A Monoclonal Antibody to the Carboxyterminal Domain of Procollagen Type I Visualizes Collagen-synthesizing Fibroblasts

Detection of an Altered Fibroblast Phenotype in Lungs of Patients with Pulmonary Fibrosis

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Abstract

Excessive collagen deposition plays a critical role in the development of fibrosis, and early or active fibrosis may be more susceptible to therapeutic intervention than later stages of scarring. However, at present there is no simple method for assessing the collagen-synthesizing and secreting activity of fibroblasts in human tissues. Type I procollagen carboxyterminal domains are proteolytically removed during collagen secretion. Thus, antibodies to these domains should stain fibroblasts synthesizing type I collagen but not extracellular collagen fibrils which could mask the signal from the cells. We developed and characterized a monoclonal antibody (Anti-pC) specific for the carboxyterminal propeptide of type I procollagen. To determine the relationship between Anti-pC staining and collagen synthesis, we stained embryonic and adult chicken tendon. Embryonic chick tendon fibroblasts actively synthesizing type I collagen stained heavily with Anti-pC, whereas quiescent tendon fibroblasts did not stain with Anti-pC. Wounded adult tendons developed fibroblasts that stained with Anti-pC at the wound site. Thus, Anti-pC specifically visualized fibroblasts actively synthesizing collagen. Lung biopsies from patients with fibrotic lung disease were stained with Anti-pC. Interstitial and intraalveolar fibroblasts in biopsies from patients with active fibrosis stained intensely with Anti-pC, while normal human lung was unstained. The absence of staining in normal lung supports the hypothesis that fibrosis is associated with an altered collagen-synthesizing phenotype of tissue fibroblasts. Anti-pC may provide a useful clinical tool for assessing fibrogenic activity at sites of tissue injury.

Introduction

The fibrotic lung diseases are a heterogeneous group of disorders reflecting a common pathophysiologic response of the lung to injury. Whereas in many cases the etiology is clear, cases are frequently of unknown cause, thus creating a significant therapeutic dilemma. Although the pathophysiologic mechanisms remain unclear, progressive connective tissue deposition characterizes fibrotic lung disorders (1). Histopathological and clinical findings currently guide therapy (2) but there is a clear need for more objective methods to assess disease activity. Determining collagen content of lung biopsies is not a reliable guide to disease activity. In addition to the obvious methodological difficulties, lung collagen may accumulate slowly or explosively, or even exist in excessive quantities as a result of prior disease activity (1,3,4). Furthermore, chemically detectable increases in collagen content may occur relatively late and thus may not be detectable as a result of focal fibroproliferative reactions. On the other hand, detecting the presence of cells actively secreting collagen in the lung would suggest active connective tissue deposition, i.e., fibrosis. Accordingly, some measure of the numbers and collagen-secreting activities of fibroblasts in fibrotic lung could be useful to guide therapy. However, no simple tool to assess the collagen-secreting activity of lung fibroblasts in biopsy specimens has emerged to date.

Because type I collagen propeptides are rapidly removed by proteolysis after collagen secretion (5), antibodies to these peptides might detect cells actively synthesizing collagen by yielding specific intracellular staining without the problems of background staining of preexisting extracellular collagen. Such antibodies might provide results similar to in situ hybridization of collagen mRNA with a cDNA. To test this hypothesis, we developed and characterized a monoclonal antibody to the carboxy terminal domain of human type I procollagen (Anti-pC)¹ and validated its use in a model system of collagen synthesis, embryonic and adult chicken tendon. In the chicken model, Anti-pC yielded selective intracellular staining of fibroblastic cells in embryonic tendon fibroblasts actively synthesizing type I procollagen (6). Adult tendons, where collagen synthesis was minimal, were not stained. However, after laceration and repair, abundant fibroblasts staining for intracellular procollagen appeared at the wound site. We then used Anti-pC to stain normal and fibrotic human lung. Healthy adult human lung did not stain with Anti-pC, whereas fibrotic lung from patients with clinically active pulmonary fibrosis contained abundant staining fibroblasts. These results suggest that Anti-pC selectively detects fibroblasts synthesizing type I procollagen and therefore represents a useful tool for the study of fibroproliferative disorders and wound healing in general.

