The src Protein Contains Multiple Domains for Specific Attachment to Membranes

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The proteins encoded by the oncogene v-src and its cellular counterpart c-src (designated generically here as pp60"src") are tightly associated with both plasma membranes and intracellular membranes. This association is due in part to the amino-terminal myristylation of pp60"src", but several lines of evidence suggest that amino-terminal portions of the protein itself are also involved. We now report that pp60"src" contains at least three domains which, in conjunction with myristylation, are capable of mediating attachment to membranes and determining subcellular localization. We identified these domains by fusing various portions of pp60"src" to pyruvate kinase, which is normally a cytoplasmic protein. Amino acids 1 to 14 of pp60"src" are sufficient to mediate both myristylation and the attachment of pyruvate kinase to cytoplasmic granules. In contrast, amino acids 38 to 111 mediate association with the plasma membrane and perinuclear membranes, whereas amino acids 204 to 259 mediate association primarily with perinuclear membranes. We conclude that pp60"src" contains independent domains that target the protein to distinctive subcellular locations and thus may facilitate diverse biological functions of the protein.

The products of the retroviral oncogene v-src and its cellular counterpart c-src (designated generically here as pp60"src") are protein tyrosine kinases that are associated with membranes. Association of pp60"src" with membranes is essential for transformation by v-src (19). In various cell types, the viral and cellular src proteins are associated with plasma membranes (7, 8, 21, 44), with perinuclear membranes (36), with secretory organelles in both chromaffin cells and platelets (30, 33), and with growth cones in developing neurons (27, 41). Little is known about how pp60"src" is specifically targeted to these subcellular locations.

The amino terminus of pp60"src" is covalently coupled to a 14-carbon fatty acid, myristic acid (4, 39). The myristyl moiety promotes the association of pp60"src" with membranes (2, 11). On the other hand, not all myristylated src proteins are associated with membranes (3, 11), and some nonmyristylated src proteins are membrane associated, albeit weakly (12, 20). These results suggest that pp60"src" contains amino acid sequences which, in conjunction with amino-terminal myristylation, mediate the association of the protein with membranes.

Previous work has implicated the amino-terminal 10 kilodaltons (kDa) of pp60"src" in attachment to membranes (22, 35). Here we report work that dissect the amino-terminal domain into several regions that independently target and attach the protein to membranes in distinctive subcellular locations.

MATERIALS AND METHODS

Materials. The following reagents were used: monoclonal anti-pp60"src" antibody 327 (MAb327), from J. Brugge (25); rabbit antibodies specific for the chicken M1 pyruvate kinase (PK), from B. Roberts; simian virus 40 expression vectors containing the chicken M1 PK cDNA (RL142PK10X, RL18PK8X, and RL18PK12X, which differ by the reading frame at a polylinker sequence), from B. Roberts (16); and a plasmid containing the chicken fibroblast c-src cDNA, pSH, from H. Hanafusa. The E3 src gene is a deletion mutant which lacks the sequence encoding amino acids 8 to 37 of pp60"src", as previously described (17). Reagents were obtained as follows: protease inhibitors, paraformaldehyde, bovine serum albumin (fraction V), and detergents from Sigma Chemical Co.; fluorescein-conjugated phalloidin from Molecular Probes Inc.; 1-[35S]methionine from ICN Radiochemicals; [3H]myristic acid, 22.4 Ci/mmole (50 mCi/ml in dimethyl sulfoxide) from Du Pont, NEN Research Products; restriction enzymes, Escherichia coli DNA polymerase I, and T4 DNA ligase from New England Biolabs, Inc.; human serum fibronectin from Collaborative Research Inc.; and Gold Seal glass cover slips (18 by 18 mm; thickness, 1.5 mm) from VWR Scientific.

