

Agrin Binds to the Nerve–Muscle Basal Lamina via Laminin

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Abstract. Agrin is a heparan sulfate proteoglycan that is required for the formation and maintenance of neuromuscular junctions. During development, agrin is secreted from motor neurons to trigger the local aggregation of acetylcholine receptors (AChRs) and other proteins in the muscle fiber, which together compose the postsynaptic apparatus. After release from the motor neuron, agrin binds to the developing muscle basal lamina and remains associated with the synaptic portion throughout adulthood. We have recently shown that full-length chick agrin binds to a basement membrane-like preparation called Matrigel™. The first 130 amino acids from the NH₂ terminus are necessary for the binding, and they are the reason why, on cultured chick myotubes, AChR clusters induced by full-length

agrin are small. In the current report we show that an NH₂-terminal fragment of agrin containing these 130 amino acids is sufficient to bind to Matrigel™ and that the binding to this preparation is mediated by laminin-1. The fragment also binds to laminin-2 and -4, the predominant laminin isoforms of the muscle fiber basal lamina. On cultured myotubes, it colocalizes with laminin and is enriched in AChR aggregates. In addition, we show that the effect of full-length agrin on the size of AChR clusters is reversed in the presence of the NH₂-terminal agrin fragment. These data strongly suggest that binding of agrin to laminin provides the basis of its localization to synaptic basal lamina and other basement membranes.

EFFICIENT synaptic transmission requires a high local specialization of pre- and postsynaptic cells. At the neuromuscular junction (NMJ),¹ these specializations include aggregates of acetylcholine receptors (AChRs) and acetylcholinesterase (AChE) in the muscle cell membrane and accumulated vesicles containing the neurotransmitter acetylcholine in the nerve terminal (for review see Hall and Sanes, 1993). Differentiation of both pre- and postsynaptic cells is directed by molecules that are localized to the synaptic portion of the muscle cell basal lamina (Sanes et al., 1978; Burden et al., 1979). The capability of the synaptic basal lamina to control and maintain synaptic

differentiation makes it distinct from the extrasynaptic basal lamina.

Collagen type IV and laminin are the major components of cell basement membranes. Both molecules assemble from three separate chains. Individual chains are encoded by a family of homologous genes giving rise to collagen type IV and laminin isoforms that differ in their chain composition (Timpl, 1996). The collagen type IV molecules and the laminins are thought to form independent networks by self assembly that are linked by entactin/nidogen (Yurchenco and O'Rear, 1994). This scaffold builds the framework with which several other proteins like perlecan, fibulin-1, and fibulin-2 associate (for review see Timpl and Brown, 1996). While perlecan and nidogen are found both at synaptic and extrasynaptic sites of the muscle basal lamina, specific isoforms of collagen type IV and laminin are localized to the NMJ (Sanes, 1995). Other molecules that are concentrated at the NMJ include the neuregulins, AChE, and agrin, some of which have been shown to regulate different aspects of synapse formation (for review see Ruegg, 1996).

Among the best characterized molecules associated with synaptic basal lamina is agrin, a heparan sulfate proteoglycan (HSPG) with an apparent molecular mass on SDS-PAGE of 400–600 kD (Denzer et al., 1995; Tsen et al., 1995a). When added to cultured muscle cells, agrin induces the aggregation of AChRs and several other pro-

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1. *Abbreviations used in this paper:* AChE, acetylcholinesterase; AChR, acetylcholine receptor; ECM, extracellular matrix; HS-GAG, heparan sulfate glycosaminoglycan; HSPG, heparan sulfate proteoglycan; NMJ, neuromuscular junction; NtA, NH₂-terminal domain in agrin; VF, vitreous fluid.

teins that are also enriched at the NMJ *in vivo* (Nitkin et al., 1987). Several lines of evidence have led to the hypothesis that agrin is released from the nerve terminal of motor axons and causes the local accumulation of AChRs and other postsynaptic proteins in muscle fibers (McMahan, 1990). Consistent with this hypothesis, agrin-deficient mutant mice lack AChR clusters, and no functional NMJs are formed (Gautam et al., 1996). In chick agrin, three sites of alternative splicing have been described. One site is located near the NH₂ terminus (Denzer et al., 1995; Tsen et al., 1995b), and two sites, called A and B (y and z in rodents), are located in the COOH-terminal half of the protein. Splicing at site B strongly influences agrin's AChR-aggregating activity. Agrin isoforms with amino acid inserts at the B-site are highly active in inducing myotubes to aggregate AChRs, while agrin isoforms lacking inserts at this site are only marginally or not at all active (Ruegg et al., 1992; Ferns et al., 1993; Gesemann et al., 1995). Transcripts encoding the individual splice variants are differentially distributed. While AChR-aggregating isoforms are selectively expressed by neuronal cells, isoforms lacking inserts at site B are predominantly expressed by nonneuronal cells (for review see McMahan et al., 1992; Bowe and Fallon, 1995).

The local immobilization of motor neuron-derived agrin in the developing muscle basal lamina is thought to be important to maintain postsynaptic structures throughout adulthood (McMahan, 1990). This becomes evident as AChR-aggregating activity and agrin-like immunoreactivity remain localized to synaptic basal lamina for several weeks after degeneration of the cells at the NMJ (Burden et al., 1979; Reist et al., 1987). In addition to the NMJ, agrin-like immunoreactivity is also detected in basement membranes of other nonneuronal tissues, such as blood capillaries, kidney, and lung (Reist et al., 1987; Godfrey et al., 1988a; Magill-Solc and McMahan, 1988; Rupp et al., 1991). Hence, agrin must strongly bind to components of synaptic basal lamina and other basement membranes.

In a first attempt to characterize the binding of agrin to extracellular matrix (ECM), we have recently shown that full-length chick agrin binds selectively to Matrigel™ (Kleinman et al., 1982), a solubilized basement membrane extracted from the Engelbreth-Holm-Swarm mouse sarcoma. This binding required a 130–amino acid–long region at the NH₂-terminal end of agrin (Denzer et al., 1995). AChR clusters on cultured myotubes were also affected by this region as clusters induced by full-length agrin were more numerous but considerably smaller than those induced by any fragment of agrin without this NH₂-terminal region (Denzer et al., 1995). We now show that a fragment comprising this region of agrin is sufficient to bind to Matrigel™ and that the binding to this basement membrane preparation is mediated by laminin-1. The fragment also binds to laminin-2 and -4, the laminin isoforms expressed by muscle fibers, and it is concentrated in AChR clusters on cultured chick myotubes. Moreover, an excess of this fragment is sufficient to reverse the effect of full-length chick agrin on the size of AChR clusters. Searches on databases revealed that the NH₂-terminal region is highly conserved in mouse and human agrin. Based on these data, the 130 amino acids from the NH₂ terminus of agrin define a laminin-binding domain.

Materials and Methods

Cell Culture, Transfection, Protein Labeling, Immunoprecipitation, and Immunoblot

Culturing of primary chick myotubes and transfections of COS-7 (Gluzman, 1981) or HEK 293 cells (Graham et al., 1977) were carried out as described by Gesemann et al. (1995). Iodination of purified protein was performed as described (Gesemann et al., 1996). ³⁵S labeling, immunoprecipitation, and immunoblots were essentially done as described (Denzer et al., 1995). Recombinant cAgrin₇ and cΔN15Agrin (see Fig. 1 a) in the supernatant of transiently transfected COS cells were immunoprecipitated with the antiserum raised against cΔN15Agrin (Denzer et al., 1995), whereas cN25₇Fc and r21Fc (see Fig. 1 a) were directly bound to protein A-Sepharose (Pharmacia, LKB Biotechnology Inc., Piscataway, NJ). ³⁵S-labeled proteins were analyzed by SDS-PAGE on a 3–12% gradient gel followed by fluorography. For immunoblots, 3 μg of purified cAgrin₇ or agrin purified from vitreous fluid (VF agrin) and 5 μg of total chick heart laminins (Brandenberger and Chiquet, 1995) was separated by SDS-PAGE on a 3–12% gradient gel, transferred to nitrocellulose membrane, and analyzed as described in Denzer et al. (1995).

Expression Constructs

cDNA constructs pcAgrin and pcΔN15Agrin have been described previously (Denzer et al., 1995). The sequence encoding parts of the constant region of mouse immunoglobulin gamma heavy chain (Fc) were obtained by PCR using the primers sFc_XhoI_BamHI (GGCAGCTCGAGGATCCTCGTGCCCGAGGGATTGTGGTTG) and asFc_XbaI (GGCCTCTAGATCATTACCAGGAGAGTGGG). As template, the Fc part of mouse immunoglobulin (Bowen et al., 1996) was used. Amplification introduced an XhoI and a BamHI site at the 5' end and a XbaI site at the 3' end. The PCR product was digested with XhoI and XbaI and ligated into the expression vector pcDNAI (InVitrogen, San Diego, CA) to yield pFc. To generate pcN25₇Fc, a second PCR was performed using EcoRI_{s-289} (GCATAGAATTCGGCTGCGGGCGATGGG) and as469_BamHI (CACGAGGATCCCTCTGCACAGGGGTCCITG) as primers and pcAgrin_{7A4B8} (Denzer et al., 1995) as template. The product coded for the NH₂-terminal domain of chick agrin and contained an EcoRI site at the 5' end and a BamHI site at the 3' end. The BamHI site enabled the subsequent in-frame fusion of this PCR fragment to the Fc sequences by digestion with EcoRI and BamHI and ligation into pFc. pr21Fc was similarly constructed. The cDNA part encoding ray agrin was isolated by PCR using the primers EcoRI_{s3426} (AGCTTGAATTCAGCCAGTGGAAAGTGAATC) and as3967_BamHI (CACGAGGATCCCTTTCTTGGCTGGACAGTG), and r100_{A4B8} (Gesemann et al., 1995) as template. After digestion of the PCR product with EcoRI and BamHI, it was ligated into pFc. The sequence of pcN25₇Fc and r21Fc was verified by DNA sequencing.

