [13], then incubated with 2 nm 125I-labeled goat anti-rat IgG for 3 h at room temperature. After washing in 0.5% Triton with NaNO3, blots were visualized by autoradiography. Proteins of the sizes expected of QAP linker β3–26 (3.3×10^{-4} Da) and corresponding free subunit δβ211L cyt (5.5×10^{-4} Da) were obtained without signs of proteolytic degradation.

**Appendix**

**Table S1**

| Construct | Description |
|-----------|-------------|
| 1 | α6 nicotinic acetylcholine receptor |
| 2 | β3 nicotinic acetylcholine receptor |
| 3 | δβ211L cyt 4* cyt |
| 4 | δβ211L cyt 4* cyt 5* cyt |
| 5 | δβ211L cyt 4* cyt 5* cyt 6* cyt |

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**Author Contributions**

Conceived and designed the experiments: JML AK CKL. Performed the experiments: CKL AK. Analyzed the data: CKL AK. Contributed reagents/materials/analysis tools: AK CKL. Wrote the paper: the manuscript: JML CKL JW AK. Created the figures: CKL. Accepted funding for the research: JML.
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