Metabolic labeling, immunoprecipitation, and subcellular fractionation. COS7 cells were labeled with [35S]methionine for 18 h in Dulbecco modified Eagle medium containing 10% the normal concentration of L-methionine, 10% dialyzed fetal calf serum, and 200 μCi of L-[35S]methionine per ml. COS7 cells were labeled with [3H]myristic acid for 18 h in Dulbecco modified Eagle medium with 10% fetal calf serum, 1% dimethyl sulfoxide, and 500 μCi of [3H]myristic acid per ml (5). Labeled cell extracts were prepared with lysis buffer (20 mM Tris hydrochloride [pH 8.0], 150 mM NaCl, 1% Nonidet P-40, 50 μg of soybean trypsin inhibitor per ml, 50 μg of aprotenin per ml, 20 μg of leupeptin per ml, and 1 mg of fraction V bovine serum albumin per ml). Immunoprecipitations with polyclonal rabbit sera and monoclonal antibodies were performed as described previously (21). Crude cell extracts prepared by Dounce homogenization of hypotonically swelled cells were fractionated into cytosol and membrane fractions by differential centrifugation, as previously described (21). Subcellular fractions

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were constructed by cutting the src gene with restriction endonucleases and fusing the repaired ends. The restriction sites used to create internal deletions of src genes are indicated. Residue numbers correspond to the sequence of pp60^c-src encoded by the Prague C strain of Rous sarcoma virus (39). Myristylation (Mys) of hybrid proteins, when analyzed, is indicated by +. When src-PK proteins associated with membranes, myristylation was assumed [and denoted (+)] but not tested. Membrane association (MA) of hybrid proteins, as determined by biochemical fractionations, is indicated by +. The intracellular distribution of hybrid proteins was determined by immunofluorescence (IF). Association of hybrid proteins with plasma membranes (PM), with the cytoplasm (CYT), with perinuclear membranes (PN), or with cytoplasmic granules (G) is indicated. At the bottom of the figure the N-terminal 259 amino acids of pp60^c-src are divided into functional domains. The first 7 amino acids of pp60^c-src act as a recognition sequence for myristylation. Amino acids 1 to 14 cause association with cytoplasmic granules (G). Amino acids 38 to 111 ( ), together with the myristylation signal, cause association with both the plasma membrane and perinuclear membranes (PM+PN). Amino acids 204 to 259 ( ), together with the first 14 amino acids, cause association solely with perinuclear membranes (PN). The myristyl moiety is indicated by the crooked line in the diagram of src259PK (top of figure).

Construction of recombinant src-PK genes. The N-terminal amino acid sequence of the hybrid src-PK proteins is derived from pp60^c-src (B77 strain of Rous sarcoma virus), whereas the C-terminal amino acids are derived from PK. The src-PK proteins are designated by the C-terminal boundary of the domain of pp60^c-src that is fused to PK, and in cases in which this domain contains internal deletions, the amino acids that are lost are indicated in parentheses. Residue numbers reported correspond to the sequence of the Prague C strain of Rous sarcoma virus (40). The amino acids derived from the chicken M1 PK (residues 120 to 529) are the same in all cases (26). The nucleotide sequence encoding PK was derived from plasmid RL142PK10X, RL18PK8X, or RL18PK12X as required to maintain an open reading frame. The src7PK plasmid (previously known as R7-PK) was described previously (17). Sequences encoding various N-terminal amino acids of pp60^c-src were fused to the coding region of PK by cutting both genes with restriction endonucleases, repairing the ends with E. coli DNA polymerase I, and fusing the resulting blunt ends. Internal deletions of src sequences were constructed by cutting the src gene with restriction endonucleases and fusing the repaired ends. The restriction sites used in these constructions are indicated in Fig. 1. The Stul site at codon 7 is a restriction polymorphism which was created by converting lysine 7 to arginine (17). A simian virus 40 vector expressing the chicken fibroblast pp60^c-src pSV5H, was constructed by replacing the PK cDNA of RL142PK10X with the c-src cDNA derived from p5H. Further details about the construction of these genes will be provided upon request.