Antiserum against β and γ Chains of Chick Laminins

Antiserum 648 against the β/γ chains of chick laminin-2 and -4 was generated as follows. 1 mg of chick heart laminin-2 and -4, purified as described (Brandenberger and Chiquet, 1995), was loaded on a preparative 3–15% gradient SDS-PAGE, run under reducing conditions, and blotted onto nitrocellulose (BA-85; Schleicher & Schuell Inc., Keene, NH). The protein band of 200 kD, containing the β and the γ chains, was cut out, suspended in 0.2 ml PBS, and sonicated on ice until homogenization. The sample was mixed with 0.2 ml Freund's complete adjuvant (GIBCO BRL, Gaithersburg, MD) and injected intracutaneously into one rabbit. The rabbit was boosted 1 mo later with the same preparation, suspended in incomplete Freund's adjuvant. Antiserum 648 was obtained 14 d later. In immunoblots, the antiserum recognized the 200-kD band of laminin isoforms from chick heart, chick gizzard, and mouse laminin-1 (Brandenberger and Chiquet, 1995; Ferris et al., 1996). As the only chain common to all these laminins is the γ1 chain, antiserum 648 must recognize at least this subunit. As the immunogen also contained the β chain, the antiserum is likely to recognize this subunit as well.

Protein Purification

Recombinant cAgrin₇ was obtained from stably transfected HEK 293 cells, whereas VF agrin was derived from eyes of day 14 chick embryos (Ruegg et al., 1989). 100 ml of serum-free conditioned medium or 50 ml of vitreous fluid was passed over a Mono Q-Sepharose (Pharmacia Diagnostics

AB, Uppsala, Sweden). The column was washed with 20 mM Tris-HCl, pH 7.2, 500 mM NaCl, and bound proteins were eluted with a linear gradient of NaCl from 500 mM to 2 M. Individual fractions with agrin immunoreactivity, as determined by an ELISA (Gesemann et al., 1995), were analyzed on SDS-PAGE (3–12% gradient) and visualized by silver staining (Morrissy, 1981). Agrin-containing fractions were pooled and dialyzed three times against PBS. Agrin concentration of such a preparation was determined by ELISA. cN25₇Fc and r21Fc from transiently transfected COS-7 cells were purified with protein A-Sepharose (Pharmacia) according to the manufacturer's advice. Purity of recovered protein was checked by SDS-PAGE and silver staining. The protein concentration was determined as described (Lowry et al., 1951), using the DC Protein assay kit (BioRad Labs, Hercules, CA) and BSA as a standard. Mouse laminin-1, purified from mouse Engelbreth-Holm-Swarm sarcoma as described (Timpl et al., 1979), was provided by Th. Schulthess (Biozentrum, University of Basel, Basel, Switzerland). Mouse nidogen, isolated from conditioned medium of stably transfected HEK 293 cells (Fox et al., 1991), and mouse perlecan (Timpl et al., 1979) were provided by Dr. R. Timpl (Max Planck Institute, Munich, Germany). Human collagen type IV (Weber et al., 1984; Ries et al., 1995) was obtained from Dr. K. Kühn (Max Planck Institute) and chick α -dystroglycan, isolated from skeletal muscle (Brancaccio et al., 1995), was a gift of Dr. A. Brancaccio (Biozentrum, University of Basel). Chick laminin isoforms (laminin-2 and -4) were purified from chick heart as described by Brandenberger and Chiquet (1995). Only laminin preparations functional in a neurite outgrowth assay (Brandenberger and Chiquet, 1995) were used for agrin-binding studies.

Solid-Phase Radioligand Binding Assay

Proteins were diluted to 10 μ g/ml with 50 mM sodium bicarbonate, pH 9.6 (coating buffer), and immobilized on 96-well plates (Becton-Dickinson, Bedford, MA) by incubation overnight at 4°C. Remaining binding sites were blocked for 1 h with TBS containing 3% BSA, 1.25 mM CaCl₂, and 1 mM MgCl₂ (blocking solution). ¹²⁵I-cAgrin₇ or ¹²⁵I-cN25₇Fc, diluted in blocking solution, was added and incubated for 3 h. After washing with TBS, 1.25 mM CaCl₂, 1 mM MgCl₂ four times, bound radioactivity in each well was counted with a gamma counter. Solubilization with SDS sample buffer and subsequent analysis on SDS-PAGE followed by silver staining confirmed that the proteins were indeed coated onto the plastic surface.

Alternatively, mAb 5B1 diluted to 10 μ g/ml in coating buffer was first immobilized by overnight incubation at 4°C. Remaining binding sites were saturated with blocking solution for 1 h. Then 3 μ g/ml agrin was added. After 1 h of incubation, the plates were washed three times with blocking solution and processed with ¹²⁵I-laminin-1 as described above.

To calculate half-maximal binding (EC₅₀) of cAgrin₇ to laminin-1, individual data points of the dose-response were fit by the following equation: $Y = (X/EC_{50}) / (1 + X/EC_{50}) \times PI$. This equation assumes a single class of equivalent and independent binding sites, where Y represents cpm, X represents the concentration of agrin, and PI represents cpm at saturation.

Immunocytochemistry

Double staining of agrin in COS-7 cells: COS cells, transiently transfected with cDNAs encoding cAgrin₇ and c Δ N15Agrin, were grown on Matrigel™ (Becton-Dickinson) and stained for agrin as described elsewhere (Denzer et al., 1995). Cells transfected with cDNAs encoding cN25₇Fc and r21Fc were also stained as described in Denzer et al. (1995), except that fluorescein-conjugated goat anti-mouse IgG (1:200; Cappel, Organon Teknika Corp., West Chester, PA) was used for the extracellular staining and that, before permeabilization and staining with rhodamine-conjugated goat anti-mouse IgG (1:200; Cappel), residual binding sites were blocked with unlabeled goat anti-mouse IgG (1:50; Cappel).

Primary chick myotubes were incubated with 200 nM c21_{B8} (Gesemann et al., 1995) for 16 h at 37°C. To localize cN25₇Fc, 20 nM of this fragment was included. Cultures were rinsed with culture medium, and to visualize individual components, they were stained with the following reagents: AChRs with 4×10^{-8} M rhodamine- α -bungarotoxin (Molecular Probes, Eugene, OR); cN25₇Fc with 5 μ g/ml biotinylated goat anti-mouse IgG (Molecular Probes) followed by 3 μ g/ml fluorescein-conjugated streptavidin (Jackson ImmunoResearch Laboratories, Inc., West Grove, PA); β/γ subunits of laminin with antiserum 648 (1:1,000) followed by Cy3™-conjugated goat anti-rabbit IgG (1:200; Jackson ImmunoResearch Laboratories, Inc.); and β 2 chain of laminin with mAb C4 (10 μ g/ml) followed by 5 μ g/ml biotinylated goat anti-mouse IgG and 3 μ g/ml fluorescein-conjugated streptavidin. The first incubation was done for 1 h at 37°C. Cultures were then washed with culture medium and fixed for 30 min at room tem-

perature with 4% paraformaldehyde, 11% sucrose in 0.1 M potassium phosphate, pH 7.2. After rinsing the cells with PBS and PBS + 20 mM glycine, the cultures were incubated for 1 h with the secondary reagents indicated before, diluted in PBS + 2% normal goat serum (PBSN). After rinsing the cells with PBS, cultures were either dehydrated with 95% ethanol at -20°C or incubated once more with the tertiary reagent diluted in PBSN. Coverslips were mounted with Vectashield™ (Vector Laboratories, Inc., Burlingame, CA), and cultures were examined with a microscope equipped for epifluorescence (Leica Inc., Deerfield, IL).

Competition Experiment and AChR Aggregation

Chick muscle cells were preincubated with 500 nM of cN25₇Fc for 4 h at 37°C. Conditioned medium from transiently transfected COS cells containing 100 pM cAgrin_{7A4B8} or c Δ N15Agrin_{A4B8} was added for 12 h at 37°C. AChRs were visualized and analyzed as described in Gesemann et al. (1995). Since cAgrin_{7A4B8}-induced AChR clusters were very small, aggregates with the longer axis of ≥ 1 μ m were included.

Results

The NH₂-terminal End of Agrin Is Sufficient to Bind to ECM

We have recently shown that recombinant full-length chick agrin (cAgrin; Fig. 1 *a*) binds to a solubilized mixture of ECM molecules, called Matrigel™. In contrast to full-length agrin, a fragment that lacks the first 130 amino acids from the NH₂ terminus (c Δ N15Agrin; Fig. 1 *a*) did not bind to this ECM preparation (Denzer et al., 1995). In this 130-amino acid-long stretch, a site of alternative mRNA splicing is found (Denzer et al., 1995; Tsen et al., 1995*b*). Both splice variants are capable of binding to Matrigel™, although with different binding strengths (Denzer et al., 1995). In the current report we have investigated the binding of the splice variant that includes the seven-amino acid-long insert (Fig. 1 *a*). This splice variant is selectively expressed by embryonic chick motor neurons (Denzer et al., 1995) and is also highly expressed in embryonic chick brain (Tsen et al., 1995*b*).

