The Glycoprofile Patterns of Endothelial Cells in Usual Interstitial Pneumonia

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Abstract

Background: The pathological classification of cryptogenic fibrosing alveolitis has been a matter of debate and controversy for histopathologists.

Objective: To identify and specify the glycotypes of capillary endothelial cells in usual interstitial pneumonia (UIP) compared to those found in normal tissue.

Methods: Sections of formalin-fixed, paraffin-embedded blocks from 16 cases of UIP were studied by lectin histochemistry with a panel of 27 biotinylated lectins and an avidin-peroxidase revealing system.

Results: High expression of several classes of glycan was seen de novo in capillary endothelial cells from patients with UIP including small complex and bi/tri-antennary bisected complex N-linked sequences bound by Concanavalin A and erythro-phytohemagglutinin, respectively, GalNAc residues bound by Helix pomatia and Maclura pomifera agglutinins, and L-fucosylated derivatives of type II glycan chains recognized by Ulex europaeus agglutinin-I. Glycans bound by agglutinins from Lycopersicon esculentum and Wisteria floribunda (GalNAc) as well as GlcNAc oligomers bound by Phytolacca americana and succinylated Wheat Germ agglutinin were also seen in the capillary endothelial cells of UIP. In contrast, L-fucosylated derivatives of type I glycan chains were absent in cells from cases of UIP when Anguilla anguilla agglutinin was applied, unlike the situation in normal tissue.

Conclusion: These results may indicate existence of two distinct populations of endothelial cell in UIP with markedly different patterns of glycosylation, reflecting a pattern of differentiation and angiogenesis, which is not detectable morphologically.

Keywords: Endothelial cells; Capillaries; Lung diseases, interstitial; Idiopathic pulmonary fibrosis; Polysaccharides; Concanavalin A glycotye; Lectins; Lycopersicon esculentum; Wisteria; Phytolacca americana; Succinylated wheat germ agglutinin

Introduction

The pathological classification of idiopathic pulmonary fibrosis (IPF), also known as cryptogenic fibrosing alveolitis, has been a matter of debate and controversy for histopathologists. A current classification¹ would include usual interstitial pneumonia (UIP), desquamative interstitial pneumonia (DIP), respiratory bronchiolitis interstitial lung disease (RB-ILD), acute interstitial pneumonia (AIP, Hamman-Rich disease) and non-specific interstitial pneumonia (NSIP). An analysis of deaths from 1979 to 1988 caused by UIP, showed the highest mortality rates in England and Wales followed by Scotland, New Zealand, Australia and Canada, with...
Germany and the USA having a lower incidence. The term cryptogenic fibrosing alveolitis has been redundant from histological point of view since the consensus classification of 2002; the appropriate histological pattern, presumably UIP, should be used, instead.

UIP is the most common idiopathic interstitial pneumonia accounting for over 60% of cases in different studies. It occurs between the ages of 40 and 60 years, has a mortality rate of 68% and men are affected nearly twice as often as women.

