Postsynaptic scaffolds for nicotinic receptors on neurons

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Complex postsynaptic scaffolds determine the structure and signaling capabilities of glutamatergic synapses. Recent studies indicate that some of the same scaffold components contribute to the formation and function of nicotinic synapses on neurons. PDZ-containing proteins comprising the PSD-95 family co-localize with nicotinic acetylcholine receptors (nAChRs) and mediate downstream signaling in the neurons. The PDZ-proteins also promote functional nicotinic innervation of the neurons, as does the scaffold protein APC and transmembrane proteins such as neuroligin and the EphB2 receptor. In addition, specific chaperones have been shown to facilitate nAChR assembly and transport to the cell surface. This review summarizes recent results in these areas and raises questions for the future about the mechanism and synaptic role of nAChR trafficking.

Keywords: nicotinic receptor; postsynaptic; scaffold; synapse; PSD-95; trafficking; neuroligin

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Introduction

Nicotinic acetylcholine receptors (nAChRs) are widely distributed throughout the central nervous system, and participate in numerous higher order functions[1,2], neurological disorders[3,4] and, of course, addiction[5]. The receptors comprise a family of subtypes in vertebrates, all of which are cation-selective ligand-gated ion channels[6–8]. At least 12 neuronal nAChR genes have been identified (α2-10, β2-4) which can form hetero- and homopentameric nAChRs[7–10]. Activation of nAChRs is able to produce diverse effects because of receptor location and because of downstream signaling pathways engaged by the receptors in the postsynaptic cell. The signaling pathways often employ calcium because nAChR activation depolarizes the cell and can activate voltage-gated calcium channels[11]. In the case of homomeric α7-containing receptors (α7-nAChRs), significant calcium can enter directly through the receptor itself[12–14]. Increasing evidence indicates that nAChRs in general, and α7-nAChRs in particular, are concentrated both pre- and postsynaptically at a variety of glutamatergic and GABAergic synapses[15,16–18]. Critical determinants for nicotinic cholinergic transmission, therefore, are the mechanisms that target and anchor nAChRs at synaptic locations and couple the receptors to specific signal transduction machinery. Little is known about such mechanisms for neuronal nAChRs.

The best understood mechanisms determining receptor localization and function on neurons are those operative postsynaptically at glutamate spine synapses. In this case, a vast number of components have been identified, linked directly or indirectly to the postsynaptic AMPA and NMDA receptors responsible for excitatory neurotransmission (Figure 1). Central are the membrane associated guanylate kinases (MAGUKs) comprising the PSD-95 family, which contain PDZ domains that bind other proteins[19]. PSD-95 itself binds directly to the intracellular C-terminal of NMDA receptors, and together with other associated PSD-95 molecules, links numerous components in an elaborate postsynaptic scaffold. Included are AMPA receptors, bound via a TARP link, as well as components important for signal transduction such as calcium/calcmodulin-dependent protein kinase II (CaMKII). SAP102 and PSD-93 are related members of the PSD-95 family and perform similar functions at glutamate synapses depending on the developmental stage and location of the synapse[20–22]. The fourth member of the family, SAP97, plays a different role, facilitating AMPA receptor trafficking to the surface membrane, for example[23]. Trafficking of AMPA receptors to the surface is a fundamental feature of synaptic plasticity[24,25]. Trafficking within the surface membrane has also recently emerged as a critical determinant of synaptic responses[26–28]. Whether similar
mechanisms might control the fate and function nAChRs has only recently emerged as a possibility.

**Postsynaptic scaffolds at nicotinic synapses**

Because nAChRs do not have intracellular N- or C-terminals, they were thought not likely to interact with PSD-95 family members. Surprisingly, the receptors do participate in PDZ-scaffolds in a variety of systems. It is not clear whether the interactions between nAChRs and PSD-95 family members are direct or indirect in those cases, but it is clear that the scaffold proteins are essential for mediating nAChR function.