To see whether the NH₂-terminal region of agrin alone is sufficient to bind to Matrigel™, we engineered a cDNA construct that encodes an agrin fragment comprising the NH₂-terminal region and the first follistatin-like domain. To facilitate detection and purification of the fragment, it was fused to the constant region of a mouse IgG (Fc; Bowen et al., 1996), giving rise to the fragment termed cN25₇Fc (Fig. 1 *a*). A chimeric construct of the most COOH-terminal, 21-kD fragment of agrin from the marine ray (Smith et al., 1992) and the Fc part (r21Fc; Fig. 1 *a*) was used as a control. All recombinant proteins were efficiently synthesized and secreted from COS cells as shown by precipitation of ³⁵S-labeled proteins from conditioned medium of transiently transfected cells (Fig. 1 *b*). The high apparent molecular mass of 400–600 kD of cAgrin₇ and c Δ N15Agrin on SDS-PAGE is a result of heparan sulfate glycosaminoglycan (HS-GAG) chains that are attached to the 225-kD core protein (Denzer et al., 1995). The chimeric constructs, cN25₇Fc and r21Fc, were also secreted from the transfected cells, and they displayed the expected molecular mass of ~69–65 and 61 kD, respectively (Fig. 1 *b*). SDS-PAGE of the chimeric proteins under nonreducing conditions demonstrated that the intermolecular disulfide bonds of the Fc region dimerize the fragments (data not shown).

The binding to Matrigel™ was assayed using transiently

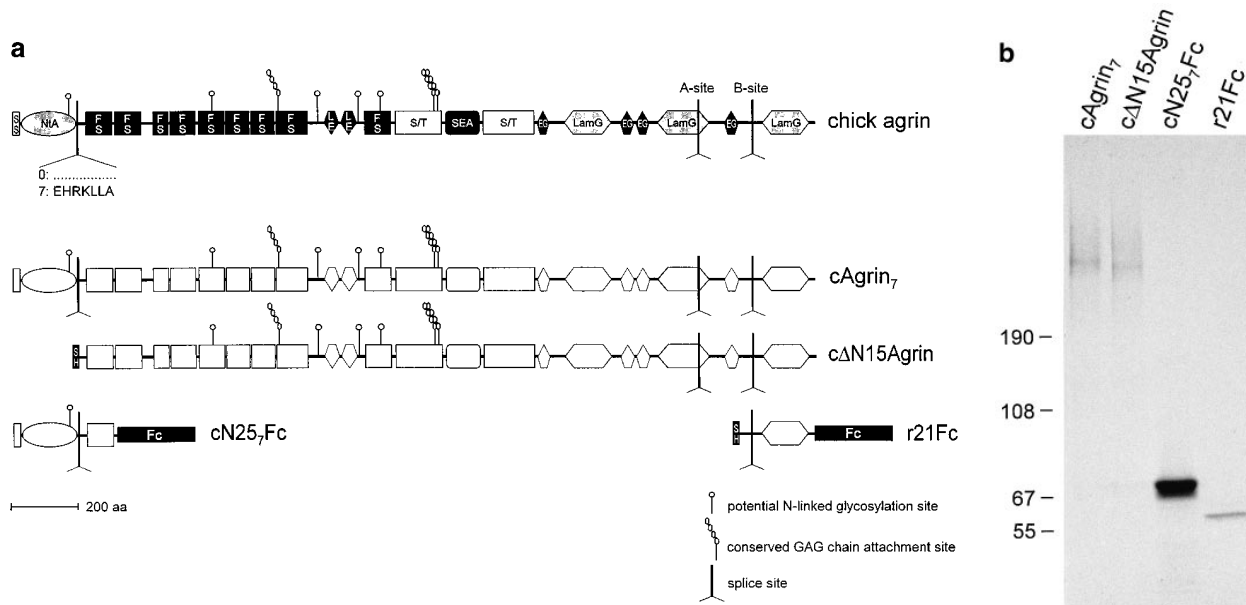


Figure 1. Structure and biochemical analysis of agrin constructs. (a) Structural organization of chick agrin and of fragments used in this study. Symbols and designations of individual domains are according to Bork and Bairoch (1995). *FS*, follistatin-like module; *EG*, EGF-like module; *LE*, laminin EGF-like module; *SEA*, module first found in sea urchin sperm protein, enterokinase and agrin; *LamG*, laminin G-like module. Furthermore, the fragment of the constant region of mouse immunoglobulin gamma heavy chain (*Fc*), the chick agrin signal sequence (*SS*), the hemagglutinin signal sequence (*SH*), the serine/threonine rich regions (*S/T*), potential N-linked glycosylation sites, conserved GAG side chain attachment sites, and the sites of alternative mRNA splicing are indicated. The NH₂-terminal region of agrin characterized in this study is named NtA domain. Notes: (1) Inserts at splice sites A and B are not specified. They are mentioned in the text if relevant for the experiment; and (2) the construct cΔN15Agrin was previously called cFull (Denzer et al., 1995; Gesemann et al., 1995) and covers the coding region of chick agrin as described by Tsim et al. (1992). (b) Autoradiogram of the ³⁵S-labeled agrin fragments depicted in a after precipitation from conditioned medium of transiently transfected COS cells. cAgrin₇ and cΔN15Agrin were immunoprecipitated with anti-cΔN15Agrin antibodies (Denzer et al., 1995). cN25₇Fc and r21Fc were precipitated with protein A-Sepharose. Proteins were separated by 3–12% SDS-PAGE. The two protein bands of cN25₇Fc are probably due to inefficient stop of protein translation at the COOH-terminal end of the construct. Molecular masses in kD of standard proteins are indicated.

transfected COS cells that were grown on tissue culture dishes coated with this basement membrane preparation. As previously shown (Denzer et al., 1995), cAgrin₇ was deposited on Matrigel™ (Fig. 2, a and a'), and no extracellular agrin-like immunoreactivity was observed with COS cells expressing cΔN15Agrin (Fig. 2, b and b'). Extracellular deposits of the recombinant proteins, cN25₇Fc and r21Fc, were visualized with fluorescein-conjugated goat anti-mouse IgG antibodies added to nonfixed, nonpermeabilized cells. The cells were subsequently permeabilized and stained with rhodamine-conjugated goat anti-mouse IgG antibodies to identify transfected cells. As shown in Fig. 2, c and c', cN25₇Fc was deposited on Matrigel™ around the transfected COS cells, which demonstrates that the NH₂-terminal region is sufficient to bind to this ECM mixture. The binding is not caused by the Fc part, as no deposits were seen with r21Fc-transfected cells (Fig. 2, d and d').

Laminin-1 Is a Binding Partner for Agrin

Matrigel™ consists of a mixture of several ECM molecules, such as collagen type IV, laminin-1, nidogen/entactin, and perlecan (Kleinman et al., 1982). To identify the components to which agrin binds, a solid-phase radioligand binding assay was used (Gesemann et al., 1996). Laminin-1

and perlecan were purified from the Engelbreth-Holm-Swarm mouse tumor (Timpl et al., 1979), and collagen type IV was isolated from human placenta (Weber et al., 1984; Ries et al., 1995). After coating the purified proteins onto the plastic surface of a microtiter plate, wells were incubated with 5 nM ¹²⁵I-cAgrin₇ or ¹²⁵I-cN25₇Fc and bound radioactivity was measured. Both agrin constructs bound to laminin-1 (Fig. 3), while no or only little binding was seen to perlecan and collagen type IV (data not shown). Since laminins bind to nidogen with high affinity ($K_d \sim 1$ nM; Fox et al., 1991), most laminin preparations also contain nidogen (Paulsson et al., 1987). To test whether the binding of agrin to laminin-1 was due to nidogen, we also measured binding of cAgrin₇ and cN25₇Fc to recombinant mouse nidogen, expressed in stably transfected HEK 293 cells (Fox et al., 1991). No binding was observed for both agrin constructs (Fig. 3). Hence, the binding of agrin to the laminin–nidogen complex is based on a direct interaction with laminin-1. As expected from previous experiments (Bowe et al., 1994; Campanelli et al., 1994; Gee et al., 1994; Sugiyama et al., 1994; Gesemann et al., 1996), cAgrin₇ also bound to α-dystroglycan, purified from chick skeletal muscle (Brancaccio et al., 1995; Fig. 3 a). In contrast, the NH₂-terminal fragment, cN25₇Fc, did not bind (Fig. 3 b). This is consistent with the notion that the binding site to this peripheral membrane protein resides in its COOH-terminal