Methods

Monoclonal antibody production. Procollagen type I was purified from ascorbate-supplemented human fetal lung fibroblast (IMR-90) cultures by ammonium sulfate precipitation and DEAE-cellulose chromatography of spent medium (7). BALB/c mice were immunized with intact type I

1. Abbreviations used in this paper: Anti-pC, monoclonal IgG to carboxyterminal domain of type I procollagen; ELISA, enzyme-linked immunosorbent assay; IMR-90, human fetal lung fibroblast; TBS, 7.5 mM Tris-HCl–150 mM NaCl, pH 7.3, containing 1% bovine serum albumin.
procollagen, spleen cells fused with Sp2/0-Ag14 myeloma cells, and growth-positive clones screened for type I procollagen reactivity by enzyme-linked immunosorbent assay (ELISA). The protocols used for monoclonal antibody production with Sp2/0-Ag14 myeloma cells including immunization, cloning, and ELISA assays have been published previously (8–10). Anti-pC, which stained IMR-90 matrix and immunoprecipitated procollagen type I from labeled fibroblast medium, was further characterized.

**Metabolic labeling and immunoprecipitation.** Metabolic labeling of IMR-90 fibroblasts with [35S]cysteine and methionine or [14C]proline and immunoprecipitation of IMR-90 medium with Anti-pC were performed as previously described (9). In some experiments, fibronectin was removed from labeled medium by gelatin-Sepharose absorption before immunoprecipitation to prevent its nonspecific precipitation.

**Immunological reagents.** Polyclonal rabbit anti-human plasma fibronectin and preimmune rabbit IgG were purified as described (11). Monoclonal mouse IgG was purified from ascites by precipitation with ammonium sulfate, molecular sieve chromatography on Sephacryl S-200 (Pharmacia, Inc., Piscataway, NJ), and DEAE chromatography. Ascites fluid (20–25 ml) containing protease inhibitors (1 mM phenylmethylsulfonyl fluoride, 1 mM N-ethylmaleimide, 20 mM EDTA, and 20 mM e-amino caproic acid in 50 mM Tris-HCl, pH 7.4 at 25°C) was clarified by centrifugation (20,000 g, 30 min). An equal volume of 50 mM sodium borate containing 200 mM NaCl, pH 7.5, was added to the supernate followed by solid ammonium sulfate (39 g/100 ml; 50% saturation at 0°C). After 4 h on ice, the precipitate containing IgG was collected by centrifugation and dissolved in 20 ml of 7.5 mM Tris-HCl–150 mM NaCl, pH 7.4. After centrifugation, the sample was applied to a 5 × 120 cm column of Sephacryl S-200 equilibrated with 50 mM Tris-HCl, pH 7.4. The peak containing IgG was then applied to a 2.5 × 10 cm column of DEAE-Sephacryl (Pharmacia, Inc.). The column was washed with 50 mM Tris-HCl, pH 7.4, and eluted with a linear gradient of NaCl from 0 to 0.5 M in the same buffer. Anti-pC eluted at 780 mg M NaCl with a typical recovery of 100 mg IgG from 20–25 ml of ascites. SDS-polyacrylamide gel electrophoresis (12), Coomassie Blue staining, and quantitative laser scanning densitometry showed that all monoclonal and polyclonal Ig preparations used in this study were at least 90–95% pure.

**Mapping of the Anti-pC binding domain using procollagen digestion products.** Procollagen was subjected to digestion with pepsin, skin collagenase, and bacterial collagenase according to published methods (9, 13–15). Crystalline pepsin (Sigma P 6887) and highly purified bacterial collagenase (type VII; see reference 9) were purchased from Sigma Chemical Co., St. Louis, MO.

**Immunohistochemistry.** Double label immunofluorescence staining of unpermeabilized fibroblasts for fibronectin and procollagen was performed as described (16) using rhodamine isothiocyanate–conjugated, affinity-purified rabbit anti-human plasma fibronectin IgG and Anti-pC, followed by fluorescein isothiocyanate–conjugated mouse-specific goat anti-mouse IgG. Control human lung was obtained from uninvolved areas of three lobectomy specimens performed for localized peripheral lung carcinomas. Samples of interstitial disease came from four open biopsies and two autopsies, as indicated in Table I. Immunofluorescence staining of lung was carried out on fresh frozen sections.

