Vitronectin (serum spreading factor): its localisation in normal and fibrotic tissue

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SUMMARY The distribution of vitronectin and fibronectin in normal and fibrotic tissue was directly compared using indirect immunofluorescence. Both glycoproteins were ubiquitously localised to loose connective tissue. Vitronectin, unlike fibronectin, was not detected in basement membranes, in normal renal glomeruli, or around smooth muscle cells of both musculares mucosae and propria of the gastrointestinal tract. The presence of vitronectin could not be shown in washed permeabilised platelets. Vitronectin was very much increased in reactive and fibrotic tissue, as was fibronectin. This was observed in lymph nodes affected by both nodular sclerosing Hodgkin's disease and by metastatic carcinoma as well as in myelofibrotic bone marrow and sclerotic glomeruli. These findings suggest that vitronectin may have an important role in the processes of inflammation and repair.

Vitronectin, also known as serum spreading factor and S-protein, is a glycoprotein that promotes the attachment and spreading of a variety of epithelial cells and fibroblasts in vitro.\(^1\)\(^2\) It is an \(\alpha\)-globulin\(^1\) that has been isolated in the form of two non-covalently associated polypeptides with molecular weights of 65,000 and 75,000. Like fibronectin,\(^4\) another adhesive glycoprotein, vitronectin has been localised to the extra-cellular matrices of various tissues,\(^5\) in addition to being associated with platelets\(^2\) and circulating in plasma at a concentration of 250–300 \(\mu\)g/ml.\(^6\)

Biologically important quantities are also present in amniotic fluid and urinary protein. It is, however, biochemically, immunologically, and functionally distinct from fibronectin as well as from laminin and chondronectin. Vitronectin interacts with several substances including heparin,\(^7\) glycosaminoglycans,\(^8\) collagen\(^9\) and glass.\(^1\)

The function of plasma vitronectin is complex and involves regulation of both the complement and coagulation systems. Podack and Miller have shown that vitronectin binds to the terminal components of the complement system (C5b–9), thereby preventing complement mediated cell lysis.\(^10\) Vitronectin also binds to heparin and neutralises its anticoagulant activity in the inhibition of coagulation factors by antithrombin III.\(^11\)\(^12\) Recently vitronectin, like fibronectin, has been shown to modulate the function of monocytes.\(^13\)

The role and source of vitronectin in tissues is at present unknown. The localisation within mesenchymal tissues suggests a possible role in the attachment of cells to the extra-cellular matrix, and that the fibroblast is the cell of origin. In this study immunofluorescence techniques were used to localise vitronectin in normal and fibrotic human tissues and to compare its distribution with that of fibronectin.

Material and methods

Mouse monoclonal anti-human vitronectin was obtained from Calbiochem (681110) and mouse monoclonal anti-human fibronectin from ICN Biomedicals Ltd (FN15-63-640-1). Fluorescein isothiocyanate (FITC) conjugated goat anti-mouse IgG was obtained from Dakopatts. Antibodies were diluted (vitronectin 1:100, fibronectin 1:200, FITC conjugate 1:10) in phosphate buffered saline (PBS), pH 7.4, before use.

Most tissues used in this study were obtained from specimens sent for routine surgical histological examination. Where such material was unavailable fresh necropsy tissue was examined. Immunofluorescence staining was performed on sections of fresh frozen tissue, subsequently fixed in methanol for 20 minutes. Myelofibrotic bone marrow was processed as previously described.\(^14\) Fixed sections were washed twice for 10 minutes in PBS and treated with primary antibody for 30 minutes. The PBS washes were repeated before treatment for 30 minutes with conjugated antibody. After two washes in PBS the

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preparations were mounted using a solution containing equal volumes of glycerol and veronal-buffered (50 mM, pH 8.6) sodium chloride (100 mM). Fluorescence was examined using a Zeiss standard RA microscope fitted with an epifluorescence condenser and HBO 50W mercury lamp. Platelets were prepared as described by Wencel-Drake et al.\textsuperscript{15} Photographs were taken with Kodachrome 400 film.

Substitution of primary antibody with non-immune mouse IgG or PBS produced negative staining. Negative results were also observed with inappropriate primary antibodies.

Results

Specific vitronectin immunoreactivity was shown in the loose connective tissue of all tissues examined. Strong positivity was noted in the supportive stroma of hepatic portal tracts and splenic cords as well as in the perimysium of skeletal and cardiac muscles. Vascular staining was located subendothelially, although it was most strong in the elastic layers (fig 1). Large vessels were more strongly reactive than capillaries. In contrast to fibronectin, vitronectin was not detected in basement membranes or around individual smooth muscle cells, being limited to the surrounding connective tissue septa. Normal renal glomeruli were consistently negative (fig 2a), contrasting with the strong positivity of mesangial cells for fibronectin (fig 2b). Both glycoproteins were present in lymph nodes, being found as an intricate network throughout except in the germinal centres where only a few fibrils were present. Visceral capsular tissue stained intensely for both vitronectin and fibronectin.
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We have previously reported a similar deposition of fibronectin in myelofibrosis and nodular sclerosing Hodgkin’s disease.\(^2\) Recently vitronectin was shown to have collagen binding site(s) which, unlike that of fibronectin, preferentially recognise(s) triple-helical collagen.\(^9\) This interaction is probably of relevance in vivo as vitronectin has been reported to co-distribute with collagens type I and III.\(^9\) Further studies are in progress to determine the relation of vitronectin and the various collagen types in fibrotic tissue.

Barnes et al have published evidence for human platelet-associated vitronectin and postulated that it may have a role in platelet adhesion or aggregation.\(^5\) We have not been able to detect platelet vitronectin using indirect immunofluorescence. Fibronectin, however, was localised to platelet \(\alpha\) granules as previously reported.\(^2\) The reason for this apparent discrepancy remains unclear.

The determination of vitronectin localisation in normal and pathological tissue is important for the understanding of its physiological function. The now reported immunoreactivity for fibrotic tissue, and the fact that vitronectin has a role in cell-substrate interaction, the coagulation and complement systems, as well as in monocyte function, suggests that it may be important in the pathophysiology of inflammation and repair.

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