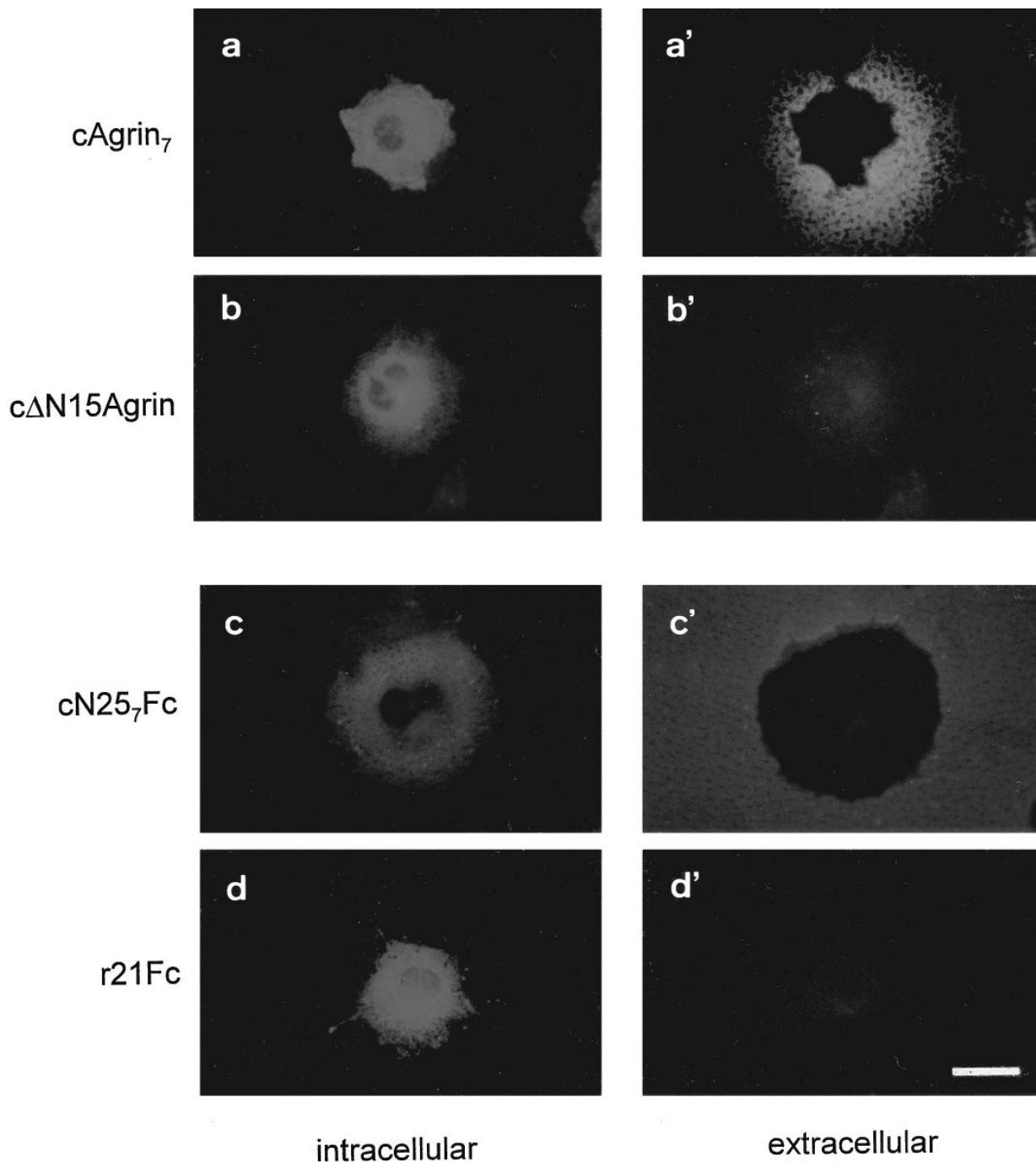


Figure 2. The NH₂-terminal end of agrin is sufficient to bind to MatrigelTM. COS-7 cells, which had been transiently transfected with cDNAs encoding the agrin constructs indicated on the left of each row, were grown on MatrigelTM. Deposition of recombinant protein onto MatrigelTM (right column) was monitored by staining living cells with the appropriate antibodies. To identify agrin-expressing cells, antibodies were added after permeabilizing the cells (*left*). To detect cAgrin₇ and cΔN15Agrin deposits, antiagrin antiserum followed by a fluorescein-conjugated secondary antibody was used (*a'* and *b'*). cN25₇Fc and r21Fc deposits were visualized with a fluorescein-conjugated goat anti-mouse IgG (*c'* and *d'*). Permeabilized cells were incubated with antiagrin mAb 5B1 followed by a rhodamine-conjugated secondary antibody (*a* and *b*) or with rhodamine-conjugated goat anti-mouse antibody (*c* and *d*). Only cells synthesizing cAgrin₇ and cN25₇Fc show extracellular deposits of recombinant protein (*a'* and *c'*). Bar, 40 μm.

half of agrin (Bowe et al., 1994; Campanelli et al., 1994; Gee et al., 1994; Sugiyama et al., 1994; Gesemann et al., 1996). Accordingly, only the laminin-binding site is common to cN25₇Fc and cAgrin₇. In addition, no binding of ¹²⁵I-r21Fc was observed to any of the components tested

(data not shown), which excludes a contribution of the Fc part in the binding of cN25₇Fc to laminin-1.

The strength of the binding of agrin to laminin-1 was measured by dose-response curves using ¹²⁵I-cAgrin₇. On immobilized laminin-1, half-maximal binding (EC₅₀) was

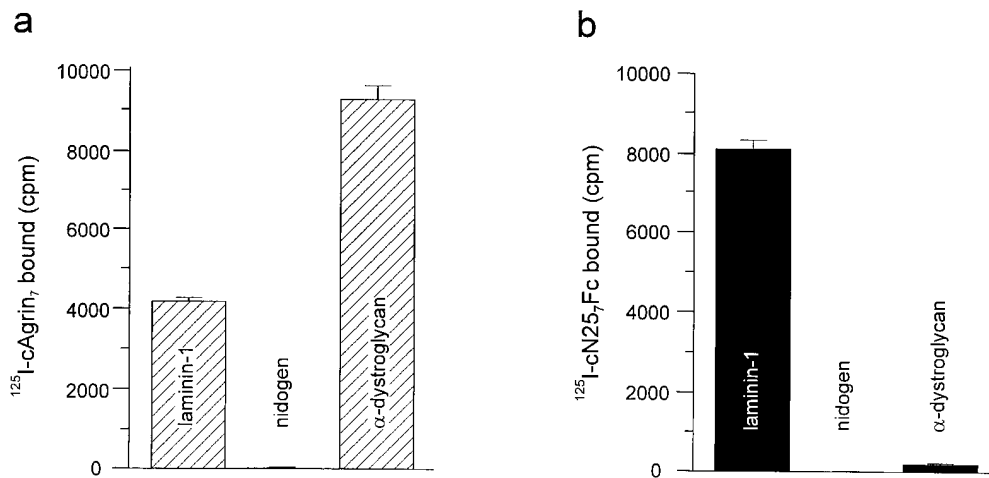


Figure 3. Agrin binds to laminin-1. Solid-phase radioligand binding assay with purified proteins. Mouse laminin-1, nidogen, and chick α -dystroglycan were immobilized in microtiter plates and subsequently incubated with ^{125}I -cAgrin₇ or ^{125}I -cN25₇Fc. Each value represents binding after subtraction of the background (BSA-coated wells). The values given are the mean \pm SD from one representative experiment with three independent measurements in the case of ^{125}I -cN25₇Fc and two measurements for ^{125}I -cAgrin₇. (a) 5 nM of ^{125}I -cAgrin₇ bound to α -dystroglycan and to laminin-1, but not to nidogen. Background counts were 89 ± 15 cpm. (b) 5 nM ^{125}I -cN25₇Fc bound to laminin-1 but not to α -dystroglycan and nidogen. Background counts were 65 ± 3 cpm. These experiments show that the binding of agrin to laminin is mediated by the NH₂-terminal end of agrin. Counts measured in a and b cannot be compared because cAgrin₇ and cN25₇Fc are iodinated to different extents.

reached at ~ 5 nM (Fig. 4), suggesting a rather strong interaction of agrin with laminin-1. This binding is of similar strength as the binding of laminin-1 to nidogen ($K_d \sim 1$ nM; Fox et al., 1991). At 4 nM ^{125}I -cAgrin₇, $90.3 \pm 0.5\%$ (mean \pm SEM; $n = 3$) of the binding was competed with 100 nM unlabeled cN25₇Fc, indicating that the majority of the binding of full-length agrin to laminin-1 is mediated by the NH₂-terminal end. Since agrin is an HSPG (Denzer et al., 1995; Tsen et al., 1995a) and laminin-1 is known to bind to heparin (Ott et al., 1982; Yurchenco et al., 1993), the residual binding of cAgrin₇ to laminin-1 could be due to the binding of the HS-GAG side chains of agrin to the heparin-binding site of laminin-1. Indeed, only the HS-GAG side chain-carrying construct cAgrin₇, but not cN25₇Fc, binds to the elastase fragment E3, which contains the major heparin-binding site of laminin-1 (Ott et al., 1982; data not shown). In summary, our data show that laminin-1, the laminin isoform present in MatrigelTM, is a basement membrane binding partner for agrin. Furthermore, they show that the interaction is mediated by the NH₂-terminal end of agrin, and we therefore propose that this part of agrin forms a structural unit (see also Discussion).