For immunoperoxidase staining, acetic acid–ethanol fixative (17) was injected through the pleura into the lung specimen to inflate the collapsed lung. The lung was immersed in the fixative overnight and embedded in paraflin wax. Sections were collected on gelatin-coated slides and air dried. Deparaffinized slides were rehydrated in 7.5 mM Tris-HCl–150 mM NaCl, pH 7.3, containing 1% bovine serum albumin (TBS) for 20

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*Figure 1. Immunoprecipitation of pepsin and collagenase digests of [14C]proline-labeled IMR-90 fibroblast medium with Anti-pC. This is a fluorogram of a 7.5% SDS-PAGE gel electrophoresis run reduced with dithiothreitol (lane 7 is unreduced). The relative mass of the standards is shown at left in kilodaltons (220, fibronectin monomer; 94, phosphorylase b; 68, bovine serum albumin; 43, ovalbumin; 25.7, chymotrypsinogen; 14.6, lysozyme). The positions of procollagen and collagen α-chains are indicated. Lanes 1–4 are from the pepsin digestion experiment. Lane 1, [14C]proline-labeled medium proteins from IMR-90 after removal of gelatin-binding polypeptides. Lane 2, sample displayed in lane 1 immunoprecipitated with Anti-pC. Lane 3, sample in lane 1 digested with pepsin. Lane 4, Anti-pC immunoprecipitate of the sample in lane 3. Note that pepsin digestion converts the procollagen chains to collagen α-chains (lane 3), and that Anti-pC does not immunoprecipitate the pepsinized collagen (lane 4). Lanes 5–7 are from the bacterial collagenase digestion experiment. Lane 5, digest of unfractonated fibroblast medium shown in lane 1. The procollagen bands disappear, but other noncollagenous polypeptides seen in lane 1 remain. Lane 6, immunoprecipitate of the sample in lane 5 containing 35- and 32-kD polypeptides (reduced). The lower polypeptide is not easily seen in this figure. Lane 7, immunoprecipitate run unreduced (this lane is from a parallel gel). The polypeptide immunoprecipitated by Anti-pC runs with a molecular mass of 110 kD unreduced. Lanes 8 and 9 show digests of labeled fibroblast medium displayed in lane 1 with purified human skin collagenase. The total digest is displayed in lane 8; the immunoprecipitated sample is in lane 9. The total digest contains pairs of labeled polypeptides of 120 and 105 kD representing the pro-TCA fragments and 56 and 63 kD representing the pro-TCB fragments resulting from mammalian collagenase cleavage of type I procollagen (13). Anti-pC immunoprecipitated the pro-TCB fragments (lane 9)."

*Figure 2. Isolation of type I procollagen and partially processed collagens from fibroblast medium by DEAE-cellulose chromatography. [35S]Methionine and [3H] cysteine-labeled IMR-90 fibroblast medium was precipitated with ammonium sulfate and chromatographed on a DE-52 DEAE-cellulose column in 50 mM Tris-HCl, 2 mM urea, pH 8.3. After elution of the nonbound material in peak A, the column was eluted with a 0 to 0.2 M NaCl gradient (dashed line). Peak A contains type I pC-collagen, fully processed collagen, and small amounts of pT-collagen. Peak B contains intact type I procollagen and pC-collagen (see Fig. 3). Peak C contains type III procollagen, which was not immunoprecipitated with Anti-pC (not shown)."
Table I. Patient Characteristics

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min and reacted for 30 or 60 min with Anti-pC (25 µg/ml) or normal mouse serum. After rinsing in TBS, bound mouse Ig was visualized by the avidin–biotin complex method with a commercial kit (Vectastain ABC Kit, Vector Laboratories, Burlingame, CA) in accordance with the manufacturer’s recommendations. In some cases, the sections were counterstained for 1 min with hematoxylin to reveal cellular detail.

Patient population. Patients referred to the Pulmonary Disease Divisions at the Washington University Medical Center, St. Louis, MO, for evaluation of unexplained dyspnea and abnormal chest roentgenograms underwent open lung biopsy for diagnostic purposes. Evidence for clinical progression included one or more of the following signs or symptoms (2): (a) increasing dyspnea with no other discernable cause; (b) deteriorating pulmonary function tests over the previous 3–6 mo; (c) increasing roentgenographic abnormalities on serial chest x-rays; (d) acute respiratory failure with no previous history of respiratory disease and a fatal outcome. Patients were judged to have quiescent disease when there was no change in categories 1–3 for a minimum of 6 mo after the biopsy.