Best characterized are the roles of PSD-95 family members in regulating nAChR function on autonomic neurons. PSD-93 co-localizes with nAChRs in mouse superior cervical ganglion neurons and submandibular ganglion neurons, and apparently tethers guanylate kinase-associated protein (GKAP) and Shank at the sites as it does at glutamate synapses. Moreover, immunoprecipitation of solubilized components shows that PSD-93 forms a complex with ganglionic nAChRs. Most importantly, denervation studies demonstrate that PSD-93 promotes synaptic stability; synaptic clusters of nAChRs disperse much more rapidly in mice lacking PSD-93.

All four PSD-95 family members are expressed by chick

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*Figure 1.* Model postsynaptic scaffold at a glutamate spine synapse. The postsynaptic density is comprised of membrane receptors and ion channels, scaffold and adaptor proteins, signaling proteins, cell-adhesion molecules and components of the cytoskeleton. Glutamate receptors, such as NMDARs (N-methyl-D-aspartate receptors) and AMPARs (α-amino-3-hydroxy-5-methyl-4-isoxazole propionic acid receptors), are located in the postsynaptic membrane, with the NMDARs at the center of the synapse and the AMPARs more peripheral. The PDZ-domain-containing scaffold proteins PSD95 (also known as DLG4) and the Src-homology domain 3 (SH3) and multiple ankyrin repeat domains (Shank) family form a two-layer protein network below the postsynaptic membrane, which is bridged by guanylate kinase-associated protein (GKAP). PSD95 forms membrane-perpendicular and roughly equally spaced filamentous structures, with its amino terminus attached to the membrane. Other signaling molecules occupy the spaces in the PSD95–GKAP–Shank protein web. Shank-family scaffolds are further linked to actin filaments. The domains of PSD95 and Shank (PDZ, SH3, guanylate kinase (GK), sterile-alpha motif (SAM) and ankyrin repeats (ANK) (see key)) are shown; other proteins are represented by simple shapes and are labeled. The presynaptic and postsynaptic membranes are connected by cell-adhesion molecules. Reprinted by permission from Macmillan Publishers Ltd: Nat Rev Neurosci 10(2), Feng W, Zhang M. Organization and dynamics of PDZ-domain-related supramodules in the postsynaptic density, 87-99, copyright 2009.
ciliary ganglion (CG) neurons\textsuperscript{[29]}. Three of them – PSD-93, PSD-95, and SAP102 – co-assemble with heteromeric nAChRs, as judged by immunoprecipitation of complexes solubilized from heterologous expression systems. PDZ-containing puncta co-distribute both with \(\alpha7\)-nAChRs and \(\alpha3\)-containing heteromeric receptors (\(\alpha3^*\)-nAChRs) on CG neurons. Dispersing the puncta disrupts nicotinic downstream signaling pathways in the neurons; this was done by transfecting the cells with a construct encoding a 9 amino acid peptide from cysteine-rich PDZ-binding protein (CRIP\(_T\)), that blocks PDZ-mediated protein-protein interactions. Receptor activation is no longer able to activate the transcription factor CREB and alter gene expression in the cells\textsuperscript{[29]}. A more surprising result is that CRIP\(_T\) dispersal of the PDZ-puncta also constrains functional innervation of the neurons. Neurons expressing CRIP\(_T\) received fewer spontaneous excitatory postsynaptic potentials (EPSCs) than controls, though the mean EPSC amplitude and nAChR immunostaining on the somata did not appear to be substantially reduced (Figure 2). The results suggested that the postsynaptic PDZ-scaffold anchors components that exert trans synaptic effects, perhaps aligning presynaptic release sites over postsynaptic nAChR clusters or boosting presynaptic release capabilities.

