Biological Function of PDGF-induced PI-3 Kinase Activity: Its Role in αPDGF Receptor-mediated Mitogenic Signaling

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Abstract. The tyrosine phosphorylation sites in the human αPDGF receptor (αPDGFR) required for association with PI-3 kinase have been identified as tyrosines 731 and 742. Mutation of either tyrosine substantially reduced PDGF-induced PI-3 kinase activity but did not impair the receptor-mediated mitogenic response. We sought to determine whether PDGF-induced PI-3 kinase activity could be further ablated so as to exclude a low threshold requirement for PDGFR signal transduction. Thus, we mutated both tyrosine 731 and 742 and expressed the double mutant (Y731F/Y742F) in 32D hematopoietic cells. In such transfectants, PDGF induced no detectable receptor-associated or anti-P-Tyr recoverable PI-3 kinase activity. Under the same conditions, neither mobility shift of raf-1 nor tyrosine phosphorylation of either PLCγ or MAP kinase was impaired. 32D transfectants expressing the double mutant showed wild-type αPDGFR levels of mitogenic and chemotactic responses to PDGF. To examine the effect of the double mutation in cells that normally respond to PDGF, we generated chimeras in which the cytoplasmic domains of wild-type αPDGFR, Y731F, and Y731F/Y742F were linked to the extracellular domain of colony-stimulating factor-1 (CSF-1) receptor (fms). After introduction of the chimeric receptors into mouse NIH/3T3 fibroblasts, the ability of CSF-1 to stimulate growth of these transfecteds was examined. Our data show that all these chimeric receptors exhibited similar abilities to mediate CSF-1-stimulated cell growth. These findings lead us to conclude that PDGF-induced PI-3 kinase activity is not required for PDGF-stimulated mitogenic pathway in both NIH/3T3 fibroblasts and 32D hematopoietic cells.

Interaction of PDGF with its receptors causes activation of the receptor tyrosine kinase, which leads to a cascade of biochemical events culminating in mitogenesis (Williams, 1989). Two PDGF receptor (PDGFR) genes, designated α and β, encode related proteins that can act independently to perform major PDGF signaling functions, including proliferation and chemotaxis (Matsui et al., 1989). Several cellular proteins, including phospholipase Cγ (PLCγ), GTPase-activating protein (GAP), and phosphatidylinositol-3- (PI-3) kinase become associated with and tyrosine-phosphorylated by the activated PDGFR (Meisenhelder et al., 1989; Molloy et al., 1989; Kaplan et al., 1990; Morrison et al., 1990; Kumjian et al., 1989; Coughlin et al., 1989; Kawaugh et al., 1992). These interactions are mediated by high affinity binding between the src homology 2 (SH2) domains of the substrates and specific phosphorylated tyrosine residues of receptor (Cantley et al., 1991). For example, in the human βPDGFR, phosphorylation of tyrosine 771 is necessary for association with GAP (Kashishian et al., 1992; Kazlauskas et al., 1992). In contrast, phosphorylation of tyrosine 1021 in the carboxy-terminal region is required for binding of PLCγ (Vaius et al., 1993), while PI-3 kinase interaction sites are tyrosine 740 and tyrosine 751 (Kashishian et al., 1992; Kazlauskas et al., 1992).