Animal models. Anti-pC recognized both chicken and calf procollagen type I, so we used the chicken flexor tendon as a model tissue synthesizing type I collagen to examine the relationship between staining of fibroblasts with Anti-pC and collagen synthesis. Immunohistochemical studies were performed on 16-d chick embryos and adult white male leghorn chickens. Intact embryos and excised adult tendons were fixed in acetic acid–ethanol fixative and embedded, and the paraffin sections processed identically to those of lung specimens. For studies of tendon healing, we used the animal model of Birdsell et al. (18), who demonstrated more than 10-fold increases in collagen synthesis at the wound site of lacerated and repaired chicken flexor digitorum profundus tendons, and much smaller increases in collagen synthesis in the same tendon distal to the wound site (18). The tendon was exposed, lacerated, sutured, and replaced in

**Figure 3.** Immunoprecipitation of partially processed type I procollagen with Anti-pC. This is a fluorogram of samples run on a 5% SDS-PAGE gel. Lanes 1–6 (left) are run reduced, and 7–9 (right) are run unreduced. Molecular mass standards are as shown in Fig. 1. Lanes 1–3 are from peak A of Fig. 2 before and after immunoprecipitation with Anti-pC or preimmune IgG. Lane 7 displays the total labeled polypeptides, lane 2 is an Anti-pC immunoprecipitate of the sample shown in lane 1, and lane 3 is a preimmune IgG control. Anti-pC immunoprecipitates only the upper pair of disulfide-bonded polypeptides, and not the nondisulfide crosslinked type I collagen chains. The preimmune control in lane 3 contains only traces of pC-collage α-chains. Lanes 4–6 are peak B from Fig. 2 run alone (lane 4), immunoprecipitated with Anti-pC (lane 5), or preimmune IgG (lane 6). Note that Anti-pC immunoprecipitated both type I procollagen and pC-collage α-chains. Lanes 7–9 are the unreduced samples corresponding to lanes 1–3. Note that the polypeptides immunoprecipitated with Anti-pC are all disulfide crosslinked, the major polypeptide migrating near the top of the resolving gel as expected for type I procollagen. The lower weight polypeptides are compatible with type I procollagen processing intermediates (see references 5 and 13).
Figure 4. Staining of embryonic, adult, and healing adult chicken tendon with Anti-pC. 4A is a tendon section from a 16-d chick embryo. The cytoplasm of fibroblasts is stained black by Anti-pC, whereas the extracellular matrix, scanty at this stage, is unstained. 4B is a tendon from an adult chicken. There is no staining with Anti-pC. 4C is a lacerated and repaired adult tendon at the site of laceration 21 d earlier. The fibroblast cytoplasm is stained. The fine strands in the background are presumably delicate extensions of cytoplasm. All are immunoperoxidase stained. Bar, 10 μm.
the tendon sheath. After 21 d, the tendon was removed, fixed, and processed for immunohistochemistry.

**Results**

*Characterization of procollagen I antibody*

Anti-pC is an IgG that specifically immunoprecipitated procollagen type I from fibroblast medium (Fig. 1, lanes 1 and 2). To map the binding site of Anti-pC, we used several selective enzyme digestions of procollagen to release specific domains and isolated physiologically processed collagen precursors from IMR-90 medium. Pepsin treatment of culture medium converts type I procollagen to collagen by removing the amino and carboxyterminal propeptides as well as telopeptide extensions from procollagen. Anti-pC did not immunoprecipitate any polypeptides from pepsin-treated fibroblast culture medium. Therefore, Anti-pC bound to either the propeptide or telopeptide regions of type I procollagen (Fig. 1, lanes 3 and 4).

Digestion of type I procollagen with purified bacterial collagenase selectively degraded triple helical domains of procollagen, thus releasing intact the carboxyterminal telopeptide and disulfide crosslinked propeptide domains (14, 15). After digestion of labeled fibroblast medium with highly purified bacterial collagenase, several noncollagenous polypeptides remained (Fig. 1, lane 5). Immunoprecipitation of this mixture yielded two polypeptides with apparent reduced molecular masses of 35 and 32 kD (Fig. 1, lane 6). Unreduced, these polypeptides migrated with apparent molecular mass of 110 kD (Fig. 1, lane 7). The apparent molecular mass of this disulfide crosslinked polypeptide is similar to previous estimates of the molecular mass of the trimeric carboxyterminal propeptides of type I procollagen (14, 15).