### Table 1: Lectins used in this study and their major specificities

| Acronym | Source | Major specificity |
|---------|--------|-------------------|
| GNA     | Galanthus nivalis Snowdrop | Nonreducing terminal α-D-mannose, especially the mannosyl α1,3-mannose linkage |
| NPA     | Narcissus pseudonarcissus Daffodil | α1,6-Mannose |
| HHA     | Hippeastrum hybrid Amaryllis | α1,3 and α1,6-Mannose |
| CON A   | Canavalia ensiformis Jackbean | α-D-glucosyl and α-D-mannosyl (terminal or 1,2 linked) in high mannose, intermediate and small complex N-linked sequences |
| PSA     | Piaum sativum Garden Pea | α-D-mannose in non-biased bi/tri-antennary, complex N-linked sequences |
| LCA     | Lens culinaris Lentil | Similar to, but not identical with PSA |
| e-PHA   | Phaseolus vulgaris (erythroagglutinin) Kidney Bean | Bi/tri-antennary bisected complex N-linked sequences |
| l-PHA   | Phaseolus vulgaris (leukoagglutinin) Kidney Bean | Tri/tetra-antennary, non-biased complex N-linked sequences |
| BSA-II  | Bandeiraea simplicifolia Griffonia | Terminal α and βGlcNAc1,3 > α1,6 or β1,3 >> β1,6 |
| LEA     | Lycopersicon esculentum Tomato | β1,4GlcNAc oligomers |
| ECA     | Erythrina cristagalli Coral Tree | Gal β1,4GlcNAc β1- |
| AHA     | Arachis hypogaea Peanut | Gal β1,3GalNAc β1- >Gal β1,4GlcNAc β1- |
| MPA     | Maclura pomifera Osage orange | Gal β1,3GalNAc1- >GalNAc1- |
| HPA     | Helix pomatia Roman snail | Terminal GalNAc1- |
| SBA     | Glycine max Soybean | Terminal GalNAc1- >Galβ1- |
| VVA     | Vicia villosa Hairy vetch | GalNAc1- Ser/Thr and GalNAc1,3Gal β1- |
| WFA     | Wisteria floribunda Wisteria | GalNAc1,6Gal β1- >GalNAc1,3Gal β1- |
| DBA     | Dolichos biflorus Horse Gram | GalNAc1,3(L-Fuc1,2)Gal- β1,3,GlcNAc β1- |
| UEA-1   | Ulex europaeus-1 Gorse | H type 2 antigen (αL-Fuc(1,2)-Gal β1,4GlcNAc β1-1) and Leα |
| LTA     | Tetragonolobus purpureus Lotus | L-fucosyl terminals (especially where clustered), Fuc1,6GlcNAc >Fuc1,2-Gal β1,4(Fuc1,3)-GlcNAc β, Leα |
| AAA     | Anguilla anguilla Eel | H type 1 antigen, Leα |
| BSA-1B | Bandeiraea simplicifolia Griffonia | Galβ1,3Gal β1,4GlcNAc β1- |
| SNA-1   | Sambucus nigra Elderberry Bark | NeuNAcα2,6Gal/GalNAc- |
| PTL-1   | Psophocarpus tetragonolobus Winged bean | αGalNAc |
| PTL-11  | Psophocarpus tetragonolobus Winged bean | βGalNAc, H type 1 antigen, Galβ1-3GalNAcSer/Thr |
| MAA     | Maackia amurensis | NeuNAcα2,3Galβ1- |
| PAA     | Phytolacca americana Pokeweed | Poly-N-acetyllactosamine, GlcNAc oligomers |
| s-WGA   | Succinylated- Triticum vulgaris Wheatgerm | N-acetylgallosamine |
ral heterogeneity so that low magnification shows histological variation from one field to another. Therefore, zones of interstitial fibrosis, inflammation, honeycomb change and normal lung exist in a single slide. Deposition of collagen causes thickness of the alveolar septa, which accompanies areas of honeycomb change. This is characterized by enlarged air spaces, either empty or containing admixed mucin and inflammatory cells, lined by hyperplastic alveolar pneumocytes.

More than 3000 new cases of UIP are likely to occur in the UK each year and these tend to be concentrated in areas of the country that traditionally had high levels of employment in manufacturing industries.\textsuperscript{10,11} Alveolar capillary endothelial cells are the most susceptible lung cells to non-specific alveolar injury, toxins, noxious agents and chemical substances. Knowing the patterns of glycosylation and the glycoprofiles of cells is likely to have major implications for an understanding of disease mechanisms. Generally, the study of glycans in cells and tissues and their localization is only of recent interest.

Although there is a body of work describing the lectin histochemistry of tissue sections of pulmonary carcinomas and adjacent “unaffected” lung, most studies have investigated the binding of particular cells with only one particular lectin or with a very limited number of lectins.\textsuperscript{12-14} Also these investigations have used different visualization techniques, staining procedures and scoring systems, which produce, potentially, variable and contradictory results. Although the glycan expression of capillary networks and endothelial cells of larger vessels of normal human lung have been investigated,\textsuperscript{10,15} little if any attention has been paid to the glycan expression of capillary endothelium in patients with UIP. The present study has been conducted to obtain a comprehensive view of the glycotypes of the capillary endothelium of pulmonary tissues, using lectin histochemistry to examine samples taken from both morphologically normal and diseased human lungs.

**Materials and Methods**

Formalin-fixed, paraffin-embedded lung tissue blocks from 16 patients with UIP were obtained from the archive of the Manchester Royal Infirmary. Sections (5-mm
thick) were cut as near serially as possible, dewaxed, blocked for endogenous peroxidase, rehydrated and stained with a panel of 27 biotinylated lectins (see Table 1 for their origin and binding specificities) after trypsinization according to the method of Jones and Stoddart. Briefly, biotinylated lectins were applied at a concentration of 10 or 20 mg/mL at room temperature for 30 minutes. After washing, sections were treated with avidin-conjugated peroxidase at 5 mg/mL in 0.125 M TBS, pH 7.6, containing 0.374 M sodium chloride for one hour. Subsequently, sites of lectin binding were revealed by 3,3-diaminobenzidine tetrahydrochloride and then sections were routinely counterstained with methyl green. The biotinylated lectins were obtained from Sigma apart from SNA and MAA, which were from Boehringer Mannheim, AAA from EY Laboratories Inc. and GNA, NPA, and HHA from Vector Laboratories. Removal of terminal sialyl residues was carried out, after trypsin digestion, by incubating the sections at 37 °C in a solution of neuraminidase (type VI, from Clostridium perfringens, Sigma) at 0.1 U/mL in 0.01 M sodium acetate buffer, pH 5.5, containing 1% (w/v) calcium chloride, for one hour and repeating once more with fresh enzyme. Controls included substitution of buffer for the lectin, lectin staining in the presence of competing sugars (0.2 M) and, for SNA and MAA, the neuraminidase pre-treatment. Staining intensity was ranked ‘0’ for none; ‘1’ for detectable, but weakly stained; ‘2’ for moderate, clearly stained; ‘3’ for strong staining; and ‘4’ for intense staining.