\begin{figure}
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\includegraphics[width=\textwidth]{figure2.png}
\caption{PDZ proteins codistribute with CG synaptic nAChRs; disrupting PDZ interactions in postsynaptic cells diminishes synaptic transmission. (A) Freshly dissociated E15 CG neurons were immunostained with a monoclonal antibody (mAb) that recognizes the PSD-95 family of PDZ proteins and costained with either goat polyclonal antisera for \(\alpha7\)-nAChRs (B) or with mAb 35 for \(\alpha3^*\)-nAChRs (E), and the paired images merged (C, F). Both \(\alpha7\)-nAChR and \(\alpha3^*\)-nAChR clusters colocalize with PDZ. Scale bar: 10 \(\mu\)m. (G) CG neurons in culture for 7 days develop \(\alpha3^*\)-nAChR clusters that colocalize with PDZ proteins. CG neurons were transfected on day 1 in culture with GFP-CRIP\(_T\), which disrupts all PSD-95 family PDZ interactions. Whole-cell patch-clamp recording revealed many spontaneous EPSCs in control cells transfected with GFP (upper trace) but only relatively few EPSCs in cells transfected with GFP-CRIP\(_T\) (bottom trace). Reprinted from Neuron, 38(5), Conroy WG, Liu Z, Nai Q, Coggan JS, Berg DK, PDZ-containing proteins provide a functional postsynaptic scaffold for nicotinic receptors in neurons, 759-71, 2003, with permission from Elsevier.}
\end{figure}
Postsynaptic PDZ-scaffolds at nicotinic synapses are not likely to be confined to autonomic neurons. In hippocampal neurons, α7-nAChRs appear to co-localize with PSD-95\textsuperscript{[30]}. What the PDZ-protein is doing there and how it might be associated with α7-nAChRs remain interesting questions. The same report showed that a Wnt-7a signaling pathway promoted accumulation of presynaptic α7-nAChRs co-localized with the scaffold protein adenosomatous polyposis coli (APC). Jacob and co-workers demonstrated that postsynaptically APC is localized with α3*-nAChRs on chick CG neurons, rather than with α7-nAChRs\textsuperscript{[32]}. PSD-93 forms part of the APC complex with α3*-nAChRs on CG neurons. In addition, the complex contains End binding protein 1 (EB1), macrophin, IQ motif containing GTPase activating protein 1 (IQGAP1), and 14-3-3 which together link α3*-nAChRs to the cytoskeleton and stabilize the postsynaptic complex\textsuperscript{[33]}.  

**Trans synaptic regulation**

The ability of CRIP to reduce nicotinic innervation of CG neurons suggested that the postsynaptic PDZ-scaffold organized components required for trans synaptic control of synapse formation. Neuroligin (NL) fulfills some of the criteria for such a component. CG neurons express several forms of NL and their binding partners α- and β-neurexins\textsuperscript{[34, 35]}. Overexpression of tagged NL demon-

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**Figure 3.** Required NL domains for induction of nicotinic synapses. (A) NL constructs were prepared for transfection of E8 CG neurons in culture. NL, rat NL-1 full length with an 8-amino acid FLAG epitope fused on the N-terminus; ExFmsCyNL, NL-1 construct in which the extracellular domain was replaced by the extracellular domain of Fms; ExFmsCyNL-pdz, ExFmsCyNL lacking the C-terminus PDZ-binding domain of NL-1; NL-pdz, NL-1 lacking the 5-amino acid C-terminus representing a PDZ-binding motif; NLc54 and NLc14, NL-1 truncated after the first 54 and 14 amino acids, respectively, of the cytoplasmic domain; ExNLgpi, extracellular AChE-like domain of NL-1 with a gpi linkage site at the C-terminus; GFPc54, GFP fused to the first 54 amino acids of the cytoplasmic domain of NL-1. Flag tag, 8 amino acids attached at the N-terminus; AChE homology domain, extracellular domain of NL-1 homologous to the equivalent region of AChE; TM, transmembrane domain; Cytoplasmic domain, cytoplasmic portion of NL-1; PDZ motif, PDZ-binding motif; GFP, green fluorescent protein sequence. (B) Changes in mEPSC frequency caused by individual NL-1 constructs. The frequency and amplitude of mEPSCs recorded (in TTX) from neurons 5−8 days after transfection with the indicated NL-1 constructs were expressed as a percent of the values obtained from control neurons in the same cultures and then averaged across experiments to obtain mean±SEM for the normalized values (n= # of neurons). CRIP refers to a GFP-CRIPT construct that disperses PDZ-scaffold proteins. Chick NL, chick full-length NL-1. All of the normalized values for frequency, except for ExNLgpi, were significantly different (P<0.05) from control (100%) where control represents cells transfected with GFP and untransfected cells; none of the normalized values for amplitude were significantly different from control (100%). The results indicate that extracellular and proximal cytoplasmic sequences of NL are necessary to enhance mEPSC frequency, while dominant-negative effects are observed for constructs having only the NL cytoplasmic sequence or the cytoplasmic sequence attached to an inappropriate extracellular sequence. PDZ interactions are not required for the effects. Reprinted from Dev Biol 307 (1), Conroy WG, Nai Q, Ross B, Naughton G, Berg DK, Postsynaptic neuroligin enhances presynaptic inputs at neuronal nicotinic synapses, 79−91, 2007, with permission from Elsevier.
strates that it co-localizes with nAChRs and can transcellularly induce accumulation of presynaptic components in adjacent neurites overlying the nAChR clusters[34]. Electrophysiological analysis of synaptic events indicates that NL increases the frequency of spontaneous miniature EPSCs (mEPSCs) recorded in the neurons without increasing mEPSC amplitude. This is most readily consistent with a presynaptic effect, though other possibilities remain. A dissection of the NL subdomains revealed that both the extracellular and intracellular domains were required for maximal mEPSC frequency, and further that the intracellular domain by itself functioned as a dominant negative (Figure 3). Unexpectedly, overexpression of NL boosted mEPSC frequency even if the construct lacked the PDZ-binding domain. The results suggested that high levels of NL can function as their own synapse-nucleating event and need not tether directly to a PDZ-scaffold.