PI-3 kinase phosphorylates the inositol ring of PI, PI-4P, and PI-4,5P2 at the D3 position (Auger et al., 1989; Whitman et al., 1988). Purification of PI-3 kinase revealed that it consists of an 85-kD (p85) regulatory and a 110-kD (p110) catalytic subunit (Skolnik et al., 1991; Escobedo et al., 1991; Ostu et al., 1991; Hiles et al., 1992). The p85 subunit lacks PI-3 kinase activity and functions as a regulator, which binds to the activated tyrosine kinase (Hu et al., 1992; McGlade et al., 1992). Although the precise role of PI-3 kinase in regulating cell growth remains unknown, evidence that transforming p60src and polyoma middle-T mutants invariably show associated PI-3 kinase activity has suggested that this enzyme may be necessary for mitogenic signaling (Fukui and Hanafusa, 1989).
In the αPDGFR, mutation of either tyrosine 731 or tyrosine 742 substantially impairs PDGF-induced PI-3 kinase activity, but not PDGF-stimulated mitogenesis in 32D cells (Yu et al., 1991). Similar results have been observed using analogous mutants of the βPDGFR in epithelial cells (Fanti et al., 1992). These results have implied that PDGF-induced PI-3 kinase is not required for mitogenic signal transduction. However, we could not exclude the possibility that the low levels (∼5-15%) of residual PDGF-induced PI-3 kinase activity observed with such mutants were sufficient to transduce the PDGF mitogenic signal. In addition, neither the hematopoietic nor epithelial cell lines used in these investigations represented a physiological system for PDGF signaling because they do not normally express PDGFRs. To better define the role of PI-3 kinase in αPDGFR signal transduction, we generated a double mutant (Y731F/Y742F) by substituting phenylalanines for tyrosines 731 and 742 within the αPDGFR. The present studies describe the effects of these mutations on PDGF-induced PI-3 kinase activity, other downstream signaling molecules, as well as biological functions of the αPDGFR in two different cell systems.

Materials and Methods

Site-directed Mutagenesis and Generation of fms/αPDGFR Chimeras

Construction of the single mutant Y731F clone has been described previously (Yu et al., 1991). To generate the double mutant, an M13 subclone containing the cytoplasmic region of the Y731F was used as a template, and an oligonucleotide (5'-ACTACACAGTTTGTCCCCAT-3') encompassing the carboxy terminal domain of the fms tyrosine kinase (Y742F) was used to direct mutagenesis. The Y731F construct each contain different drug-resistant markers (Beckman et al., 1987a). Cells were fed with DME containing 10% calf serum as described previously. For determination of directed cell migration in soft agar, 10^5 cells were plated at 3 x 10^5 cells/ml into 24-well plates (Costar Corp., Cambridge, MA) for 24 h followed by labeling with 5 μCi/ml of [3H]thymidine for 5 h. Cells were harvested and processed for measurement of [3H]thymidine incorporation, as previously described (Heidaran et al., 1991). For measurement of in vivo αPDGFR-associated PI-3 kinase activity, quiescent 32D cells were exposed to PDGF-AA (100 ng/ml) for 5 min at 37°C, incubated with 5 mM DFP at 4°C for 5 min, and lysed in a buffer containing 20 mM Tris (pH 8), 137 mM NaCl, 2.7 mM KCl, 1 mM MgCl₂, 1 mM CaCl₂, 1% Nonidet P-40, 10% glycerol, 1 mM Na₃VO₄, 5 mM DFP, 1 mM PMSF, and 10 μg/ml aprotinin and leupeptin. Soluble lysates (2 mg) were immunoprecipitated with monoclonal anti-human αPDGFR that recognizes human, but not murine αPDGFR (Yu et al., 1994). After SDS-PAGE and electrophoretic transfer, the filter was immunoblotted with a polyclonal anti-human αPDGFR antiserum, which reacts with both human and murine αPDGFR equally well (Yu et al., 1991).

PI-3 Kinase Assay

For measurement of PI-3 kinase activity, quiescent 32D cells were exposed to PDGF-AA (100 ng/ml) for 5 min at 37°C, incubated with 5 mM DFP at 4°C for 5 min, and lysed in a buffer containing 20 mM Tris (pH 8), 137 mM NaCl, 2.7 mM KCl, 1 mM MgCl₂, 1 mM CaCl₂, 1% Nonidet P-40, 10% glycerol, 1 mM Na₃VO₄, 5 mM DFP, and 10 μg/ml aprotinin and leupeptin. Soluble lysates (2 mg) were immunoprecipitated with monoclonal anti-αPDGFR antibody (Genzyme Corp., Boston, MA) or anti-P-Tyr antibody (Upstate Biotechnology, Inc.). Immunoprecipitates were resolved on SDS-PAGE and immunoblotted with monoclonal anti-PLCγ (Upstate Biotechnology, Inc.) or monoclonal anti-MAP kinase (Sigma Immunochemicals, St. Louis, MO) or monoclonal anti-PLCγ (Upstate Biotechnology, Inc.). Immunoprecipitates were recovered with protein G-Sepharose and assayed for PI-3 kinase activity as measured by their ability to phosphorylate PI to yield PIP (Gutkind et al., 1990).