Agrin Expressed in Developing Chick Retina Binds to Laminin-1

The results presented so far show that recombinant chick agrin, expressed in mammalian cells, binds to laminin-1. If this binding were important for agrin's function in vivo, agrin purified from tissue should have the same binding properties. To test this, agrin was isolated by anion exchange chromatography from the vitreous fluid of embryonic day 14 chick eyes, a rich source for agrin and other molecules secreted from retinal cells (e.g., Ruegg et al., 1989; Halfter, 1993; Denzer et al., 1995). As a control, cAgrin₇ from conditioned medium of stably transfected HEK 293 cells was purified by the same procedure. When agrin-containing fractions were assayed by Western blot, agrin-like protein from vitreous fluid and cAgrin₇ had a high apparent molecular mass (Fig. 5 a, left), which is consistent with

previous results (Denzer et al., 1995). The bands with an apparent molecular mass of ~ 115 kD are probably due to proteolytic degradation similar to what has been reported elsewhere (e.g., Nitkin et al., 1987; Godfrey, 1991; Rupp et al., 1991; Denzer et al., 1995). The same samples were also probed with antiserum 245, raised against native chick heart laminin (Brubacher et al., 1991). No laminin-like immunoreactivity was observed in the samples containing cAgrin₇, derived from stably transfected HEK 293 cells, whereas immunoreactive bands were visible in the sample containing agrin from the vitreous fluid (Fig. 5 a, right). The strong immunoreactive band with the apparent molecular mass of 200 kD most likely corresponds to the β/γ

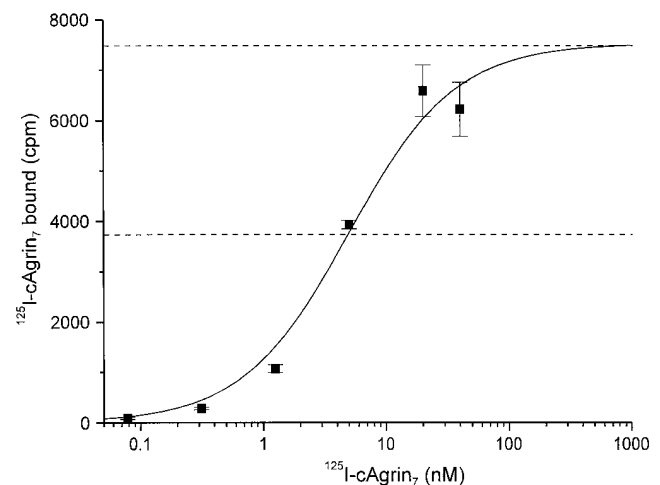
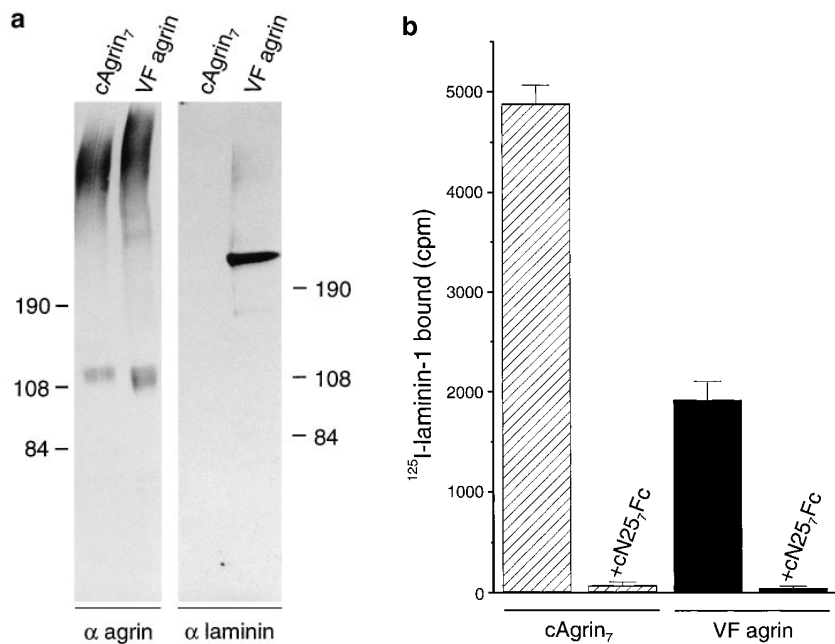


Figure 4. Binding of agrin to laminin-1 is of high affinity. Dose-response curve of the binding of ^{125}I -cAgrin₇ to laminin-1. Half-maximal binding was reached at ~ 5 nM, suggesting a rather strong binding of agrin to laminin-1. The binding curve shown results from one representative experiment. Background values (BSA-coated wells) are subtracted from each data point. Values shown are the mean \pm SEM from three measurements.



representative experiment and are the mean \pm SD of three measurements. Background counts (no agrin added; 132 ± 15 cpm) was subtracted. 100 nM cN257Fc (+ cN257Fc) inhibits binding of 125 I-laminin-1 to cAgrin₇ and VF agrin by more than 97%, indicating that the NH₂-terminal domain is responsible for the binding.

subunits of laminin isoforms, while the faint bands with the molecular masses of 400 and 150 kD could represent α subunits and nidogen, respectively (Brubacher et al., 1991). The copurification of chick laminin isoforms from the vitreous fluid with agrin by a procedure that is selective for highly charged anions is unexpected and may reflect the binding of laminin to agrin. Consistent with this idea, Halfter (1993) reported copurification of laminin with a chick HSPG, which later was shown to be agrin (Tsen et al., 1995a).

To test directly whether VF agrin is capable of binding to laminin-1, we performed a modified solid-phase radioligand binding assay. In this assay, purified cAgrin₇ or VF agrin was immobilized on microtiter plates that had been first coated with the antiagrin mAb 5B1 (Reist et al., 1987). To these wells, 5 nM of 125 I-laminin-1 was added and bound radioactivity was measured. As shown in Fig. 5 b, 125 I-laminin-1 bound to cAgrin₇ and VF agrin. In both cases, binding was competed by more than 97% by including 100 nM of unlabeled cN257Fc during the incubation with 125 I-laminin-1 (Fig. 5 b). These experiments show that agrin synthesized and secreted by cells of the chick retina can bind to laminin-1 and that this binding is also mediated by the NH₂-terminal domain of agrin. Although the same amount of agrin was added to the 5B1-coated microtiter wells, the amount of 125 I-laminin-1 bound to VF agrin was only half of that bound to cAgrin₇. The most likely explanation for this difference is that some of the binding sites in VF agrin may already be occupied by laminin isoforms from the vitreous fluid that copurified with VF agrin (see Fig. 5 a). In summary, chick agrin synthesized in vivo has the same laminin-binding properties as recombinant agrin, and its binding is also mediated by the NH₂-terminal domain.

Figure 5. Chick agrin synthesized in vivo contains the laminin-binding domain. (a) Western blot of recombinant cAgrin₇ and agrin purified from vitreous fluid (VF agrin). Transferred protein were stained with anti-agrin antibodies (α agrin) or antilaminin antibodies (α laminin). Agrin purified from vitreous fluid has approximately the same apparent molecular mass as recombinant full-length agrin. The band detected at \sim 115 kD most likely results from proteolytic degradation. No laminin-like immunoreactivity is associated with recombinant agrin (right). In contrast to this, the agrin-containing fractions from vitreous fluid are positive for laminin-like protein. Based on their apparent molecular mass and the specificity of the antiserum used (Brubacher et al., 1991), they may represent nidogen (\sim 150 kD), the β and γ chain (200 kD), and α chain (400 kD). Molecular masses of standard proteins are given in kD. (b) cAgrin₇ and agrin from vitreous fluid were bound to immobilized antiagrin mAb 5B1 and incubated with 125 I-laminin-1. The data result from one repre-

The Laminin-binding Domain Is Highly Conserved in Mouse and Human Agrin

The NH₂-terminal region required for the binding of agrin to laminin-1 has so far only been described in chick (Denzer et al., 1995). Full-length cDNA encoding rat agrin lacks this region and instead, the first follistatin-like domain is preceded by a sequence that has been proposed to serve as signal sequence (Rupp et al., 1991). To see whether the NH₂-terminal region of chick agrin is found in other species, we searched for homologous sequences using the BLAST algorithm (Altschul et al., 1990). Four expressed sequence tags, isolated from different tissues in mouse and in human (Lennon et al., 1996), closely matched this amino acid sequence. Sequencing of the clones confirmed that the deduced amino acid sequences of mouse and human agrin are highly homologous to each other and to chick agrin (Fig. 6). The homology starts at residue 26 of chick agrin, which corresponds to the predicted cleavage site for the signal sequence (Denzer et al., 1995). In the stretch from residue 26 to 149, 96% of the amino acids are identical between mouse and human, and 90% are identical between chick and the mammalian sequences (Fig. 6). This is by far the most highly conserved region in agrin (see also Tsim et al., 1992), suggesting that this domain may also confer binding to laminin in mammals.

Agrin Binds to Laminin Isoforms Expressed at the NMJ

The data presented so far show that agrin binds to mouse laminin-1. The laminins are a family of heterotrimeric glycoproteins composed of α , β , and γ chains (Burgeson et al., 1994; Timpl, 1996). Each laminin is characterized by its chain composition, for example laminin-1 is a trimer with the α 1, the β 1, and the γ 1 chains. Recent data suggest that

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CHICK  m g g s g a a a t l a l g l a l g l a l g g w a * N C P E R E 30
HUMAN  . . . . . a a c v l p g a g g T . . . . . A
MOUSE  . . . . . p g a s g T . . . . . A

CHICK  L Q R R E E E E A N V V L T G T V E E I M N V D P V H H T Y S 60
HUMAN  . E . . . . . L . . . . . Q . . . . .
MOUSE  . E . . . . . L . . . . . Q . . . . .

