CASK Participates in Alternative Tripartite Complexes in which Mint 1 Competes for Binding with Caskin 1, a Novel CASK-Binding Protein

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CASK, an adaptor protein of the plasma membrane, is composed of an N-terminal calcium/calmodulin-dependent protein (CaM) kinase domain, central PSD-95, Dlg, and ZO-1/2 domain (PDZ) and Src homology 3 (SH3) domains, and a C-terminal guanylate kinase sequence. The CaM kinase domain of CASK binds to Mint 1, and the region between the CaM kinase and PDZ domains interacts with Velis, resulting in a tight tripartite complex. CASK, Velis, and Mint 1 are evolutionarily conserved in Caenorhabditis elegans, in which homologous genes (called lin-2, lin-7, and lin-10) are required for vulva development. We now demonstrate that the N-terminal CaM kinase domain of CASK binds to a novel brain-specific adaptor protein called Caskin 1. Caskin 1 and a closely related isoform, Caskin 2, are multidomain proteins containing six N-terminal ankyrin repeats, a single SH3 domain, and two sterile α motif domains followed by a long proline-rich sequence and a short conserved C-terminal domain. Unlike CASK and Mint 1, no Caskin homolog was detected in C. elegans. Immunoprecipitations showed that Caskin 1, like Mint 1, is stably bound to CASK in the brain. Affinity chromatography experiments demonstrated that Caskin 1 coassembles with CASK on the immobilized cytoplasmic tail of neurexin 1, suggesting that CASK and Caskin 1 coat the cytoplasmic tails of neurexins and other cell-surface proteins. Detailed mapping studies revealed that Caskin 1 and Mint 1 bind to the same site on the N-terminal CaM kinase domain of CASK and compete with each other for CASK binding. Our data suggest that in the vertebrate brain, CASK and Velis form alternative tripartite complexes with either Mint 1 or Caskin 1 that may couple CASK to distinct downstream effectors.

Key words: CASK; synapse; scaffold; lin-2; Mint 1; Caskin; neurexin; syndecan; Velis

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Velas bind to a sequence between the CaM kinase and PDZ domains (Butz et al., 1998). In *C. elegans*, the lin-2–lin-7–lin-10 complex is required for the correct targeting of epidermal growth factor (EGF) receptors to the basolateral plasma membrane of vulval precursor cells (Kaech et al., 1998). In mice, a transgenic insertion mutant that changes CASK expression causes a cleft palate syndrome, suggesting that CASK also has a role in vertebrate development (Laverty and Wilson, 1998); however, its precise function and mechanism of action are unknown.

In addition to Mint 1 and Velas, CASK binds to calmodulin in a Ca$^2+$-dependent manner via its N-terminal CaM kinase domain (Hata et al., 1996) and to protein 4.1 via a C-terminal sequence (Cohen et al., 1998; Biederer and Sudhof, 2001). The resulting CASK–cell-surface protein–protein 4.1 complex nucleates actin assembly and may mediate the attachment of actin microfilaments to the cytoplasmic tails of cell-surface receptors (Biederer and Sudhof, 2001). Furthermore, the SH3 domain of CASK binds to Ca$^2+$–calmodulin (Maximov et al., 1999), and the SH3 domains and guanylate kinase domains of CASK andDlg interact intramolecularly and intermolecularly (Nix et al., 2000). Finally, the guanylate kinase domain of CASK binds to the transcription factor Tbr1-1 and may translocate from the synapse into the nucleus, in which it could function as a coactivator of Tbr1-1 in transcription (Hsueh et al., 2000).

The interactions of CASK support the notion that it functions as a scaffolding protein, possibly by binding sequentially to different proteins. Such a function would imply that multiple proteins should bind competitively to the same domains. In the present study, we describe a novel protein called Caskin 1 that binds to the CaM kinase domain of CASK in competition with Mint 1, suggesting that CASK participates in alternative complexes with Mint 1 or Caskin 1.