Mitogenesis and Chemotaxis Assays

For measurement of DNA synthesis, cells were washed twice with PBS and plated at 3 x 10⁵ cells/ml into 24-well plates (Costar Corp., Cambridge, MA) in RPMI-1640 medium containing 15% fetal calf serum in the absence or presence of increasing concentrations of PDGF-AA or murine IL-3 (Genzyme Corp.) for 24 h followed by labeling with 5 μCi/ml of [3H]thymidine for 5 h. Cells were harvested and processed for measurement of [3H]thymidine incorporation, as previously described (Heidaran et al., 1991; Matsui et al., 1989). For determination of directed cell migration in response to PDGF-BB, modified Boyden chambers and Nucleopore filters (5-μm pore size; Costar Corp.) were used as described above (Heidaran et al., 1991; Matsui et al., 1989).

Soft Agar Assay

For analysis of proliferation in semisolid medium, 1 x 10⁵ of NIH/3T3 transfected cells were suspended in 0.4% agarose (SeaPlaque, FMC BioProducts, Rockland, ME) in DME containing 10% calf serum as described elsewhere (Di Fiore et al., 1987a). Cells were fed with DME containing 10% calf serum in the presence or absence of 100 ng/ml of human CSF-I once per week. Colonies were stained with 4′,6-diamidino-2-phenylindole (Sigma Immunochemicals, St. Louis, MO) and scored after 2 wk.

Results

Effects of Double Mutations on PDGF-induced PI-3 Kinase Activity

We have previously shown that mutation of tyrosine 731 or 742 within the kinase insert domain of the αPDGFR impairs...
PDGFR-associated PI-3 kinase activity 95% or 80%, respectively (Yu et al., 1991). In an effort to further ablate αPDGFR-associated PI-3 kinase activity, we generated a double mutant αPDGFR (Y731F/Y742F), and cloned it into a long terminal repeat-driven expression vector (Di Fiore et al., 1987a). To analyze the double mutant, 32D cells were transfected by electroporation, and mass populations were marker selected. The level of expression of wt or mutant αPDGFR in independent 32D transfectants was found to be similar, ∼5 × 10⁵ receptors/cell (Matsui et al., 1989, and data not shown). To assess the effects of the Y731F/Y742F mutant on PDGF-induced PI-3 kinase activity, cell lysates from PDGF-stimulated or unstimulated 32D transfectants were immunoprecipitated with a monoclonal anti-αPDGFR antibody. The immune complexes were subjected to an in vitro PI-3 kinase assay (Gutkind et al., 1990).

As shown in Fig. 1 A, addition of PDGF caused >50-fold increase in phosphatidylinositol phosphate (PIP) formation in PDGFR immunoprecipitates from 32D cells expressing the wt αPDGFR (Fig. 1 A, lanes 3 and 4). As previously reported (Yu et al., 1991), the PI-3 kinase activity associated with the Y731F upon PDGF triggering was ∼5-10% of that associated with the wt receptor (Fig. 1 A, lanes 5 and 6). Under the same conditions, 32D cells expressing the Y731F/Y742F receptor showed no detectable increase in PIP level upon PDGF triggering (Fig. 1 A, lanes 7 and 8). As a control, a portion of each preparation of immunoprecipitated receptors was resolved by SDS-PAGE and immunoblotted with anti-αPDGFR antisera. The levels of receptors among the preparations varied less than twofold (Fig. 1 B).

PI-3 kinase activity has also been observed in anti-αP-Tyr immunoprecipitates after ligand stimulation (Auger et al., 1989; Kaplan et al., 1987). Moreover, the p85 subunit has been shown to be tyrosine-phosphorylated in PDGF-triggered cells (Kavanaugh et al., 1992). Thus, we also performed PI-3 kinase assays using anti-αP-Tyr immune complexes after PDGF stimulation. As shown in Fig. 1 C, the Y731F mutant showed ∼5% of wt αPDGFR immune complexes of anti-αP-Tyr recoverable PI-3 kinase activity (Fig. 1 C, lanes 5 and 6), while the double mutant resulted in complete loss of recoverable activity (Fig. 1 C, lanes 7 and 8). Anti-αP-Tyr immune complexes from cells expressing wt or mutant αPDGFRs were resolved by SDS-PAGE and immunoblotted with polyclonal anti-p85 antibody. Consistently, cells expressing αPDGFR (Y731F/Y742F) showed no increase in anti-αP-Tyr recovery of p85 upon PDGF stimulation, while cells expressing the Y731F mutant showed a low but detectable level of anti-αP-Tyr recovery of p85 as compared to that of wt αPDGFR (data not shown). All of these results confirm that mutations affecting both tyrosines 731 and 742 resulted in complete loss of PDGF-stimulated receptor-associated or anti-αP-Tyr recoverable PI-3 kinase activity.