CHICK  C K V R V W R Y L K G K D I V T H E I L L D G G N K V V I G 90
HUMAN  . . . . . L . A R . S . . . . . S
MOUSE  . . . . . V . A Q . S . . . . .

CHICK  G F G D P L I C D N Q V S T G D T R I F F V N P A P O Y M W 120
HUMAN  . . . . . . . . . . . . . . . . . . . . . . . . . . . . . . . . .
MOUSE  . . . . . . . . . . . . . . . . . . . . . . . . . . . . . . . . .
RAT    . . . . . M P P L P L E H R P R Q E

CHICK  P A H R N E L M L N S S L M R I T L R N L E E V E H C V E E 150
HUMAN  . . . K . . . . . . . . . . . . . . . . . . . . . . . . . . . . .
MOUSE  . . . K . . . . . . . . . . . . . . . . . . . . . . . . . . . . .
RAT    . G A S M L V R Y F M I P C N I C . I L . A T S T L G F A V

CHICK  H R K L L A D K P N S Y P T Q T P P T P R D A C 173
HUMAN  - - - - - . . . . . G T H F . P V . . . . . P . . . .
MOUSE  - - - - - . . . . . G I H F . A A . S M . P . V . .
RAT    L L S . S N Y . . G I H F . P A . . . . . P . V .

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Figure 6. The laminin-binding domain is present in mouse and human agrin. Deduced amino acid sequences of the expressed sequence tags (Lennon et al., 1996) from mouse and human agrin were aligned to the first 173 amino acids of chick agrin. In addition, the first 67 amino acids of rat agrin are shown. Amino acids identical to chick agrin are denoted with dots. The small letters in the chick sequence represent the proposed signal sequence (Denzer et al., 1995) with the initiator methionine (*bold*) and the proposed signal sequence cleavage site (*). Underlined amino acids show the position of the tryptic peptide derived from a HSPG of bovine kidney (Hagen et al., 1993). Homology between chick and human starts after the signal sequence cleavage site of chick (Cys 26). Amino acids between Cys 26 and Glu 149 are almost 90% identical between human, mouse, and chick. Note that the amino

laminin-1 is not expressed in muscle basal lamina but instead is replaced by laminin-2 and -4 (Schuler and Sorokin, 1995). Laminin-2 ($\alpha 2$, $\beta 1$, $\gamma 1$) is present early in development throughout the extracellular matrix of muscle fibers. In adult muscle, it is expressed in the extrasynaptic region of the basal lamina. Laminin-4 (also called s-laminin; Hunter et al., 1989), in which the $\beta 1$ chain is replaced by the $\beta 2$ chain, is expressed later in development and localizes to the synaptic portion of the muscle cell basal lamina (Sanes et al., 1990).

Since we were interested in whether agrin's NH₂-terminal domain would also mediate the tethering to muscle cell basal lamina, we measured binding of cAgrin₇ and cN25₇Fc to laminin-2 and -4. The source for laminin-2 and -4 was a laminin preparation isolated from adult chick heart after EDTA extraction and sequential purification by wheat germ agglutinin Sepharose and immunoaffinity chromatography (Brandenberger and Chiquet, 1995). As shown by Western blot analysis using antiagrin antibodies, this laminin preparation contained substantial amounts of agrin-like protein (Fig. 7 *a*). As this laminin preparation did not

acid sequences of mouse and human agrin that precede the proposed signal peptide cleavage site (*) are not homologous to chick and consist mainly of hydrophobic amino acids. Thus, as in chick these stretches might be part of a signal sequence (*lowercase letters*). The homology between chick and rat begins at Asp 157 immediately after the boundary where chick is alternatively spliced. The human and mouse sequences lack the seven-amino acid-long insert at this splice site (*dashes*). These sequence data are available from GenBank/EMBL/DBJ under accession number U84406 (human agrin) and U84407 (mouse agrin).

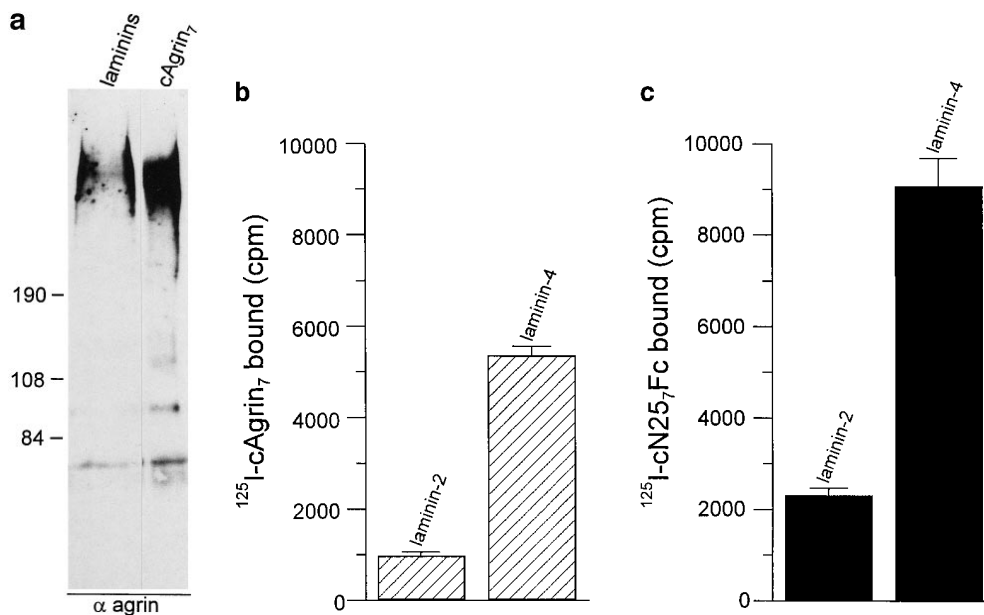


Figure 7. Agrin binds to laminin isoforms expressed in muscle. (*a*) Immunoblot using antiagrin antibodies of laminin isoforms purified from EDTA extracts of adult chick heart by immunoaffinity column with the anti- $\gamma 1$ subunit-specific mAb 11B7 (Brandenberger and Chiquet, 1995). Agrin-like protein with the same apparent molecular mass as cAgrin₇ is detected in the laminin preparation. Since no α -dystroglycan was detected in the same laminin preparation (data not shown), the copurification of agrin suggests that laminins are associated with agrin in muscle tissue. (*b* and *c*) Agrin binds directly to purified chick laminin-2 and -4. Equal amounts of the two

laminins were immobilized on microtiter plates and incubated either with 5 nM of iodinated cAgrin₇ (*b*) or cN25₇Fc (*c*). Values given are the result of one representative experiment and represent the mean \pm SD of three measurements where the background (BSA-coated wells) has been subtracted. Background values were 163 ± 24 cpm in *b* and 75 ± 11 cpm in *c*. The binding of cAgrin₇ to laminin-4 is fivefold stronger than to laminin-2 (*b*) and a similar difference in the binding (fourfold) is observed with cN25₇Fc (*c*). Note that the difference in the counts measured with ¹²⁵I-cAgrin₇ and ¹²⁵I-cN25₇Fc is due to different iodination efficiencies.

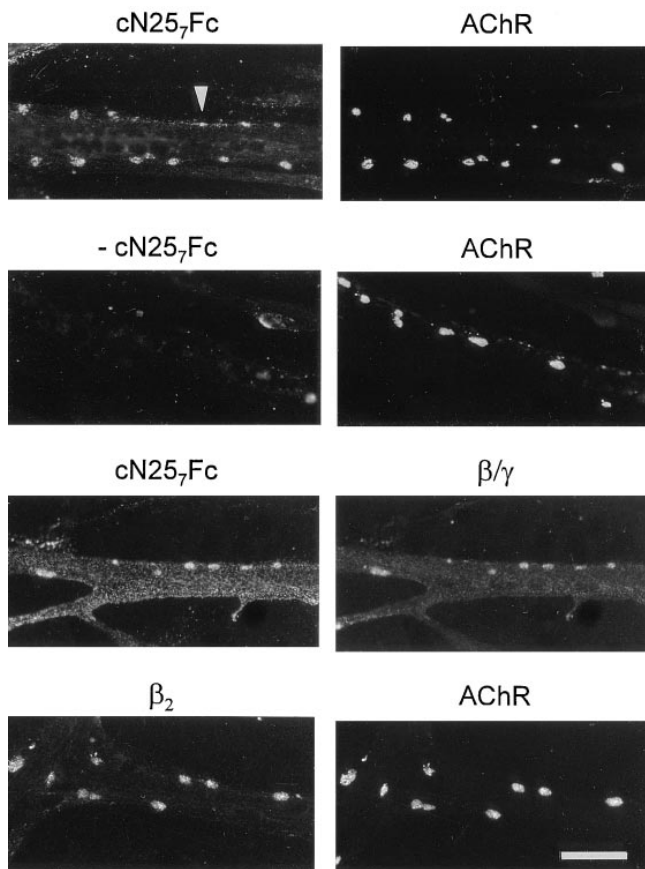


Figure 8. Binding sites for agrin on cultured chick myotubes colocalize with AChRs and laminin. Cultured chick myotubes were induced to form AChR clusters with 200 nM c21_{B8}. Simultaneously, 20 nM cN25₇Fc were included. AChR aggregates were stained by rhodamine- α -bungarotoxin, and cN25₇Fc bound to the myotubes was visualized with biotinylated goat anti-mouse IgG followed by fluorescein-conjugated streptavidin. cN25₇Fc is concentrated in AChR clusters and is distributed along the edges of the myotubes (*arrowhead*). No staining is seen in the absence of this fragment (-cN25₇Fc). Consistent with the idea that cN25₇Fc binds to laminin, the distribution of myotube-bound cN25₇Fc resembles the staining pattern obtained with anti- β / γ -specific antiserum 648 (β / γ). The β 2 chain of laminin, also called s-laminin, is concentrated in AChR clusters. In light of the colocalization of cN25₇Fc and AChR clusters, laminin-4 (α 2, β 2, γ 1) is a binding partner of agrin in these clusters. Bar, 40 μ m.