**MATERIALS AND METHODS**

**Purification of CASKIN.** Rat brain proteins were affinity-purified on immobilized glutathione-3-transferase (GST)–CASK fusion protein essentially as described previously (Butz et al., 1998). Three proteins were detectable on Coomassie-stained gels: Mint 1, a 180 kDa protein called Caskin 1 in this study, and a 225 kDa protein. The 180 kDa and 225 kDa bands were eluted from the gel and digested by trypsin, and tryptic peptides were purified by HPLC and sequenced by Edman degradation essentially as described previously (Hata et al., 1993). The 180 kDa peptide sequences did not match any protein in the database except for KIAA and expressed sequence tag (EST) sequences (see Results), whereas the 225 kDa protein was identified as the rat homolog of NEDD4-like ubiquitin ligase 1 (GenBank accession #BAB13352).

cDNA cloning and sequencing. GenBank searches identified a random human brain cDNA (KIAA1306) and human EST clones that contained the sequences of the peptide fragments. To determine the complete structure of Caskin 1, PCR products were performed with rat brain cDNA and degenerate oligonucleotides based on the human sequences. The products were used to screen a rat brain cDNA library in A-ZAP II by standard cDNA cloning methods (Sambrook et al., 1989). Fifty positive clones were isolated, and the full-length sequence was assembled from their combined nucleotide sequences and translated into amino acid sequences. Human Caskin 1 (KIAA1306), human Caskin 2 (KIAA1306), and mouse Caskin 2 (AW321773.1, AA100308.1, AA166316.2, AA678547.1, A146644.1, AA216903.1, A14253971.1, B1837871, and BG872937) clones were identified in the EST databank using basic local alignment search tool searches with rat Caskin 1 sequence. The cDNA sequences were submitted to GenBank (accession # AF451975–AF451978).

**Construction and expression of bacterial and eukaryotic expression vectors.** Expression vectors for GST fusion proteins were constructed in pGEX-4G (Ggen and Dixon, 1991) by standard procedures (Sambrook et al., 1989). The following GST fusion proteins were produced by the indicated vectors (residue numbers in parentheses): Caskin 1: GST–Caskin 1–437, pGEX–Caskin 1 Neol–BglII–4 (1–378); GST–Caskin 1 378–471, pGEX–Caskin 1–SS (375–471); GST–Caskin 1 374–546, pGEX–Caskin 1–sterile e motif 1 (SAM1) (374–540); GST–Caskin 1 359–659, pGEX–Caskin 1 BglII–SacI–1 (374–659); CASK: GSTCASK, pGEX–CASK full-length; GST–CASK 1–275, pGEX–Cas9–3 (1–275); GST–CASK 1–310, pGEX–CASK ΔCaM (1–310); GST–CASK 1–337, pGEX–Cash3–13 (1–337); GST–CASK 98–337, pGEX–CASK BglII–CaM (98–337); GST–CASK 129–328, pGEX–CASK–P–CaM (237–337); GST–CASK–25 (328–432); GST–Caskin 1 1 (116–432); and Neurexin I: GST–Neurexin I (NIX–CaM) was described previously (Biederer and Sudhof, 2000). For maltose-binding protein (MBP)–Caskin 1 fusion protein, the BamHI–HindIII fragment from pGEX–Caskin 1 BglII–SacI–1 (374–659) was subeloned into the same site of pMalC2 (New England Biolabs, Beverly, MA). A full-length eukaryotic expression vector for Caskin 1 (pCMV–CB16) was constructed by cloning the EcoRI insert from clone pBlue–CB16 into the same site of pCMV5. Bacterial expression of GST fusion proteins and COS cell transfections have been described previously (Okamoto and Sudhof, 1997). MBP–Caskin 1 fusion proteins were immobilized on amyllose resin and purified by 10 mM maltose/PBS elution.

**GST-pulldown experiments.** GST fusion proteins were expressed in *Escherichia coli* and immobilized on glutathione agarose (Sigma, St. Louis, MO) using standard procedures. Rat forebrains were homogenized in a pestle tissue grinder at slow speed in solubilization buffer (25 mM HEPES–NaOH, pH 7.4, 125 mM K-acetate, 5 mM MgCl2, 0.32 M sucrose, and 1% Triton X–100). Proteins solubilized from rat forebrain were incubated with immobilized GST fusion proteins for 14 hr at 4°C under mild agitation, and bound proteins were analyzed after extensive washing with the solubilization buffer as described previously (Butz et al., 1998; Biederer and Sudhof, 2000).