**Effects of Tyrosine Mutations on PDGF-stimulated Activation of Other Cellular Signaling Molecules**

To study the effects of the mutations on receptor interactions with other signaling molecules, we analyzed PDGF-stimulated tyrosine phosphorylation of PLCγ and mitogen-activated protein (MAP) kinase as well as activation of raf-1. Accordingly, cell lysates of PDGF-treated or untreated 32D transfectants were immunoprecipitated with anti-αP-Tyr followed by immunoblotting with PLCγ monoclonal antibody. The levels of expression and tyrosine phosphorylation of αPDGFRs upon triggering were similar among the transfectants analyzed (Fig. 2, A and B). Fig. 2 C shows that PLCγ was tyrosine-phosphorylated at comparable levels in PDGF-stimulated cells expressing wt or mutant receptors (Fig. 2 C, lanes 4, 6, and 8). These results imply that mutation of either Y731 or Y731/Y742 impaired in vivo kinase activity of the αPDGFR for this well-characterized substrate. Such results are consistent with recent findings that the tyrosine required for βPDGFR to associate with PLCγ is tyrosine 1021 within its carboxy-terminal domain (Valius et al., 1993).

Both raf-1 and MAP kinase are cytoplasmic serine/threonine protein kinases that are indirectly activated in response to PDGF (Morrison et al., 1989; L'Allemand et al., 1991; Felech and Sanghera, 1992). The raf-1 protein kinase is re-
Figure 2. Effect of tyrosine-mutated αPDGFRs on activating downstream signal molecules in PDGF-stimulated 32D transfectants. 32D cells (lanes 1 and 2) or 32D cells transfected with wt αPDGFR (lanes 3 and 4), Y731F (lanes 5 and 6), or Y731F/Y742F (lanes 7 and 8) were either untreated (−) or treated (+) with 100 ng/ml of PDGF-AA. Clarified lysates (100 μg/lane) were resolved by SDS-PAGE and transferred to Immobilon-P (Millipore). The transferred blot was immunoprobed with anti-αPDGFR serum (A), monoclonal anti-P-Tyr antibody (B), or polyclonal anti-raf-1 antisem (D). In C and E, 2 mg of clarified lysates were subjected to immunoprecipitation using anti-P-Tyr antibody followed by SDS-PAGE. The transferred blot was immunoblotted with anti-PLCγ antibody (C) or monoclonal anti-MAP kinase antibody (E).

Mitogenic and Chemotactic Responses of Mutant αPDGFRs in 32D Cells

Expression of PDGFRs in IL-3-dependent 32D cells has previously been shown to allow efficient coupling with intracellular pathways of mitogenic and chemotactic signaling (Matsui et al., 1989). To test the biological effects of the double mutant in 32D cells, we compared mitogenic activation by wt and mutant αPDGFRs. As shown in Fig. 3 A, PDGF induced a dose-dependent increase in DNA synthesis in 32D transfectants containing either wt or double-mutant αPDGFRs with ~10–20 ng/ml required for half maximal responses. Moreover, the maximal stimulation in DNA synthesis observed with 32D cells expressing the Y731F/Y742F receptor was similar to that of transfectants expressing the wt receptor (Fig. 3 A). Thus, despite complete abrogation of PDGF-induced receptor-associated or anti–P-Tyr recoverable PI-3 kinase activity, the double mutation was not associated with any significant reduction in mitogenic signaling in 32D cells.