Vertebrate collagenase (purified human skin fibroblast collagenase, generously provided by Dr. Howard F. Welgus, Dermatology Division, The Jewish Hospital of St. Louis, St. Louis, MO) cleaves type I procollagen at a site in the collagenous domain approximately three quarters of the length from the amino terminus, releasing a trimeric disulfide crosslinked fragment comprised of the carboxyterminal one quarter of the collagenous domains and the intact carboxyterminal propeptides (13). Fibroblast collagenase released from labeled fibroblast medium two major pairs of polypeptides ($M_r = 120, 150, 63$, and 56 kD reduced) representing the aminoterminal and carboxyterminal mammalian collagenase cleavage products of type I procollagen, respectively (Fig. 1, lane 8). The smaller of these pairs represents the carboxyterminal vertebrate collagenase cleavage product was selectively immunoprecipitated with Anti-pC (Fig. 1, lane 9). Thus, both bacterial and vertebrate collagenase digestion results were compatible with Anti-pC binding to a site either in the carboxyterminal telopeptide or propeptide domain of type I procollagen.

To further evaluate the Anti-pC epitope, we isolated intact and partially processed procollagen using DEAE-cellulose chromatography (Fig. 2) (7). Anti-pC specifically precipitated pC- and intact type I procollagen but not pN- or fully processed collagen (Fig. 3). Because processing of procollagen by cells and tissues to pN-collagen results in cleavage between the carboxyterminal telopeptide and propeptide domains, these data suggest that Anti-pC binds to the carboxyterminal domain of type I procollagen (5).

*Immunostaining results*

**Relationship of Anti-pC staining to the collagen-synthesizing activity of embryonic and adult tissues.** As expected from previous results (16, 19), Anti-pC stained cultured IMR-90 matrix identically to anti-fibronectin antibodies (not shown). This staining, as well as the staining of fibrotic lung, was completely prevented by formalin fixation. Anti-pC also stained cells in embryonic chick tissues known to be actively synthesizing type I collagen, including osteoblasts and tendon fibroblasts (6) (Fig. 4 A). To explore the relationship between onset of collagen synthesis induced by wounding and the acquisition of fibroblast staining by Anti-pC, we stained normal and wounded adult chicken tendon. As shown in Fig. 4 B, unwounded chicken tendon did not stain with Anti-pC. Three weeks after wounding, when collagen synthesis was shown to be greatly increased (18), the wound site contained abundant fibroblasts with heavy intracellular staining (Fig. 4 C), whereas the same tendon exhibited no staining 1 cm distal to the wound site (not shown).

**Staining of normal and fibrotic lung with Anti-pC.** Although normal lung (Fig. 5) lacked detectable staining with Anti-pC,
Figure 6. Spectrum of staining observed with Anti-pC in biopsies from patients with pulmonary fibrosis. A and B are from patient C.S. with rapidly progressive pulmonary fibrosis and rheumatoid disease. A is a low power view showing numerous cells in the thickened alveolar septae stained black with Anti-pC immunoperoxidase stain. Bar, 50 μm. B is a higher magnification of the same preparation as A demonstrating the fibroblast morphology of the stained cells and the failure of Anti-pC to stain extracellular collagen. The slight darkening of the alveolar lining cells...
control experiments with antibodies to fibronectin, laminin, and type IV collagen (not shown) gave the expected localization to basal lamina (20). Thus, Anti-pC stained cultured lung fibroblast matrix, previously demonstrated to contain procollagen type I (16), as well as embryonic and healing adult tendons but not normal adult tendon or lung.

Lung biopsies from six patients with pulmonary interstitial fibrosis, either idiopathic or related to other diseases (Table I), were stained with type I procollagen with Anti-pC. The spectrum of staining patterns observed is illustrated in Fig. 6, which contains low and high power views of results obtained in three different patients. Fig. 6, A and B is a biopsy obtained from patient C.S. who had rheumatoid disease and accelerated fibrosis with a fatal outcome (21). Anti-pC stained myriad fibroblastic cells in the greatly thickened interstitium. A higher magnification image of this specimen shown in Fig. 6 B displays the fibroblastic morphology of the stained cells as well as the total absence of extracellular staining. Patient D.G., who had a more slowly progressive course, showed clusters of positively stained cells (Fig. 6 C), which at higher power were fibroblastic in appearance (Fig. 6 D). However, Anti-pC was not a general stain for all lung fibroblasts, as results in clinically quiescent disease demonstrated. For example, patient E.M. had idiopathic pulmonary fibrosis for many years. The disease progressed very slowly and culminated in respiratory failure and death. The lung, massively fibrotic and honeycombed, contained abundant fibroblasts but no fibroblastic cells stained with Anti-pC (Fig. 6, D and E).