**Immunoprecipitations.** Immunoprecipitations were performed from rat brain homogenates as previously (Butz et al., 1998). To characterize the CASK–Caskin 1 interaction further, immunoprecipitations were subjected to different washing conditions; after five washes using lysis buffer (20 mM imidazole, pH 6.8, 0.1 mM NaCl, 5 mM EDTA, 1 mM DTT, 0.5% Triton X–100, 1 mM PMSE, 10 mg/l leupeptin, 1 mg/l aprotinin), bound immune complexes were treated with lysis buffer supplemented with final concentrations of either 0.2, 0.5, or 1 mM NaCl, 0.6 mM KI, or 0.5% SDS/2% Triton X–100.

**Antibodies.** Most antibodies have been described previously (Okamoto and Sudhof, 1997; Butz et al., 1998). Antibodies against Caskin 1 were raised in rabbits against a peptide corresponding to the last 15 residues of Caskin 1 with additional cysteine at the N terminus (sequence: CSFMDDLADQDLAMDE). CASK and Mint monoclonal antibodies were obtained from Transduction Laboratories (Cincinnati, OH).

**RNA blotting experiments.** RNA blotting experiments were performed using multiple tissue blots purchased from Clontech (Cambridge, UK). Northern blots were hybridized using random primer-labeled cDNA probes. Distinct bands (Hata et al., 1996) with a DNA fragment encoding residues 59–232 of Caskin 1 were observed.

**Miscellaneous procedures.** SDS–PAGE was performed as described previously (Laemmli, 1970). Protein concentrations were determined by commercial protein assay kits obtained from Bio-Rad (Hercules, CA) and Pierce (Rockford, IL) and by comparison of samples run on SDS–PAGE with known amounts of BSA standards analyzed on the same gels. Immunocytochemistry experiments with cultured hippocampal neurons and rat brain sections were performed as described previously (Rosahl et al., 1995; Hsueh et al., 1998).

**RESULTS**

**Identification and cloning of Caskin 1**

Affinity purification of rat brain proteins with a GST fusion protein of the N-terminal CaM kinase domain of CASK revealed that in addition to Mint 1 (~170 kDa), a second, slightly larger protein (~180 kDa) was captured in relatively large amounts (data not shown; Butz et al., 1998). To identify the 180 kDa protein, we isolated preparative amounts of binding proteins by affinity chromatography on immobilized CASK and obtained sequences of tryptic peptides by Edman degradation. Ten unequivocal peptide sequences were obtained from the 180 kDa protein (1, ATPSPVPS; 2, SGEYLLDEGAPPGTP; 3, VLPSSGVSHTF?QK; 4, KTPPO?L; 5, DTTDP?GTSP; 6, AVSPPVPGKP; 7, EASAALQR; 8, AGDITVLEQHP; 9, HGTPPPVSGBP; 10, AASD?EP). Databank searches
did not uncover significant homologies of the peptide sequences with known proteins, suggesting that the 180 kDa protein has not been studied previously. However, we did identify several EST clones and two human KIAA clones (KIAA1306 and KIAA1139; GenBank accession #’s AB037727 and AB032965, respectively) that encode the 180 kDa protein or a closely related homolog. We designed PCR primers based on the human nucleotide and rat peptide sequences and obtained a PCR product with rat brain cDNA that was subsequently used to clone the respective cDNAs from rat and mouse brain cDNA libraries using standard techniques (Okamoto and Südhof, 1997). Because of the tight binding of the 180 kDa protein to CASK, we named it Caskin 1 for CASK interacting protein 1 and called its closely related homolog Caskin 2.

Figure 1 displays alignments of the rat and human Caskin 1 and the mouse and human Caskin 2 sequences. Each Caskin isoform is highly conserved evolutionarily, and the two Caskins are very similar. Databank searches indicated that Caskins are composed of two principal regions: the N-terminal half exhibits a multidomain architecture composed of six ankyrin repeats (Fig. 1, purple), a single SH3 domain (Fig. 1, green), and two SAM domains (Fig. 1, blue). The C-terminal half is made up of a long proline-rich sequence (prolines highlighted in red in Fig. 1 to illustrate their abundance) and a unique conserved C-terminal domain (CTD) (Fig. 1, yellow). The N-terminal halves of Caskins 1 and 2 are very similar, especially in the identified domains. In contrast, the C-terminal regions diverge considerably but exhibit patches of identity that are often organized around proline residues. For example, an unusual sequence (KGPPPPPPPKRxSS) that could form a polyproline helix and a protein kinase A phosphorylation site is conserved in all sequences, as is a second protein kinase A consensus site (RRRTLxSxP) and another cluster of positively charged amino acids (LTESDTVXRKPxKxE).