contain α -dystroglycan, detected by a transfer overlay assay using iodinated chick agrin (data not shown), we conclude that the copurification of agrin with laminin isoforms from cardiac muscle is most likely a consequence of the association of agrin with the laminins. To test directly whether laminin-2 and -4 are binding proteins for agrin, we performed a solid-phase radioligand binding assay using purified laminin isoforms. With the same amount of laminin-2 and -4 coated on the microtiter wells, 5 nM of ¹²⁵I-cAgrin₇ gave a clear signal on laminin-2 and a fivefold stronger signal on laminin-4 (Fig. 7 *b*). Like cAgrin₇, the NH₂-terminal fragment, cN25₇Fc, also bound approximately four times more strongly to laminin-4 than to laminin-2 (Fig. 7 *c*). The weaker signal on laminin-2 was not due to a contamination with laminin-4, since this isoform

represents less than 5% in the laminin-2 preparation (Brandenberger et al., 1996). A difference in the coating efficiency between laminin-2 and -4 was also excluded because equal amounts were immobilized when tested with an mAb specific for the α 2 chain (Brandenberger et al., 1996). In summary, full-length chick agrin binds to both laminin-2 and -4, but at this particular concentration the interaction with the extrasynaptic laminin-2 is of lower apparent affinity.

Many proteins that are highly concentrated at the NMJ in vivo are also found in AChR clusters in vitro (for review see Bowe and Fallon, 1995). These include proteins of the ECM, such as HSPG (Wallace, 1989) and laminin (Nitkin and Rothschild, 1990). Consequently, the laminin-binding fragment, cN25₇Fc, should colocalize with AChR clusters. To test this, AChR aggregation was induced on cultured chick myotubes with 200 nM c21_{B8}, the minimal COOH-terminal agrin fragment required for AChR aggregation (Gesemann et al., 1995), and 20 nM of cN25₇Fc was included. After 16 h, AChR clusters were visualized with rhodamine- α -bungarotoxin and myotube-bound cN25₇Fc was stained with biotinylated goat anti-mouse IgG followed by fluorescein-conjugated streptavidin. As shown in Fig. 8 (*first row*), cN25₇Fc localized to the agrin-induced AChR clusters. cN25₇Fc also displayed a more widespread distribution, often along the edge of the myotubes (Fig. 8, *arrowhead*). In myotubes, where AChR aggregation had been induced with c21_{B8}, but no cN25₇Fc was included, no specific staining was seen (Fig. 8, *second row*). These results illustrate that binding sites for the NH₂-terminal fragment of agrin are concentrated in AChR clusters but are also found on the remaining surface of the myotube.

To look at the relationship between the localization of cN25₇Fc and laminin, antiserum 648, directed against the β and γ chains of laminin-2 and -4, was used. Examination of many myotubes showed that the distribution of myotube-bound cN25₇Fc matched laminin-like immunoreactivity (Fig. 8, *third row*). In addition, staining with the mAb 11B7, directed against the γ 1 chain of chick laminins (Brandenberger and Chiquet, 1995), gave the same staining pattern (data not shown). Since these antibodies do not distinguish between laminin-2 and -4, we also applied mAb C4, recognizing the β 2 chain, to localize the synapse-specific laminin isoform on myotubes (Hunter et al., 1989). In Fig. 8 (*fourth row*), staining with the β 2-specific mAb C4 is shown. As reported for spontaneous AChR clusters on mouse C2 myotubes (Martin et al., 1995), the β 2 chain was concentrated in AChR clusters on chick myotubes. Interestingly, no or only little laminin β 2-like immunoreactivity was detected outside of the clusters. This differs from the pattern obtained with mAbs specific for the γ 1 or the α 2 chain of laminin, where immunoreactivity was also seen along the edge of myotubes (data not shown). In summary, these experiments demonstrate that the binding sites for agrin coincide well with the distribution of laminin on cultured myotubes, and they suggest that laminin-4 mediates binding of agrin to AChR clusters.

Binding of Agrin to Laminin Alters the Size of the AChR Aggregates

AChR aggregates induced by the active full-length splice

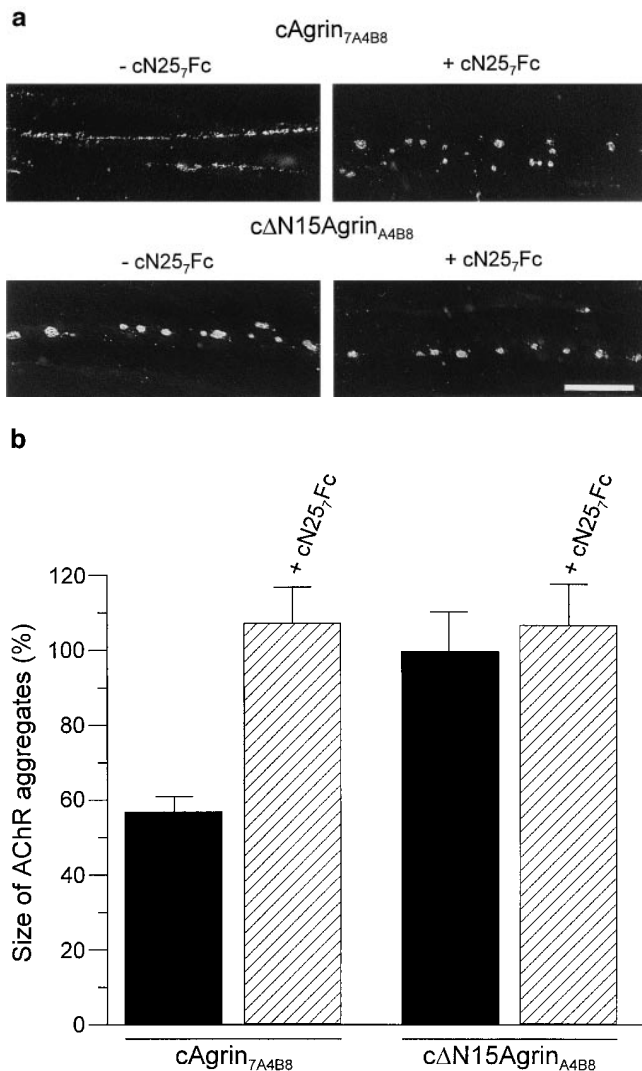


Figure 9. The NH₂-terminal, laminin-binding domain of agrin causes AChR clusters to be small. (a) Fluorescence micrographs of cultured chick myotubes labeled with rhodamine- α -bungarotoxin. Myotubes were grown for 12 h in the presence of conditioned medium of transiently transfected COS cells containing either 100 pM cAgrin_{7A4B8} or cΔN15Agrin_{A4B8}. AChR clusters induced by cAgrin_{7A4B8} are considerably smaller than those induced by cΔN15Agrin_{A4B8}. In the presence of cN25₇Fc, the size of cAgrin_{7A4B8}-induced AChR clusters increases. (b) Quantification of the effect on the size of the AChR aggregates. The presence of 500 nM cN25₇Fc (+ cN25₇Fc) increases the size of the AChR clusters induced by cAgrin_{7A4B8} twofold and makes them indistinguishable from AChR clusters induced by cΔN15Agrin_{A4B8}. In contrast, addition of 500 nM cN25₇Fc does not alter the size of cΔN15Agrin_{A4B8}-induced AChR aggregates. The result (mean \pm SEM) of one representative experiment is shown where the size of AChR clusters in 20 myotube segments was determined. Bar, 50 μ m.

variant cAgrin_{7A4B8} are more than twofold smaller than those induced by the fragment cΔN15Agrin_{A4B8} lacking the first 130 NH₂-terminal amino acids (Denzer et al., 1995). Aggregation of AChRs induced by agrin is mainly based on the lateral migration of diffusely distributed molecules (Godfrey et al., 1984). We have speculated that the bind-

ing of cAgrin_{7A4B8} to ECM influences this process by immobilizing agrin in the ECM. If this were so, inhibition of the binding of cAgrin_{7A4B8} to muscle laminins should result in AChR clusters with the same size as AChR clusters induced by cΔN15Agrin_{A4B8}. Myotubes incubated with cAgrin_{7A4B8} had considerably smaller AChR aggregates than those incubated with cΔN15Agrin_{A4B8} (Fig. 9 a). When available laminin-binding sites for full-length agrin were blocked by 500 nM cN25₇Fc, cAgrin_{7A4B8}-induced AChR clusters became indistinguishable from those induced by cΔN15Agrin_{A4B8} (Fig. 9 a). In contrast to this, 500 nM of cN25₇Fc did not alter the shape of AChR aggregates induced by cΔN15Agrin_{A4B8}. Quantification of this effect with a computerized image analysis system confirmed that the presence of cN25₇Fc increased the average size of AChR aggregates induced by cAgrin_{7A4B8} approximately twofold, while the average size of cΔN15Agrin_{A4B8}-induced AChR clusters was not affected (Fig. 9 b). These experiments provide evidence that the smaller size of the AChR clusters induced by full-length agrin is mainly based on its binding to laminin isoforms expressed by the myotubes.