Transfection of recombinant src-PK genes. COS7 cells were exposed for 18 h to calcium phosphate precipitates containing 20 μg of recombinant src-PK DNA and then subjected to a 2-min shock with Tris-buffered saline (0.8% NaCl, 0.1% glucose, 0.038% KCl, 0.2% Tris hydrochloride, 0.06% Tris base, 4.5 mg of phenol red per liter) containing 25% dimethyl sulfoxide (15). Transfected COS7 cells were labeled for 18 h with either 1-[^35]S]methionine or [3H]myristic acid starting 36 h after the transfection had begun.

Immunofluorescence. Glass cover slips were coated with human serum fibronectin (100 μg/ml) in Dulbecco phosphate-buffered saline (PBS) for 1 h at 37°C in a humidified chamber. Transfected COS7 cells were seeded onto glass cover slips 24 h after the transfection had begun. After 8 to 12 h, the cover slips were washed with PBS, cells were fixed with 3.5% paraformaldehyde in PBS for 20 minutes and

FIG. 1. Summary of the membrane association of src-PK proteins. The sequences derived from pp60^c-src in all of the src-PK proteins are depicted. The restriction sites used to create internal deletions of src genes and to fuse the src sequences to the coding region of PK are indicated. The intracellular distribution of hybrid proteins was determined by immunofluorescence (IF). Association of hybrid proteins with plasma membranes (PM), with the cytoplasm (CYT), with perinuclear membranes (PN), or with cytoplasmic granules (G) is indicated. At the bottom of the figure the N-terminal 259 amino acids of pp60^c-src are divided into functional domains. The first 7 amino acids of pp60^c-src act as a recognition sequence for myristylation. Amino acids 1 to 14 cause association with cytoplasmic granules (G). Amino acids 38 to 111 ( ), together with the myristylation signal, cause association with both the plasma membrane and perinuclear membranes (PM+PN). Amino acids 204 to 259 ( ), together with the first 14 amino acids, cause association solely with perinuclear membranes (PN). The myristyl moiety is indicated by the crooked line in the diagram of src259PK (top of figure).
permeabilized with 0.1% Triton X-100 in PBS, and cover slips were coated with 10% normal goat serum in PBS to block nonspecific binding sites. Primary antibodies used were monoclonal anti-p60src 327 (10 μg/ml in PBS) and polyclonal rabbit anti-PK (1/200 dilution in PBS). Fluorescein-conjugated antibodies were used at a dilution of 1/200 in PBS. Rhodamine-conjugated phalloidin was used as recommended by the manufacturer. Stained cover slips were washed several times with PBS and then with absolute ethanol and were embedded in glycerol containing 2% propyl gallate, an antibleaching reagent. Microscopy was done with a Zeiss Photomicroscope III or an inverted Olympus fluorescence microscope. Cells were photographed with Kodak Tr-X film at ASA 200. In general, 1 to 5% of the transfected COS7 cells gave a fluorescent signal following this procedure; this signal presumably corresponds to the fraction of cells that took up the DNA. The surrounding nonsfluorescent cells acted as an internal control for the specificity of the antibodies used.

RESULTS

To test the role of N-terminal amino acid sequences of p60src in association with membranes and in subcellular localization, we constructed a set of genes encoding hybrid src-PK proteins (Fig. 1). Since myristylation is normally required for membrane association, we included the myristylation signal in all of these hybrid genes. The association of the hybrid proteins with membranes was analyzed first by a crude biochemical fractionation and subsequently by immunofluorescence.

Myristylation is not sufficient for membrane anchorage. We have previously shown that the first 7 amino acids of p60src act as a sufficient recognition sequence for myristylation (17). Fusing sequences encoding the first 7 amino acids of p60src to the coding region of PK creates a hybrid gene, src7PK, which encodes a myristylated protein (see Fig. 6). If myristylation is a sufficient cause of membrane association, the src7PK protein should attach to membranes.

Cells expressing either the native PK protein or the src7PK protein were fractionated into soluble and membrane fractions, and these fractions were assayed for PK proteins (Fig. 2). Virtually all of the native PK protein (Fig. 2, lanes 1 to 3) and the src7PK protein (Fig. 2, lanes 4 to 6) was recovered in the soluble fraction (Fig. 2, lane 2 and 5, respectively), indicating that a second domain of p60src is required in conjunction with myristylation to cause membrane association.