Figure 2 summarizes the domain organization of Caskins. Although the N-terminal ankyrin repeats and the SH3 and SAM domains were identified by standard domain searches, such as
that of the conserved domain database of the National Center for Biotechnology Information, these domains are rather atypical. The SH3 domain in particular is only distantly related to other SH3 domains (data not shown); its closest homolog is the SH3 domain of yeast CDC25. The domain organization of Caskins is different from that of other multidomain proteins in the current databases. The structure of Caskins resembles that of Shanks, a family of postsynaptic scaffolding proteins that bind to guanylate kinase-associated protein, glutamate receptor-interacting protein homer, and cortactin (for review, see Sheng and Sala, 2001). Shanks and Caskins both contain N-terminal ankyrin repeats followed by an SH3 domain and a large C-terminal proline-rich sequence. However, in Shanks, the SH3 domain is followed by a PDZ domain that is absent from Caskins. Furthermore, in Shanks, a single SAM domain is positioned at the end of the C terminus, whereas Caskins contain two SAM domains in the center (Fig. 2). Finally, the ankyrin repeats and the SH3 and SAM domains of Caskins and Shanks exhibit little sequence identity and are more related to analogous domains in other proteins than to each other. Thus, it appears that the similar domain organization of Caskins and Shanks does not reflect an evolutionary connection but rather possibly a similar scaffolding function. Databank searches failed to uncover a direct homolog of Caskins in Drosophila or C. elegans, although multiple proteins containing N-terminal ankyrin repeats and SAM domains were observed. Because the nearly complete Drosophila and C. elegans genome sequences are available, this suggests that Caskins, unlike CASK and Mint 1, are not conserved in invertebrates.

**Tissue distribution of Caskin 1**

To examine which tissues express Caskin 1, we hybridized a multitissue RNA blot at high stringency with a Caskin 1 cDNA probe. The resulting signal revealed that among the tissues examined, Caskin 1 mRNA was detectable only in the brain (Fig. 3). We subsequently raised an antibody against a synthetic peptide corresponding to the C terminus of Caskin 1. Immunoblotting of rat brain homogenates with the antibody uncovered a single major band of ~180 kDa that was not observed with preimmune serum (Fig. 4 and data not shown). Comparison of this band with the band detected in COS cells transfected with a full-length Caskin 1 expression vector showed that the recombinant Caskin 1 comigrated with the band observed in brain homogenates (Fig. 4). This result suggests that the band observed in the brain corresponds to Caskin 1 and that the cDNA we isolated for Caskin 1 is full-length with respect to the coding region. Immunoblots of different rat tissues with the Caskin 1 antibody confirmed the conclusion from RNA blotting that among all tissues
examined, Caskin 1 is expressed at detectable levels only in the brain (Fig. 5).

We used immunocytochemistry experiments to determine the distribution of Caskin 1 in the brain (Fig. 6). Immunoperoxidase labeling of rat brain sections revealed that Caskin 1 was localized primarily to the neuropil and was enriched in synaptic areas. This pattern resembles that of synaptic vesicle proteins, such as rab3A, and is illustrated in Figure 6A,B for the cerebellum. The observed labeling was specific, as evidenced by the lack of staining obtained with preimmune serum (Fig. 6C). To ensure that Caskin 1 is indeed a neuronal protein, we stained cultured hippocampal neurons (Fig. 6D,E). Caskin 1 was highly concentrated in these neurons and was present throughout the cells. Although Caskin was also observed in synapses, we observed no apparent enrichment in synapses in hippocampal neurons (Fig. 6E and data not shown). In this regard, Caskin 1 was very similar to proteins such as CASK and syntaxin 1 that were also enriched in synapses in brain sections, but it appeared to be more widely distributed throughout the cells in cultured neurons (data not shown; also see Hsueh et al., 1998).