Discussion

Immunoprecipitation of the domains of procollagen type I, obtained by selective enzymatic digestion, and of physiologically processed forms of type I procollagen are compatible with Anti-pC binding to the carboxyterminal domain of type I procollagen. Anti-pC did not immunoprecipitate type III procollagen from the same cultures. Although we have not performed competitive binding assays, it is unlikely that Anti-pC will crossreact with other types of procollagen. Even polyclonal antisera to the carboxyterminal domain of type I procollagen are highly selective for type I procollagen (15).

Anti-pC has the properties of an antibody specifically detecting type I collagen—synthesizing cells. Embryonic chick tissues actively synthesizing type I procollagen such as osteoblasts and tendon fibroblasts exhibit heavy intracellular staining, whereas adult tendon fibroblasts do not stain. Fibroblasts at the wound site of lacerated adult chicken tendons developed marked intracellular staining, thus demonstrating that Anti-pC stains wound fibroblasts synthesizing type I collagen. Interestingly, Anti-pC stained lung fibroblasts in biopsies from patients with clinically active pulmonary fibrosis, but not normal lung tissue. Biopsies from a larger series of patients must be studied to correlate antibody staining with disease activity. However, we can conclude that cells containing procollagen type I in vivo can be easily identified with Anti-pC. Moreover, in active pulmonary fibrosis, the number of fibroblasts staining for procollagen and their average intracellular collagen content (and almost certainly their synthetic activity) is increased. Thus, it appears clear from this and related studies that unknown factors in pulmonary fibrosis stimulate both increased numbers of fibroblasts and a more active fibroblast synthetic phenotype (1, 3, 21, 22). Lung fibroblasts stained with Anti-pC could contain more procollagen than fibroblasts from quiescent lung simply because procollagen secretion is somehow impeded, as in the case of ascorbate deprecation (23). This is unlikely, as dermal fibroblasts maintained in vitro under ascorbate-replete conditions, where procollagen secretion should be optimal, have intracellular pools of type I procollagen readily detectable by similar immunohistological techniques (19). Moreover, we have observed similar intracellular staining of IMR-90 cultured with ascorbate using Anti-pC.

There are no reported studies on the immunohistochimical localization of type I procollagen in human lung fibrosis, and few studies employing antibodies specific for procollagen in any human tissue. In embryonic and adult skin, immunofluorescent staining shows persistence of types I and III collagen aminoterminal propeptides in collagen fibrils, whereas the carboxyterminal propeptide is not associated with collagen fibrils (24, 25). Antibodies to the aminoterminal propeptide of type III procollagen stained extracellular reticulin and collagen in liver (26).

Notably, hepatocytes in cirrhotic liver exhibited intracellular staining similar to that seen in this study, suggesting heightened synthesis of type III collagen by hepatocytes (26). In fibrotic human lung, immunohistochimical detection of types I and III collagen has revealed apparent increases in type III that correlate somewhat with disease activity (4), but no studies of procollagen staining were performed.

Anti-pC may prove useful in the noninvasive assessment of disease activity. The absence of extracellular staining for the carboxyterminal domain of type I procollagen in chicken embryos and adult human fibrotic lung suggests that this domain is rapidly cleaved and removed in vivo, as previously suggested (5). Hence, quantitative immunonassays of serum or bronchial lavage fluid for the carboxyterminal propeptide may be useful in assessing collagen synthesis and deposition. Several investigators have used quantitative assays for type III collagen precursor extension peptides ("propeptides") to study lung and liver fibrosis. Consistent with histochimical findings cited above, the amino terminal propeptide of collagen type III is elevated in serum of patients with alcoholic liver disease and hepatic fibrosis where increased concentrations may correlate with disease activity assessed histologically or clinically (27, 28). Elevated bronchoalveolar lavage fluid and serum levels of type III procollagen N-terminal peptide have also been found in patients with idio-
pathic fibrotic lung disease and sarcoidosis (29, 30). These peptides likely appear in plasma and bronchoalveolar lavage fluid because they are removed rapidly after collagen secretion. Because collagen type I is the major collagen synthesized in lung (31), levels of carboxyterminal propeptide in biological fluids obtained from patients with pulmonary fibrosis warrant evaluation.

Finally, the carboxyterminal antigen present in fetal lung fibroblast pericellular matrix deposited in tissue culture is not present in vivo, where the antigen was only detected intracellularly. This result indicates that the posttranslational proteolytic processing of procollagen is much more complete in lung and other organs than in the cultured IMR-90 we studied or in adult dermal fibroblasts (19, 32). Whereas fibroblast culture is a convenient and therefore frequently used model, the fidelity with which this model reflects the complex events of collagen synthesis, processing, and deposition in vivo is uncertain.

Acknowledgments

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References