Discussion

Agrin is a large, multidomain protein that is associated with basement membranes in many tissues (for review see Denzer et al., 1996). In the current study, we investigated the molecular basis of agrin's binding to basement membrane. We found that agrin binds with high affinity to laminin isoforms, including those that are concentrated at the NMJ, and we have mapped the main laminin-binding site to the NH₂-terminal end of agrin. Agrin has been shown to be a key regulator of synaptic differentiation at the NMJ (McMahan, 1990; Gautam et al., 1996), and thus we will mainly discuss the significance of the binding of agrin to laminin in this process.

The Laminin-binding Domain of Agrin Defines a Novel Module

At the outset of the current work, it was known that the binding of recombinant chick agrin to Matrigel™ requires the first 130 amino acids at the NH₂ terminus of full-length agrin (Denzer et al., 1995). We now find that laminin-1, one component of Matrigel™, serves as a binding partner for chick agrin. The binding of recombinant cAgrin₇ (Fig. 1 a) to laminin-1 is of high affinity (EC₅₀ of \sim 5 nM; Fig. 4) and is mainly carried by the NH₂-terminal end of agrin as an excess of the NH₂-terminal fragment, cN25₇Fc, competes the binding of full-length agrin by more than 90%.

Although cN25₇Fc includes the first follistatin-like domain, this domain is not necessary for the binding to laminin-1 (Denzer, A.J., and M.A. Ruegg, unpublished observation). The stretch preceding the follistatin-like domain is the most highly conserved region of agrin (Fig. 6). This high homology and the fact that this region in chick agrin mediates binding to all laminin isoforms so far tested (i.e., laminin-1, -2, and -4) strongly suggest it being important for the integration of agrin into the scaffold of ECM molecules formed by the self-assembled laminins and collagen type IV. Since no homology to any so far defined modules

was found using BLAST or FASTA algorithms (Altschul et al., 1990; Pearson and Lipman, 1988), we propose to call this novel module NtA domain (for NH₂-terminal domain in agrin; see also Fig. 1 a).

The binding of the NtA domain of agrin to laminin-1 is confined to a particular region in the upper part of the triple coiled-coil domain of laminin-1, as shown by the binding of cN25₇Fc to proteolytic fragments of laminin-1 and visualization of this interaction by electron microscopy after rotary shadowing (Denzer, A.J., T. Schulthess, C. Fauser, B. Schumacher, J. Engel, and M.A. Ruegg, manuscript in preparation). These results show that the binding of agrin to laminin-1 is mediated by specific domains in both molecules and they support the result of this study (Fig. 4) that the agrin–laminin interaction is of high affinity.

Binding of Agrin to Laminin and Its Role in AChR Aggregation

Induction and maintenance of postsynaptic differentiation at nerve–muscle contacts *in vivo* is triggered by neural agrin in a restricted area of muscle fibers (McMahan, 1990). The simplest explanation for this local action of neural agrin is that it becomes trapped in the extracellular matrix near its release site, the tip of the motor neuron's axon. Although we cannot exclude that other molecules are also involved in this process, we propose that the local immobilization of neural agrin is mainly mediated by its binding to laminin isoforms expressed by muscle fibers. Consistent with this view, the fragment cN25₇Fc binds to purified laminin-2 and -4 (Fig. 7), the two main laminin isoforms expressed in developing muscle fibers (Chiu and Sanes, 1984; Sanes et al., 1990). In addition, agrin-like protein copurifies with laminin isoforms (Fig. 7 a), and cN25₇Fc colocalizes with laminin and AChR clusters on cultured myotubes (Fig. 8). Similarly, laminin-like immunoreactivity colocalizes with agrin-like protein in chick embryo hindlimb muscle *in vivo* and *in vitro*, and both proteins are enriched in AChR clusters (Godfrey et al., 1988b; Nitkin and Rothschild, 1990). The binding of neural agrin to laminin would also explain earlier observations that motor neuron–derived agrin associates with the earliest AChR clusters in frog nerve–muscle cocultures (Cohen and Godfrey, 1992) and that it is deposited onto culture substrates containing laminin (Cohen et al., 1994).

We have shown that the size of agrin-induced AChR aggregates is affected by the NtA domain of agrin (Denzer et al., 1995; Fig. 9). On cultured chick myotubes, the size of cΔN15Agrin_{A4B8}-induced AChR clusters is indistinguishable from the size of the clusters induced by c21_{B8}, the minimal fragment required for AChR aggregation that does not bind to α-dystroglycan (see Figs. 8 and 9; Gese-mann et al., 1996). Hence, unlike the binding of agrin to α-dystroglycan, the NtA domain has a clear effect on the size of the AChR clusters. One explanation for this phenomenon may be that agrin becomes immobilized on the muscle cell surface by its binding to laminin and thereby prevents small AChR clusters from fusing (see also Discussion in Denzer et al., 1995).

We also think that the binding to laminin is the molecular basis that, after degeneration of the nerve terminals and the muscle fibers, AChR-aggregating activity and agrin-

like immunoreactivity is maintained at former synaptic sites for several weeks (Burden et al., 1979; Reist et al., 1987). At least *in vitro*, the tight association of agrin with muscle cell basal lamina requires the NtA domain; cultured chick myotubes that are incubated for only 30 min with AChR-aggregating full-length agrin (cAgrin_{7A4B8}), subsequently washed and further incubated for 15 h in agrin-free culture medium, show agrin-induced AChR aggregates. In contrast to this, no AChR clusters are induced with the same paradigm using cΔN15Agrin_{A4B8} or any other active, COOH-terminal agrin fragment (Denzer, A.J., and M.A. Ruegg, unpublished data). Our observation that cΔN15Agrin_{A4B8} is not capable of inducing AChR clusters by short-term incubation is similar to results of Wallace (1988), who showed that maintenance of AChR clusters *in vitro* needs the continuous presence of agrin. Since his agrin preparation mainly contained proteolytic fragments of the COOH-terminal half of agrin (Nitkin et al., 1987), these fragments also lack the NtA domain.

Binding of Agrin to Laminin Isoforms and Neuromuscular Junction Development *In Vivo*

Upon formation of primary muscle fibers, β1-containing laminin isoforms are expressed throughout the muscle cell basal lamina (Chiu and Sanes, 1984; Schuler and Sorokin, 1995). At later stages of synaptogenesis, the β2 chain of laminin (s-laminin) accumulates in synaptic basal lamina, and the β1 chain is displaced from this region and continues to be expressed extrasynaptically (Chiu and Sanes, 1984; Hunter et al., 1989; Sanes et al., 1990). In contrast to the β chains, the α2 and γ1 chains are found throughout the muscle fiber basal lamina (Sanes et al., 1990). These results strongly suggest that laminin-2 (α2, β1, γ1) and laminin-4 (α2, β2, γ1) are the laminin isoforms in the extrasynaptic and synaptic basal laminae, respectively. We find that, at one particular concentration, laminin-4 binds more strongly to agrin than laminin-2 (Fig. 7). Hence, agrin may preferentially bind to laminin-4, and this may be the basis of the tight association of neural agrin with synaptic basal lamina throughout adulthood.

Similarly, the phenotype of mice that are deficient of the β2 chain of laminin may, at least partially, be based on alterations of agrin. In these mice, NMJs are formed on schedule, but maturation of pre- and postsynaptic specializations is impaired, and terminal Schwann cells penetrate partially the synaptic cleft (Noakes et al., 1995). Although the loss of the β2 chain is compensated by continued expression of the β1 chain at the NMJ (Martin et al., 1996), this may not be sufficient to tether agrin to the NMJ. Since agrin has been shown to influence the formation of pre-synaptic specializations (Campagna et al., 1995; Gautam et al., 1996), alterations in the binding of agrin to the NMJ may also explain the abnormalities in presynaptic specialization in the β2-deficient mice.

For the current studies, we used agrin isoforms that include a 7–amino acid insert at the NH₂-terminal splice site. Motor neurons in the developing chick spinal cord contain agrin transcripts encoding this agrin variant, whereas the majority of agrin mRNA in nonneuronal cells codes for agrin without the insert (Denzer et al., 1995; Tsen et al., 1995b). Interestingly, the cDNA clones encoding mouse

and human agrin, which are derived from nonneuronal tissue, all encode the splice variant lacking the insert (Fig. 6). Preliminary results show that both splice variants bind to laminin-1 (Denzer, A.J., and M.A. Ruegg, unpublished observation). Since there appears to be a difference in the binding of the two splice variants to Matrigel™ (Denzer et al., 1995), it will be interesting to see whether splicing at this site influences binding of agrin to different laminin isoforms.

Little is known about the function of agrin isoforms without AChR-aggregating activity. Since these inactive isoforms bind more strongly to α -dystroglycan than AChR-aggregating isoforms and since they also bind to laminin, the inactive isoforms might have a structural role to link the basement membrane with the underlying cytoskeleton via the dystrophin-glycoprotein complex (for reviews see Campbell, 1995; Worton, 1995). We have now generated the tools that will enable studies on the physiological significance of the binding of agrin to laminin isoforms and to the different binding proteins expressed on the muscle cell membrane, like α -dystroglycan.

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