**Immunoprecipitation of a CASK–Caskin 1 complex from brain homogenates**

To obtain independent evidence that Caskin 1 and CASK form a complex in the brain, we performed immunoprecipitations. Proteins from rat brain homogenates were immunoprecipitated with preimmune or anti-Caskin 1 serum and analyzed by immunoblotting. CASK was specifically coimmunoprecipitated with Caskin 1 (Fig. 7A). Mint 1 and Munc18 were not detected in the immunoprecipitates, suggesting that Caskin 1 antibodies do not bring down Mint 1 together with CASK. We subsequently washed the immunoprecipitates with solutions of increasing ionic strength (0.2–1.0 M NaCl), a chaotropic agent (0.6 M KI), or a denaturing...
solution (0.5% SDS in 2% Triton X-100). Figure 7B shows that the CASK–Caskin complex resisted high-salt washes of up to 1 M NaCl and was disrupted only by chaotropic or denaturing agents, suggesting that it is very stable.

Caskin 1 and Mint 1 bind to the same site on CASK
We subsequently used GST pulldowns to examine the interaction of Caskin 1 with CASK. Initial experiments showed that GST–CASK efficiently captured Caskin 1 and Mint 1 from the brain, as expected (Fig. 8). Interestingly, although both GST–Mint 1 and GST–Caskin 1 bound CASK, Mint 1 did not pull down Caskin 1 with CASK, and Caskin 1 did not pull down Mint 1. Because Mint 1 also was not coimmunoprecipitated with CASK by the Caskin antibodies (Fig. 7), this result suggests the possibility that Caskin 1 and Mint 1 bind to the same site on CASK. To test this hypothesis, we expressed a Caskin 1 fusion protein with MBP in bacteria. We subsequently added increasing amounts of purified recombinant MBP–Caskin during GST–CASK pulldown experiments from brain homogenates (Fig. 9). The results show that higher Caskin 1 concentrations prevent Mint 1 binding to CASK, suggesting that CASK cannot bind Mint 1 and Caskin 1 at the same time.

The N-terminal regions of CASK and Caskin 1 bind to each other
In the next set of experiments, we examined which CASK and Caskin 1 sequences interact with each other. Binding of rat brain CASK to a series of increasingly smaller GST–CASK fusion proteins uncovered a short sequence in Caskin 1 (residues 375–471) that was fully capable of affinity-purifying CASK from rat brain homogenates (Fig. 10). As expected, Mint 1 was not copurified with CASK, but Velis (which bind to a different sequence in CASK) (Butz et al., 1998) were pulled down together with CASK. This indicates that CASK forms a tripartite complex with Caskin 1 and Velis similar to the CASK–Mint 1–Veli complex (Butz et al., 1998). The Caskin 1 sequence that binds to CASK lies between the SH3 domain and the two SAM domains and is relatively poorly conserved between Caskins 1 and 2 (Fig. 1).
Mint 1, and Velis as indicated. Signals were visualized by ECL.

Caskin fusion proteins were analyzed by immunoblotting for CASK, left at recombinant MBP MBP alone at 500/\textmu g (lane 8). Bound proteins were visualized by Coomassie blue staining (top) or immunoblotting with Mint 1 antibodies (bottom). Numbers at left indicate positions of molecular weight markers.

**Figure 9.** Caskin 1 competes with Mint 1 for CASK binding. Proteins from rat brain homogenates (Homog.) (lane 1) were bound to immobilized GST–CASK\(^{1-335}\) in the presence of increasing concentrations of recombinant MBP–Caskin 1 fusion protein up to 500 \(\mu\)g (lanes 2–7) or of MBP alone at 500 \(\mu\)g (lane 8). Bound proteins were visualized by Coomassie blue staining (top) or immunoblotting with Mint 1 antibodies (bottom). Numbers at left indicate positions of molecular weight markers.

suggesting that only Caskin 1, and not Caskin 2, binds to CASK, which is similar to the interaction of only Mint 1, and not Mints 2 and 3, with CASK (Butz et al., 1998). This hypothesis was confirmed in experiments that showed that GST–Caskin 2 fusion proteins cannot capture CASK from rat brain homogenates (data not shown).