Other transmembrane “synaptogenic” molecules are also expressed by CG neurons and can augment nicotinic innervation. These include the cell adhesion molecules L1 and SynCAM, both of which can act in CG neurons in culture to increase the number of synaptic contacts the cells receive[36]. This implies a trans synaptic effect. Electroporation studies confirmed that both endogenous NL and L1 act to provide this function in vivo. They are required for CG neurons to receive the expected number of presynaptic boutons overlying postsynaptic nAChR complexes (Figure 4). SynCAM, in contrast, is not critical for synapse formation in vivo but may nonetheless contribute to synaptic maturation[36].

Yet another transmembrane component interacting with nAChRs on CG neurons is the EphB2 receptor (EphB2R). Transsynaptic interactions between EphB2Rs on postsynaptic cells and the transmembrane protein ephrinB-1 on presynaptic neurons can influence the clustering and function of NMDA receptors[37–39]. On CG neurons, EphB2Rs co-localize with a7-nAChRs on somatic spines embedded within lipid rafts[40]. Activation of the EphB2Rs with an ephrinB-1 fragment had two effects: it physically constrained a7-nAChRs from dispersal following spine collapse or lipid raft dispersal, and it augmented nicotinic activation of the transcription factor CREB[40]. How it does this and what the physical basis is for EphB2R/a7-nAChR interactions remain open questions.

![Figure 4](https://i.imgur.com/3EJ5.png)

Figure 4. L1 RNAi show reduces innervation of a3*-nAChR clusters in ovo. (A) CG neurons transfected in culture with a construct encoding an RNAi against chicken L1 and RFP as a marker (red) showed lower levels of endogenous L1 as seen by immunostaining (green) than did nearby untransfected cells. This is most evident when viewing the images with the RFP fluorescence deleted to reveal the L1 stain (bottom). (B) An RNAi with a scrambled sequence (RNAi-control) had no effect on L1 levels. (C) Electroporation of CG neurons in ovo revealed significant reductions in SV2 levels abutting RNAi-L1 transfected cells compared to RNAi-control transfected cells, similar to the pattern seen with the LICyt-GFP construct. In contrast transfection with an RNAi construct that targets NL (RNAi-NL) had no affect on SV2 levels. (D) Quantifying the proportion of a3*-nAChR clusters with SV2 clusters apposed, to assay for potential synapses, revealed a similar pattern. Cells transfected with RNAi-L1, but not with RNAi-NL, had a significantly reduced fraction of a3*-nAChR clusters receiving SV2 puncta. *P<0.05 compared to RNAi-control by ANOVA with Bonferroni post-tests. Scale bars: 10 µm. Reprinted from Mol Cell Neurosci 39 (1), Triana-Baltzer GB, Liu Z, Gounko NV, Berg DK, Multiple cell adhesion molecules shaping a complex nicotinic synapse on neurons, 74-82, 2008, with permission from Elsevier.
Trafficking and chaperones