The gels illustrated in Fig. 2 contained variable amounts of proteins that appeared ancillary to the specific products of the transfected DNAs (marked by arrowheads). We do not know the identities of these ancillary proteins, but they were present in all specimens, including cells that had not been transfected. We therefore attribute them to nonspecific immunoprecipitation.

Anchoring PK to membranes with a domain from p60src. Since previous reports had implicated N-terminal sequences in p60src as membrane-anchoring domains (9, 12, 19, 22, 34), we fused sequences encoding the first 259 amino acids of the protein to the coding sequences of PK, creating a hybrid gene called src259PK. Cells expressing the src259PK protein were fractionated into cytosol and membrane fractions, and these fractions were assayed for the src259PK protein (Fig. 2, lanes 7 to 9). Most (ca. 90%) of the src259PK protein was recovered in the membrane fraction (lane 9). This result demonstrates that sequences in the N-terminal half of p60src, together with myristylation, are sufficient to mediate association with membranes.

The src259PK protein and p60src associate with membranes in a similar manner. Before further characterizing the membrane-anchoring domains made up of the N-terminal
259 amino acids of pp60\textsuperscript{src}, we tested whether the src259PK and pp60\textsuperscript{src} proteins associate with membranes in a similar manner. We first asked whether myristylation is required for the association of src259PK with membranes. Replacement of lysine 7 with an asparagine largely abolishes myristylation of pp60\textsuperscript{src} (17). Therefore, we converted lysine 7 to asparagine in src259PK and evaluated the subcellular distribution of the mutant protein (Fig. 3, lanes 4 to 6). The protein (designated N7-259PK) was found predominantly (ca. 87%) in the soluble fraction of cells (Fig. 3, lane 5), implying that the membrane association of src259PK, like that of pp60\textsuperscript{src} itself, requires N-terminal myristylation.

A second hallmark of the association of pp60\textsuperscript{src} with membranes is that, like other peripheral membrane proteins, pp60\textsuperscript{src} can be removed from membranes by alkaline extractions (Fig. 4, lanes 4 and 5). Similarly, the src259PK protein is extracted from membranes following extraction with base (Fig. 4, lanes 9 and 10). On the other hand, neither pp60\textsuperscript{src} (Fig. 4, lanes 2 and 3) nor src259PK (Fig. 4, lanes 6 to 8) is efficiently extracted from membranes by 0.3 M NaCl. We conclude that by several criteria, src259PK and pp60\textsuperscript{src} appear to associate with membranes in a similar manner.

The first 111 amino acids of the src protein are sufficient for membrane association. Previous work indicated that the N-terminal 10 kDa of pp60\textsuperscript{src} is required to anchor the protein to membranes (22, 34). To better define the responsible amino acid sequences, we fused the coding region of PK to sequences encoding each of the following domains: the first 204 amino acids of pp60\textsuperscript{src} (src204PK); the first 111 amino acids of pp60\textsuperscript{src} (src111PK); and the first 204 amino acids of pp60\textsuperscript{src} from which residues 8 to 111 were deleted [src204(AS-111)PK]. Cells expressing these proteins were fractionated as described above (Fig. 2).

Although 75% of the src204(AS-111)PK protein was recovered in the cytosol (Fig. 2, lanes 13 to 15), most of the src204PK protein (not shown in Fig. 2, but see Fig. 4, lanes 11 to 15) and ca. 65% of the src111PK protein (Fig. 2, lanes 10 to 12) were recovered with the membrane pellet (Fig. 2, lanes 10 to 12). The proteins encoded by both src204PK (Fig. 4, lanes 11 to 15) and src111PK (Fig. 4, lanes 16 to 20) associated with membranes in a manner similar to pp60\textsuperscript{src}.
because they were extracted only partially from membranes by 0.3 M NaCl, but were extracted quantitatively by base.

We conclude that the first 111 amino acids of pp60src are sufficient to cause association with membranes and that the association is similar to that found with native pp60src. By contrast, amino acids 111 to 204, together with N-terminal myristylation, permit only limited association with membranes.