We subsequently studied which CASK sequences bind to Caskin 1 and Mint 1 (Fig. 11). Again, we used a series of GST fusion proteins to elucidate the sequence requirement for binding (Fig. 11A). The first set of experiments showed that like Mint 1, Caskin 1 was efficiently bound by the N-terminal CaM kinase domain of CASK, consistent with competition between the two proteins for binding to CASK (Fig. 11B). Velis, in contrast, were bound by the C-terminal fragment. We subsequently used deletion analysis to determine which sequences are precisely required for Caskin 1 and Mint 1 binding. Figure 11C demonstrates that a C-terminal truncation of CASK, in which the autoinhibitory region and calmodulin-binding site of the CaM kinase homology region of CASK are included (Cask\(^{1-319}\)), still binds to both Caskin 1 and Mint 1. In contrast, removal of the autoinhibitory region and calmodulin-binding site of CASK (Cask\(^{1-275}\)) abolishes Caskin 1 and Mint 1 binding. Thus, for both Caskin 1 and Mint 1, the sequence of CASK that is C terminal to the actual kinase domain and corresponds to the regulatory sequences in CaM kinase II (including its autophosphorylation site threonine\(^{286}\)) (Hata et al., 1996) is essential for binding. N-terminal truncation of the CASK CaM kinase domain demonstrated that Caskin 1 and Mint 1 not only bind to the C-terminal peptide sequence of the domain but also require a larger region. Deletion of the N-terminal 98 residues of CASK strongly diminished Caskin 1 and Mint 1 binding, whereas removal of an additional 139 residues abolished binding (Fig. 11C). Together, these results suggest that the entire CaM kinase domain of CASK is involved in binding relatively short sequences of Caskin 1 and Mint 1. This binding reaction is specific for the catalytically inactive CASK CaM kinase domain, because genuine CaM kinase II was unable to bind to either Caskin 1 or Mint 1 (data not shown).

**Figure 10.** Mapping of the CASK-binding site on Caskin 1 by GST pulldowns. A. Position of GST–Caskin 1 fusion proteins used for pulldowns in B. B. GST pulldowns of rat brain proteins with the indicated Caskin fusion proteins were analyzed by immunoblotting for CASK, Mint 1, and Velis as indicated. Signals were visualized by ECL. Numbers at left indicate positions of molecular weight markers.

.Assembly of a CASK–Caskin 1 complex on the immobilized cytoplasmic tails of neurexins

CASK binds to the cytoplasmic tails of several cell-surface proteins, including neurexins, syndecans, and JAMs (Hata et al., 1996; Cohen et al., 1998; Hsu et al., 1998; Biederer and Südhof, unpublished observations). This reaction can be biochemically reconstituted with the immobilized recombinant tail of neurexin 1, which recruits not only CASK but also Mint 1 and Velis from brain homogenates (Biederer and Südhof, 2000). To test whether Caskin 1 can also be coassembled with CASK on the cytoplasmic tails of neurexins, we used affinity chromatography (Fig. 12). Both Caskin 1 and Mint 1 were efficiently bound together with CASK, indicating that the CASK–Caskin 1 complex could be assembled on the plasma membrane. In contrast, GDP dissociation inhibitor (GDI, used as a negative control) remained unbound.

**DISCUSSION**

In the present study, we purified a novel CASK-binding protein that we named Caskin 1, determined its primary structure from cDNA clones, and studied its interactions with CASK. A closely related second isoform of Caskin 1, called Caskin 2, was also identified and cloned. Our data reveal that Caskin 1 constitutes a multidomain protein that in adult rodents is detected only in brain, in which it binds tightly to CASK. Caskin 1 and 2 are novel proteins that are distinct from other previously reported multidomain proteins, with a structure that suggests an adaptar function. At the N terminus, Caskins contain six ankyrin repeats that in many proteins participate in protein–protein interactions, fol-
followed by an atypical SH3 domain and two SAM repeats (Fig. 2). After a long proline-rich region, the two Caskins end in a similar C-terminal sequence (Fig. 1). This sequence may correspond to a novel evolutionarily conserved domain, because a similar sequence is found in an otherwise unrelated Drosophila protein in GenBank (accession #AAF58176; data not shown). Caskins resemble many other adaptor proteins that usually contain a series of small, autonomously folded binding modules (such as the ankyrin repeats, SAM domains, and SH3 domain in Caskins) and in addition often include extended proline-rich sequences with characteristic signature motifs that may recruit the SH3 or WW domains of other proteins. The domain composition of Caskins is most similar to that of the Shank family of postsynaptic adaptor proteins (for review, see Sheng and Sala, 2001), although the precise number, type, and arrangement of domains differ (Fig. 2).