Current expectations about nAChR trafficking have been shaped in part by recent results showing that glutamate receptor trafficking both to and within the plasma membrane determines synaptic function and plasticity\(^\text{[41-43]}\). Early studies identified an "up-regulation" of functional nAChRs on the cell surface in response to chronic nicotine exposure\(^\text{[44-46]}\). It has now become clear that the up-regulation results from a variety of post-translational mechanisms including protein assembly and both trafficking and stabilization within the surface membrane\(^\text{[47-51]}\). Moreover, both the mechanism and the extent of up-regulation appear to be cell-type specific\(^\text{[52]}\).

Trafficking of nAChRs to the cell surface depends on chaperones. This may be most pronounced for \(\alpha_7\)-nAChRs which cannot be expressed by many cell types \(^\text{[53]}\). Ric-3 has been identified as a chaperone that helps assemble and traffic \(\alpha_7\)-nAChRs to the cell surface\(^\text{[54-56]}\). Yeast-2-hybrid analysis has identified other chaperones mediating assembly and transport of neuronal nAChRs. One is \(14-3-3\eta\) which interacts with \(\alpha_4\) nAChR subunits and increases the steady-state levels of \(\alpha_4\beta_2\)-nAChRs on the cell surface\(^\text{[57]}\). A second is VILIP-1 which also regulates \(\alpha_4\beta_2\)-nAChR surface expression\(^\text{[58]}\).

Receptor internalization is also likely to depend on specific scaffold components and contribute importantly to the regulation of nicotinic signaling. An interesting example is provided by the SNARE-dependent activity-induced internalization of \(\alpha_7\)-nAChRs; replacement from an internal pool is required to maintain downstream signaling\(^\text{[59]}\). Yet other scaffold proteins control \(\alpha_7\)-nAChR clustering as demonstrated by the report on PICK1\(^\text{[60]}\). The use of proteomics to identify proteins that interact specifically with individual nAChR subtypes will almost certainly divulge new and interesting players controlling nAChR trafficking and stabilization at synaptic sites\(^\text{[61]}\).

Rapid trafficking of nAChRs in the surface membrane is only beginning to be examined. Early studies on muscle nAChRs demonstrated that receptors can be mobile in the plasma membrane and traffic to sites of nerve-muscle contact\(^\text{[62-64]}\). Relatively rapid and reversible diffusion of the receptors can also occur in muscle membrane\(^\text{[65]}\). Recently it has been shown that nAChRs on autonomic neurons are capable of rapid lateral diffusion into and out of synapses depending on innervation and cellular conditions\(^\text{[66]}\). How this trafficking is regulated and what role it might play in nicotinic signaling in the CNS will be interesting issues to resolve.

Future directions

Although it is clear that the PSD-95 family of MAGUKs is critical for the maintenance of normal nicotinic synapse function, the roles of individual members have not yet been defined. Studies in glutamatergic systems have shown that PSD-95 family members exert both common and unique effects on synaptic function\(^\text{[42]}\). A similar level of complexity may exist at nicotinic synapses. Determining how and which PSD-95 family members specify nAChR expression, localization, and activity will be an important part of understanding the nature of nicotinic synapses.

The synaptic capabilities of nAChRs are likely to be determined in part by their precise spatio-temporal regulation on neurons. New technologies have recently been developed to characterize the dynamics of endogenous synaptic receptors with high temporal and spatial resolution. Prominent among these is single-particle-tracking studies using semiconductor quantum dots (QDs). Recently QD analysis has demonstrated that lateral diffusion of glutamate receptors is a critical mechanism shaping synaptic responses at glutamate synapses\(^\text{[26]}\). These new tools will be useful to resolve unanswered questions about the dynamics of nAChRs in central excitatory synapses. It will be exciting to apply these tools for the first time to the analysis of nAChRs on neurons.

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