Chemotaxis is another major biological response mediated by αPDGFR in 32D cells (Matsui et al., 1989). Thus, we also compared the chemotactic responsiveness of transfectants containing wt and mutant αPDGFRs. As shown in Fig. 3 B, cells expressing the wt αPDGFR exhibited an eight-fold increase in chemotactic response to PDGF (100 ng/ml). 32D cells expressing the mutant αPDGFR (Y731F/Y742F) demonstrated a comparable fold increase under the same conditions. Thus, despite complete inhibition of PDGF-induced receptor-associated or anti–P-Tyr recoverable PI-3 kinase activity, the Y731/Y742 mutant showed no impairment in either mitogenic or chemotactic responses to PDGF in 32D cells.

Comparison of the Transforming Activity of wt and Mutant αPDGFR in NIH/3T3 Cells Cotransfected with PDGF-A Chain

We next sought to investigate the effect of the Y731/Y742 mutation in a cell type, NIH/3T3, which normally expresses PDGFRs. To do so, we used a PDGF-A chain transformation enhancement assay (Heidaran et al., unpublished results). PDGF-A chain transformation is known to exhibit low transforming efficiency as compared to PDGF-B chain in NIH/3T3 cells, which express both α and βPDGFRs (Beclanann et al., 1988). αPDGFR levels have been shown to be limiting for PDGF-A chain transformation, since cotransfection of constructs expressing PDGF-A and the wt αPDGFR results in increased PDGF-A–transformed foci.

As shown in Fig. 4 A, transfection with PDGF-A resulted in ~20 transformed foci/μg (Fig. 4 Aa). In contrast, cotransfection of PDGF-A chain with the wt αPDGFR resulted in ~200 foci/μg of PDGF-A chain (Fig. 4 Ab). It can be ob-
Figure 3. Comparison of PDGF-induced mitogenic and chemotactic response of 32D transfectants. DNA synthesis of 32D cell expressing wt aPDGFR (A), Y731F (O), or Y731F/Y742F (©) was measured by [H]thymidine incorporation in IL-3-free medium as described in Materials and Methods (A). Each point represents the mean value of duplicate samples. Chemotaxis was determined by directed cell migration in response to 100 ng/ml of PDGF-BB (B). Modified Boyden chambers and Nucleopore filters (5-μm pore size) were used in this assay. Similar results were obtained in three independent experiments.

Figure 4. Comparison of the ability of wt and mutant aPDGFRs to enhance PDGF-A-transforming function in NIH3T3. NIH/3T3 cells were cotransfected with PDGF-A chain and LTR-gpt marker (Aa), wt aPDGFR (Ab), Y731F (Ac), Y731F/Y742F (Ad), or K627R (Ae). Transforming foci were detected 3 wk after transfection. Cells were fixed with 70% ethanol and stained with Giemsa stain. (B) The level of human aPDGFR expressed in each transfectant was analyzed. Cell lysates from marker-selected cultures were subjected to immunoprecipitation using monoclonal anti-aPDGFR antibody that recognizes human but not the murine aPDGFR. After SDS-PAGE and electrophoretic transfer, the filter was immunoblotted with a polyclonal antibody to aPDGFR antiserum. The results of several independent experiments in which we measured enhancement of PDGF-A-transforming activity coexpressed with various aPDGFR constructs. Our results show that wt aPDGFR, Y731F and Y731F/Y742F all enhance PDGF-A-transforming activity to a similar level, as determined by the number of transformed foci/number of marker-selected colonies. Under the same conditions, transfection with the receptor constructs alone did not lead to any
Table I. Summary of αPDGFR-mediated Increase in Transforming Activity of PDGF-A Chain

| DNA transfected         | Transfection 1 | Transfection 2 | Transfection 3 |
|-------------------------|----------------|----------------|----------------|
| PDGF-A + LTR-gpt        | 4/40           | 30/500         | 30/200         |
| PDGF-A + WT             | 40/20          | 300/300        | 100/50         |
| PDGF-A + Y731F          | 100/100        | 300/300        | 150/80         |
| PDGF-A + Y731F/Y742F    | 20/10          | 150/100        | 50/30          |
| PDGF-A + K627R          | 4/20           | 20/200         | 30/200         |

Transfection was performed by the calcium-phosphate precipitation method, using 1 μg of PDGF-A cDNA, 1 μg of indicated αPDGFR cDNA, and 40 μg of calf thymus DNA as carrier. Focus formation was scored 14–21 d after transfection in unselected cultures. To measure the number of cotransfected cells, marker selection was performed in paralleled cultures using medium containing both mycophenolic acid (80 mM) and geneticin (750 μg/ml). Colonies surviving double-marker selection were scored 14–21 d after transfection.