The first 111 amino acids of the src protein contain at least two independent membrane-anchoring domains. The domain represented by the first 111 amino acids of pp60src was divided into several segments, and the sequences encoding these segments were fused to the coding region of PK. The association of these hybrid proteins with membranes was analyzed by biochemical fractionation as described above.

(i) The first 14 amino acids of the src protein contain a membrane-anchoring domain. It was previously shown that fusing the first 14 amino acids of pp60src to any one of several other proteins causes these proteins to associate with membranes in crude biochemical fractions (1, 31, 32). This result was confirmed by fusing sequences encoding the first 14 amino acids of pp60src to the PK-encoding region, creating src14PK. An appreciable fraction of the src14PK protein was recovered with the membrane pellet after subcellular fractionation (Fig. 5, lanes 4 to 6), although precise quantitation was prohibited by the presence of a nearly comigrating protein.

Other hybrid proteins that contained the first 14 amino acids of pp60src also associated with membranes: 72% of src14PK (Fig. 5, lanes 7 to 9), 72% of src111(Δ15-54)PK (Fig. 5, lanes 13 to 15), and 67% of src259(Δ14-203)PK (Fig. 5, lanes 16 to 18) were recovered with the membrane pellet. These results leave open the possibility that amino acids 15 to 54, 55 to 111, and 204 to 259 also contain membrane-anchoring domains, a possibility that will be confirmed below by immunofluorescence (see Fig. 7).

(ii) Amino acids 38 to 111 of the src protein contain a membrane-anchoring domain. Two previous observations indicated that portions of pp60src outside the first 14 amino acids contribute to membrane attachment. First, removal of more than 8 kDa from the amino terminus of pp60src failed to detach the protein from membranes (22). Second, the protein encoded by the deletion mutant E3 src, which lacks amino acids 8 to 37, attaches to membranes (17).

We sought and identified a second membrane-anchoring domain by fusing the sequence encoding the first 81 amino acids of the E3 src protein to the coding region of PK, creating the src111(Δ8-37)PK gene. Most (66%) of the src111(Δ8-37)PK protein (Fig. 5, lanes 10 to 12) was recovered with the membrane pellet (Fig. 5, lane 12) following subcellular fractionation. This implies that amino acids 38 to 111 of pp60src also contain a membrane-anchoring domain.

Failure of chimeric proteins to bind membranes cannot be attributed to poor myristylation. An alternative explanation for the behavior of the cytoplasmic src7PK protein and the membrane-associated src259PK and src14PK proteins is that they differ not by the presence of a membrane-anchoring domain, but rather in the stoichiometry of myristylation. If the association of pp60src with membranes were mediated solely by the myristyl moiety, only the myristylated molecules would associate with membranes. This possibility was tested by comparing the stoichiometry of myristylation of the src7PK (Fig. 6, lanes 2 and 5), the src14PK (Fig. 6, lanes 3 and 6), and the src259PK (Fig. 6, lanes 1 and 4) proteins. Cells expressing these proteins were labeled with either L-[3H]myristate (lanes 1 to 3) or [3H]myristic acid (lanes 4 to 6), and the PK proteins were analyzed by immunoprecipitation.

Since it had previously been shown that 84% of pp60src molecules are myristylated (4), we arbitrarily used the same value for the stoichiometry of myristylation of src259PK. In comparison, approximately 60% of the src14PK protein and 17% of the src7PK protein were myristylated (estimates derived from Fig. 6). Thus, if myristylation were sufficient to cause membrane association, 17% of the src7PK protein
Virtually all of the myristate-labeled proteins were labeled with [3H]myristic acid and one hand, and the membrane-associated cytoplasmic molecules that are myristylated, all of the myristylated molecules would be membrane associated. Cells expressing these proteins were poorly visualized after staining with anti-PK antibody, and the intensity of the staining was proportional to the thickness of the cell (i.e., to the volume of underlying cytoplasm). The src204COS7 protein appeared to be equally distributed between the plasma membrane and the cytoplasm (Fig. 7G). These results confirm our interpretation of the biochemical fractionations described above.