It is generally noticeable that multidomain adaptor proteins appear to be particularly important for synaptic transmission and nuclear transcription, presumably because these reactions are most tightly regulated.

To define the protein network in which Caskin 1 participates, we mapped its interactions with CASK and compared them with those of Mint 1, a previously identified interaction partner of CASK (Butz et al., 1998). We showed that Caskin 1 binds to the N-terminal CaM kinase domain of CASK in direct competition with Mint 1. As a result, CASK participates in two independent and alternative tripartite complexes: the previously characterized CASK–Velis–Mint 1 complex (Borg et al., 1998a; Butz et al., 1998) and the new CASK–Velis–Caskin 1 complex. The coimmunoprecipitation of Caskin 1 and Mint 1 with CASK (Fig. 7) and the coassembly of Caskin 1 and Mint 1 with CASK on the immobilized cytoplasmic tail of neurexin 1 (Fig. 12) suggest that both complexes are normally present in the brain and bind to the cytoplasmic tails of neurexins and other cell-surface proteins. Viewed together with previous studies (Hata et al., 1996; Borg et al., 1998b; Butz et al., 1998; Cohen et al., 1998; Hsueh et al., 1998; Biederer and Südhof, 2001), these results show that in the vertebrate brain, CASK forms tight complexes with a multitude of distinct proteins (Fig. 13): The N-terminal CaM kinase domain of CASK binds to either Caskin 1 or Mint 1 and interacts with calmodulin; the central Veli-interacting domain binds to Velis; the following PDZ domain interacts with the C-terminal cytoplasmic sequences of cell-surface proteins such as neurexins and syndecans, which anchor the entire complex on the plasma membrane; and finally, the C-terminal region of CASK binds to protein 4.1, which in turn connects to the actin cytoskeleton.

It is striking that except for the cell-surface proteins and calmodulin, all proteins known to bind to CASK are themselves adaptor proteins composed of multiple domains. The two alternative tripartite complexes formed by CASK (CASK–Mint 1–Velis and CASK–Caskin 1–Velis) incorporate a large number of potential interaction domains: multiple PDZ, SAM, and SH3 domains, in addition to ankyrin-like repeats, a guanylate kinase region, and other unique motifs, suggesting that these complexes form the nucleus of multiple high-molecular-weight complexes.

Figure 11. Mapping of the Caskin 1 binding site on CASK. A, Domain structure of CASK and positions of GST–CASK fusion proteins used for pulldowns. B, C, GST pulldowns of proteins in rat brain homogenates with the indicated CASK fusion proteins analyzed by immunoblotting for the proteins identified at right. B, Proteins were eluted with 0.8 M K-acetate (KAc) followed by SDS sample buffer, whereas in C, proteins bound to the beads were examined. Note that in CASK, not only the CaM kinase domain but also the region homologous to the autoregulatory sequence of CaM kinase II are required for binding. Also note the separation between the common Mint 1–Caskin 1 and the Veli binding sites on CASK. Signals were visualized by ECL. Numbers at left indicate positions of molecular weight markers.
The fact that Mint 1 and Caskin 1 are expressed primarily in the brain indicates that the two tripartite complexes are assembled only in the brain. In contrast, CASK and all other interacting proteins are widely expressed outside of the brain, where they presumably participate in other complexes with as yet unidentified interactors for the CASK CaM kinase domain that are ubiquitously expressed. Finally, it is noticeable that Caskin 1 is vertebrate-specific, whereas CASK and all of its other interacting proteins are evolutionarily conserved in invertebrates. The evolutionary derivation of novel proteins by mixing and matching established, phylogenetically old domains represents a quick mechanism of inventing new proteins that contribute to the rapid emergence of molecular complexity during mammalian evolution. This, together with the apparent lack of Caskin conservation in invertebrates, suggests that CASK diversified evolutionarily toward the use of multiple effectors that include Caskin 1. Although much remains to be done to elucidate the precise functions of CASK and its alternative interactions with Caskin 1 and Mint 1, the description of the protein networks in which these proteins participate clearly represents the first necessary step toward addressing this question.

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