Cellular transformation (data not shown). Thus, all of these results indicate that tyrosine mutations that specifically abrogate PDGF-induced PI-3 kinase activity do not impair the ability of the αPDGFR to enhance PDGF-A chain transformation of fibroblasts. In this cotransfection assay, we observed that the number of transformed foci exceeded the number of marker-selected colonies when PDGF-A-transformation activity was greatly enhanced. This may be caused by the differences in threshold levels of proteins required for forming foci and for surviving double-marker selection. Alternatively, it may result from the different levels of expression of transfected ligand and receptor vs that of marker-resistant genes, since they are driven by different promoters (Beckmann et al., 1988; Di Fiore et al., 1987a).

CSF-1-stimulated Cell Growth of NIH/3T3 Cells Transfected with Chimera Containing the Extracellular Domain of fms and the Cytoplasmic Domain of wt and Mutant αPDGFR

Since endogenous αPDGFR in NIH/3T3 cells may be activated in the PDGF-A cotransfection assay, we generated chimeric receptors between the extracellular domain of fms and the cytoplasmic domains of wt αPDGFR, Y731F, and Y731F/Y742F (Fig. 5 A). The chimeric receptors (designated as fms/αRWT, fms/αR731, and fms/αR731+742, respectively) were transfected into NIH/3T3 cells, which do not normally express fms. Cell lysates of NIH/3T3 transfecants treated with or without CSF-1 were resolved by SDS-PAGE and immunoblotted with either anti-αPDGFR serum against the carboxy-terminal domain of the αPDGFR that recognizes both the human and murine αPDGFR equally well (C). Total cell lysates of NIH/3T3 stimulated or not with CSF-1 were resolved by SDS-PAGE and transferred to immobilon-P. The membrane was immunoblotted with anti-αPDGFR serum against the carboxy-terminal domain of the αPDGFR that recognizes both the human and murine αPDGFR equally well (C). Total cell lysates of NIH/3T3 stimulated or not with CSF-1 were resolved by SDS-PAGE and transferred to immobilon-P. The membrane was then immunoblotted with anti-αP-Tyr.

Figure 5. Expression and tyrosine-phosphorylation of fms/αPDGFR chimeric receptors. (A) Schematic diagram of wt αPDGFR, fms, and chimeric receptors between fms and αPDGFRs. Coding regions of the αPDGFR are represented by open boxes. Coding regions of fms are represented by shaded boxes. Black boxes correspond to signal peptide (SP) and transmembrane (TM) domains. The number indicates the site of mutated tyrosines of αPDGFR. (B) Total cell lysates of NIH/3T3 cells transfected with vector alone (lane 1), fms/αRWT (lane 2), fms/αR731 (lane 3), and fms/αR731+742 (lane 4) were resolved by SDS-PAGE and transferred to immobilon-P. The membrane was immunoblotted with anti-αPDGFR serum against the carboxy-terminal domain of the αPDGFR that recognizes both the human and murine αPDGFR equally well (C). Total cell lysates of NIH/3T3 stimulated or not with CSF-1 were resolved by SDS-PAGE and transferred to immobilon-P. The membrane was then immunoblotted with anti-αP-Tyr.

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Table II. The Activity of CSF-1–induced Cell Growth of NIH/3T3 Cells Expressing fms/αPDGFR Chimeric Receptors

| Plasmid                  | Soft agar growth (%) |
|--------------------------|----------------------|
|                          | − CSF-1 | + CSF-1 |
| LTR-gpt                  | 1       | 1       |
| fms/αRWT                 | 6       | 48      |
| fms/αR731                | 6       | 53      |
| fms/αR731+742            | 6       | 38      |