(iii) Most of the membrane-associated src-PK proteins have an intracellular distribution similar to pp60src. The src259PK protein (Fig. 7D), the src204PK protein (Fig. 7E), the src111PK protein (Fig. 7F), the src111COS7PK protein (Fig. 7H), the src54PK protein (Fig. 7I), and the src111COS7PK protein (Fig. 7J) had a distribution similar to that of pp60src itself. Often cells expressing src-PK proteins that associate with the plasma membrane appeared to have a filamentous component to the fluorescent signal (for example, see Fig. 7E). The filamentous structures also contained actin, as demonstrated by staining with rhodamine-conjugated phalloidin, an actin-specific stain (data not shown). We do not know the nature of these filamentous structures, but they conceivably could correspond to sites of adhesion.

(iv) The src259(Δ14-203)PK protein associates specifically with perinuclear membranes. Interestingly, the src259(Δ14-203)PK protein associated primarily with perinuclear membranes (Fig. 7K). Since the distribution of the src259(Δ14-203PK) and the src14PK (see below) proteins were different, we conclude that amino acids 204 to 259 contain a domain that preferentially targets proteins to perinuclear membranes.

(v) The src14PK protein associates with cytoplasmic granules. Cells expressing the src14PK protein had a punctate pattern of cytoplasmic fluorescence after staining with anti-PK antibody (Fig. 7L). These fluorescent spots had an apparent diameter of 0.2 μm and were uniformly distributed throughout the cytoplasm. We suspect that this distribution corresponds to a membranous organelle, rather than a large proteinaceous complex, because the src14PK protein does not sediment rapidly after dissolution of membranes with 1% Triton X-100 (data not shown). The photographic reproduction of Fig. 7L is not adequate to exclude a perinuclear location for some of the src14PK protein. However, other exposures of the same field and examination of numerous other cells sustain our conclusion that src14PK is located principally in cytoplasmic granules, rather than in either of the other two locations reported here.)

The src14PK is the only hybrid protein that associates principally with cytoplasmic granules. Although most of the
FIG. 7. Membrane-anchoring domains target proteins to specific subcellular locations. COS7 cell were transfected with plasmids encoding pp60^csrc (A), PK (B), src7PK (C), src259PK (D), src204PK (E), src111PK (F), src204(Δ8-111)PK (G), src111(Δ8-57)PK (H), src54PK (I), src111(Δ15-54)PK (J), src259(Δ14-203)PK (K), or src14PK (L) proteins. At 48 h after exposure to these DNAs, the cells were seeded onto glass cover slips. After an additional 24 h, the cells were fixed with paraformaldehyde, permeabilized with Triton X-100, and stained with either anti-pp60^csrc MAb327 or a polyclonal anti-PK antibody, as described in Materials and Methods. Perinuclear membranes (PN) and cytoplasmic granules (G) are indicated.
src-PK proteins that contain amino acids 1 to 14 were also found in cytoplasmic granules, proteins that contain additional portions of pp60<sup>src</sup> associated with other membranes as well. For example, both the src54PK protein (Fig. 7I) and the src111(Δ15–54)PK protein (Fig. 7J) associated with the plasma membrane and perinuclear membranes, in addition to cytoplasmic granules. Because the distribution of these proteins differs from src14PK, amino acids 15 to 54 and 55 to 111 must contain additional sorting information. These results imply that multiple domains of pp60<sup>src</sup> can affect the intracellular distribution of a protein.

**DISCUSSION**

Several lines of evidence suggest that the association of pp60<sup>src</sup> with membranes is mediated in part by interaction between pp60<sup>src</sup> and one or more membrane proteins. First, pp60<sup>src</sup> is found in diverse subcellular locations that vary from one type of cell to another. These locations include the plasma membrane in fibroblasts (8, 21, 44), particularly focal contacts (37); perinuclear membranes in fibroblasts (36); cytoplasmic granules in platelets (33) and chromaffin cells (30); and growth cones in neurons (27, 41). These specificities suggest that pp60<sup>src</sup> interacts with other proteins in the various locations. Second, reconstitution of purified pp60<sup>src</sup> into lipid vesicles in vitro requires the addition of membrane proteins (34), and the binding of pp60<sup>src</sup> to membranes utilizes at least one protein receptor that is both specific and saturable (35). Third, the protein tyrosine kinase encoded by lck associates with two transmembrane glycoproteins, the CD4 and CD8 antigens of T lymphocytes (38, 42), and the catalytic activity of lck can be modulated by cross-linking the CD4 antigen (43). Since the lck protein is closely related to pp60<sup>src</sup>, it seems likely that the latter also interacts specifically with membrane proteins and that the interaction may help govern the activity of pp60<sup>src</sup>.

If the attachment of pp60<sup>src</sup> to membranes resembles that of the lck protein, the following predictions should be borne out: (i) pp60<sup>src</sup> should behave like a peripheral membrane protein in schemes for solubilization; (ii) myristylation should not be sufficient for the association of pp60<sup>src</sup> with membranes; (iii) specific domains of pp60<sup>src</sup> should mediate its association with membranes; and (iv) these membrane-anchoring domains will target pp60<sup>src</sup> to specific subcellular locations. The work reported here fulfills each of these predictions.

We acknowledge one caveat. Our work was performed with chimeric proteins formed between portions of pp60<sup>src</sup> and the cytosolic protein PK. We cannot presently refute the possibility that the conformation (or other features) of these chimeras has influenced the outcome of our experiments in ways that are artifactual. It is reassuring, however, that the chimera src539PK (which contains all of the membrane-targeting domains that we have defined within pp60<sup>src</sup>) distributes within cells in a manner identical to that of pp60<sup>src</sup> itself. In addition, the conclusions reached here are in accord with previous observations on the roles of myristylation and other domains of pp60<sup>src</sup> in the attachment to membranes (for examples, see references 22, 34, and 35).

The src protein is a peripheral membrane protein. Previous reports have demonstrated that the association of pp60<sup>src</sup> with membranes cannot be disrupted by chelating agents or by extreme concentrations of salt, whereas nonionic detergents completely extract pp60<sup>src</sup> from membrane vesicles (21, 22). These results led to the provisional conclusion that pp60<sup>src</sup> is an integral membrane protein. In contrast, we have shown that pp60<sup>src</sup> is a peripheral membrane protein, because it can be extracted from membrane vesicles with alkali. Similarly, hybrid src-PK proteins that are associated with membranes are also extracted with alkali. Perhaps extraction from membranes with alkali will be a general feature of myristylated proteins associated with membranes.

The role of myristylation in the membrane association of pp60<sup>src</sup>. Myristylation is neither necessary nor sufficient for the association of pp60<sup>src</sup> with membranes. Several forms of pp60<sup>src</sup> are myristylated yet cytoplasmic (3, 11, 29). Conversely, nonmyristylated forms of pp60<sup>src</sup> have been described which still associate with membranes (12, 20, 22). Our results provide further support for the notion that the myristyl moiety is not sufficient for the association of pp60<sup>src</sup> with membranes: the myristylated proteins src7PK and src204(Δ8–111)PK are cytoplasmic when analyzed by either biochemical fractionation or immunofluorescence (see Fig. 1 for a summary).

What, then, is the role of myristylation in the attachment of pp60<sup>src</sup> to membranes? First, the modification may increase the affinity of the protein for membranes: forms of pp60<sup>src</sup> that lack myristylation but are nevertheless found on membranes can be solubilized by ionic strength alone. Second, myristylation may initiate the binding of pp60<sup>src</sup> to membranes: typically, nonmyristylated forms of pp60<sup>src</sup> never attach to membranes. The full association of pp60<sup>src</sup> with membranes requires one or more portions of the protein itself, however, and once the association is established, it may be independent of myristylation: cleavage of membrane-bound pp60<sup>src</sup> with protease removes 8 kDa from the amino terminus, yet fails to disrupt the association of the remainder of the protein with the membranes (22).