Marker-selected NIH/3T3 transfectants (1 × 10⁶) were suspended in 0.4% semisolid agarose in the presence of DME supplemented with 10% calf serum. Cells were then fed with this media lacking or containing CSF-1 (100 ng/ml) once per week. The number represents the percentage of the cells forming colonies of size >30 μm in diameter after 2 wk. Results represent mean values of duplicate plates.

and the cell cultures were maintained in DME containing 2% calf serum supplemented with or without human CSF-1 (100 ng/ml). As shown in Table III, transfection with vector alone did not result in any detectable focus formation. However, transfection with fms/αRWT, fms/αR731, and fms/αR731+742 resulted in significant and similar level of focus formation in the presence of CSF-1. Together, our findings clearly suggest that ligand-induced association of PI-3 kinase activity with receptor is not necessary for αPDGFR-mediated cell proliferation and transformation in a physiologically relevant cell system.

Discussion

Our present studies demonstrate that mutations of tyrosines 731 and 742 within the kinase insert domain of the αPDGFR (Y731F/Y742F) completely impaired PDGF-induced receptor-associated or anti–P-Tyr recoverable PI-3 kinase activity. Yet the mutation of these tyrosines did not affect the ligand-dependent receptor autophosphorylation, tyrosine phosphorylation of PLC-γ, and MAP kinase or mobility shift of raf-1. In 32D cells, the ectopic expression of the mutant αPDGFR led to ligand-dependent mitogenic and chemotactic responses that were comparable to that induced by the

Table III. Comparison of the Ability of fms/αPDGFR Chimeric Receptors to Mediate CSF-1–induced Transformation of NIH/3T3 Cells

| Plasmid                  | No. of transformed foci/ no. of marker-selected colonies |
|--------------------------|--------------------------------------------------------|
|                          | Transfection 1 | Transfection 2 |
| LTR-gpt                  | −CSF-1 | 0/500 | 0/500 |
|                          | +CSF-1 | 0/500 | 0/500 |
| fms/αRWT                 | −CSF-1 | 0/500 | 0/500 |
|                          | +CSF-1 | 150/500 | 100/500 |
| fms/αR731                | −CSF-1 | 0/500 | 1/500 |
|                          | +CSF-1 | 100/500 | 100/500 |
| fms/αR731+742            | −CSF-1 | 0/500 | 0/500 |
|                          | +CSF-1 | 100/500 | 100/500 |

NIH/3T3 cells transfection was performed by the calcium-phosphate precipitation method, using 1 μg of indicated fms/αPDGFR chimeric receptor cDNA and 40 μg of calf thymus DNA as carrier. Cells were grown in medium containing 2% calf serum with or without 100 ng/ml of human CSF-1 after 5 d after transfection. Focus formation was scored 2–3 wk after transfection. Marker selection was performed in medium containing mycophenolic acid (80 mM).
wt αPDGFR. In addition, the mutant receptor demonstrated the same ability as the wt αPDGFR to enhance transforming activity of PDGF-A chain in NIH/3T3 cells. Furthermore, CSF-1 induced similar levels of cell growth and transformation of NIH/3T3 cells transfected with chimeric receptor containing the extracellular domain of fms and the cytoplasmic domain of wt or mutant αPDGFRs. Therefore, these results suggest that PDGF-induced PI-3 kinase activity is not necessary in αPDGFR-mediated mitogenic, proliferative, or transforming signaling pathways.

Previously, we have shown that mutation of tyrosine 731 or 742 of the αPDGFR impaired PDGF-induced PI-3 kinase activity 95% or 80%, respectively, but not PDGF-stimulated mitogenesis in 32D cells (Yu et al., 1991). Fantl et al. reported that similar mitogenic effects were observed using analogous single mutants of mouse βPDGFR in normal murine mammary gland (NMuMG) epithelial cells (Fantl et al., 1992). However, they found that mutation of both tyrosines of βPDGFR completely abrogated PDGF-induced PI-3 kinase activity and mitogenic response in that cell line (Fantl et al., 1992). These results suggest that a low level of PDGF-induced PI-3 kinase activity is required in PDGF mitogenic signaling. This is in contrast to our findings showing that complete abrogation of PDGF-induced PI-3 kinase activity does not impair PDGF-induced mitogenic response in 32D cells. To address this discrepancy, we used NIH/3T3 cell line since it provides a biologically relevant system to study αPDGFR-mediated signaling. Using chimeric receptors containing the extracellular domain of fms and the cytoplasmic domains of wt αPDGFR, Y731F, and Y731F/Y742F, we showed that the ability of these chimeric receptors to mediate cell growth and transformation of NIH/3T3 cells induced by CSF-1 is similar. Since fms is not expressed in NIH/3T3 fibroblasts, and endogenous αPDGFR is not cross-phosphorylated by the chimeric receptor, our data strongly argue that PDGF-induced association of PI-3 kinase activity with receptor is not required for αPDGFR-mediated cell growth and transformation in a physiologically relevant cell system. However, the exact role of receptor-associated PI-3 kinase activity in mediating migration of NIH/3T3 cells remains to be established.

Recently, Kazlauskas et al. described that an analogous double mutant of human βPDGFR in dog kidney TRMP epithelial cell can mediate PDGF-induced mitogenic response when expressed at high level (6 × 10⁶ receptors/cell) (Kazlauskas et al., 1992). Since the level of receptor expression has been shown to affect the biological property of erbB-2 receptor expressed in NIH/3T3 cells (Di Fiore et al., 1987b), it is possible that the mitogenic response mediated by double mutant of βPDGFR was caused by overexpression and a physiological level of mutant receptor expression may not be sufficient for efficient coupling of receptor to mitogenic signaling pathway. However, our data show that a double mutant of αPDGFR can transduce mitogenic and proliferation signaling, even when expressed at ~5 × 10⁴ receptors/cell in both 32D cells and NIH/3T3 cells. Since the endogenous αPDGFR is expressed at similar level in fibroblast (see Fig. 5 B and Heidaran et al., 1993), our results suggest that PDGF-induced PI-3 kinase activity is not required for mitogenic signaling, even when the mutant receptor is expressed at a physiologically relevant level.

The ligand-dependent activation of c-raf and ras have been shown to be essential for PDGF-mediated biological response in NIH/3T3 cells (Kolch et al., 1991; Cantley et al., 1991). Moreover, MAP kinase has also been implicated in mitogenic signaling pathways since it has been shown to be downstream of ras-1 and to activate c-jun in the nucleus (Kyriakis et al., 1992; Howe et al., 1992; Pulverer et al., 1991). Our data showing that mutations of tyrosines 731 and 742 of αPDGFR did not affect the mobility shift of c-raf and tyrosine phosphorylation of MAP kinase suggest this important mitogenic signal pathway to the nucleus is not impaired. We also found that activation of the wt αPDGFR and Y731F/Y742F led to a similar level of tyrosine phosphorylation of Shc molecule in 32D cells (data not shown). Shc has recently been shown to be involved in mediating ras activation in v-src transformed cells (Rozakis-Adcock et al., 1993; Egan et al., 1993). Thus, our findings also imply that the double mutation may not impair the ability of PDGF to activate ras protein in vivo. These findings are consistent with results published recently by Valius and Kazlauskas, who reported characterization of a mutant βPDGFR that did not associate with PI-3 kinase, but was still capable of activating ras upon PDGF triggering (Valius and Kazlauskas, 1993). Together, all of these data suggest that abrogation of PI-3 kinase association does not block the critical mitogenic signaling pathways in vivo.

The ligand-dependent receptor association and/or tyrosine phosphorylation of GAP (Kazlauskas et al., 1992) or PLCγ (Valius et al., 1993) have recently been shown to be dispensable for PDGF mitogenic signaling in vivo. Our present report also suggests that PI-3 kinase association is not required for PDGF-stimulated mitogenic signaling. Thus, all of these findings suggest that it is possible that any one signal molecule can be dispensable for mitogenic signaling due to sufficient redundancy. Alternatively, these results suggest that none of these molecules are directly involved in mitogenic pathways, and they may have functions yet to be identified. In this regard, the sequence homology between the p110 catalytic subunit of PI-3 kinase and a yeast protein VPS34 suggest PI-3 kinase may play a protein-sorting function in mammalian cells as in yeast (Hiles et al., 1992; Herman and Emr, 1990). Future investigation of downstream signal molecules of PI-3 kinase and the biological function of its products should allow us to further dissect the role of this enzyme in vivo.

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