Membrane-anchoring domains target molecules to distinct subcellular locations. Previous reports have demonstrated that pp60<sup>src</sup> is found in plasma membranes, perinuclear membranes, and cytoplasmic granules. The results reported here are unusual in that pp60<sup>src</sup> is found in all of these locations in a single type of cell, a circumstance that we cannot presently explain but that might be due either to the use of COS7 cells or to the exceptional abundance of pp60<sup>src</sup> produced in the cells (10- to 20-fold more than found in cells transformed by Roux sarcoma virus [data not shown]). Whatever its cause, this circumstance allowed us to demonstrate that distinctive domains of pp60<sup>src</sup> are required for targeting the protein to each subcellular location.

Amino acids 1 to 14 of pp60<sup>src</sup> cause proteins to associate primarily with cytoplasmic granules in COS7 cells. Perhaps these amino acids specifically target pp60<sup>src</sup> to granules in chromaffin cells and platelets as well. Amino acids 38 to 111 cause proteins to associate with both plasma membranes and perinuclear membranes. We suspect that these amino acids make up multiple membrane-anchoring domains, because in conjunction with amino acids 1 to 14, either amino acids 15 to 54 or amino acids 55 to 111 are sufficient to cause association with plasma membranes, perinuclear membranes, and cytoplasmic granules. Amino acids 204 to 259, together with amino acids 1 to 14, cause a protein to associate primarily with perinuclear membranes. We conclude that the subcellular localization of pp60<sup>src</sup> is mediated by multiple domains within the protein.

Is the subcellular distribution of pp60<sup>src</sup> controlled by phosphorylation? How is it that in most types of cells, pp60<sup>src</sup> is localized to one subcellular compartment and not to others? It appears possible that the activity of membrane-anchoring domains is regulated by phosphorylation. For example, we have shown that amino acids 1 to 14 of pp60<sup>src</sup>
target proteins specifically to the membranes of cytoplasmic granules. Within and close to this domain are serine residues that are known to be phosphorylated by cellular kinases: serine 12 by kinase C (13, 14) and serine 17 by the cyclic AMP-dependent protein kinase (6, 10, 18). Perhaps phosphorylation of these sites helps regulate the subcellular location of pp60\textsuperscript{c-src}. There is precedent for this speculation in the finding that phosphorylation of the epidermal growth factor receptor by kinase C causes internalization of receptor molecules (24).

The viral and cellular src proteins differ in their amino acid sequence, including regions used here in chimeras. Since the chimeras contained sequences derived only from viral src, the conclusions that we have reached regarding the targeting of src protein to membranous compartments apply rigorously only to the viral version of the protein. Alternative splicing in neurons results in the synthesis of a novel form of pp60\textsuperscript{src} (pp60\textsuperscript{123}) (23, 28). Perhaps the localization of pp60\textsuperscript{src} in neurons to growth cones is in part due to these structural alterations.

Where within the cell is the transforming activity of pp60\textsuperscript{src} effected? Transformation by the viral version of pp60\textsuperscript{src} requires that the protein be associated with membranes (see reference 19 for a review), presumably because the crucial substrates of pp60\textsuperscript{src} are also membrane associated. Since pp60\textsuperscript{src} is capable of associating with various membranes, it is not clear whether transformation results from the activity of pp60\textsuperscript{src} at a single or at multiple subcellular locations.

Surprisingly, many of the mutant src proteins with deletions that begin at codon 15 retain transforming activity. None of these proteins has been analyzed by immunofluorescence. Our results indicate that these transforming src proteins would associate primarily with perinuclear membranes and cytoplasmic granules.

No mutant allele of src has been described in which all of the membrane-anchoring domains have been deleted. In most of the available mutants, the encoded proteins retain the first 14 amino acids of pp60\textsuperscript{src} among these mutant proteins, only the product of the NY18-3 allele is not associated with membranes (11). The NY18-3 src gene lacks only the sequences that normally encode amino acids 169 to 264, yet it encodes a myristylated protein that is located in the cytoplasm. Our results do not explain these properties. Perhaps the conformation of this protein precludes association with membranes.

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