Results

Lectin specificity and the validity of the lectin histochemistry were confirmed by different controls. The overall findings are shown graphically in Figure 1. Normal capillary endothelium bound only with l-PHA, AAA, PTL-II, SNA, and MAA. Lectin staining showed the proliferation of alveolar capillary endothelial cells uniformly in UIP associated with lectin binding with ConA, ePHA, LEA, PAA, sWGA, WFA, HPA, MPA, PTL-I, and UEA-I. The pattern of UEA-I and AAA completely changed in patients with UIP in whom UEA-I selectively showed strong staining of capillary endothelial cells (Fig 2). In contrast, AAA completely failed to react with UIP capillary endothelial cells though moderate...
staining was seen in normal capillary endothelial cells. The pattern of l-PHA staining was different from that of ePHA—very high and clear affinity of l-PHA was seen with capillary endothelial cells in normal and UIP lung, whereas ePHA binding to UIP tissue was strong. The binding of l-PHA (Fig 3) slightly increased in capillaries of patients with UIP. Patchy and weak staining of endothelial cells was also observed with ConA. PTL-II, however, showed strong staining of capillary endothelial cells in UIP (Fig 4). SNA and MAA, in contrast to ConA, ePHA, LEA, and PAA, consistently stained all the capillary endothelial cells, though the reaction tended to be stronger in cases of UIP. The pattern of HPA and MPA staining in UIP cases was different—capillary endothelial cells were clearly stained by these lectins. The effect of neuraminidase on the reactions of capillary endothelial cells with AHA and ECA was also seen. Thirteen lectins (GNA, NPA, HHA, PSA, LCA, AHA, ECA, DBA, VVA, SBA, BSA-1B4, LTA, and AAA) completely failed to react with endothelia capillaries in all patients with UIP.

**Discussion**

Lectins have been used as molecular probes from the early 1970s by several investigators for histochemical studies of lung tissue. A large panel of lectins with the same staining system, as used in this study, prevents inconsistency and a wide variation in results. In addition, such a large panel of lectins enables the exploration of similar glycans with subtle differences in sugar sequences, as well as the nature of the sugars present and their chemical linkages. The use of competing sugars and enzyme pre-treatments, as seen here, also confirm and support our data. The patterns of lectin binding of the endothelial cells seen in this study had some similarities to (and differences from) previous studies. The glycan expression resulting from UIP has not been investigated up till now and thus no any comparison with previous studies is possible. As seen below, the biosynthetic activity is altered in cryptogenic fibrosing alveolitis, leading to the changes in glycan structure which suggests some changes in the activity of one or more of the glycosyltransferases.

The positive and prominent constitutive staining by l-PHA (tri/tetra-antennary non-bisected forms) of capillary endothelial cells...
Glycoprofile of Endothelial Cells

Lial cells indicates the presence of some subsets of non-bisected complex N-linked glycans, while bisected bi/triantennated forms shown by ePHA staining and high mannose, intermediate and small complex N-linked sequences (Con A) were only expressed in the capillary endothelium in UIP cases. α1,3 and α1,6-mannosyl residues, bound by NPA and HHA were absent in both the normal and diseased state.

The capillary endothelial cell binding of ConA in UIP differs from that found in normal endothelium, as previously reported GalNAca1,3(LFuca1,2) Galb1,3/4GlcNAcb1- (the DBA ligand) and terminal GalNAca1- (the SBA and VVA ligand) were not detected in capillaries endothelial cells either of normal or UIP cases. The lack of staining of normal endothelium with DBA, SBA, and AHA was also in accordance with the findings of Barkhordari, et al, and Alroy, et al, in the human, but the variable staining with BSA-IB4 observed in the latter study, differed from our findings. Staining of all the endothelium with both AHA and ECA, after neuraminidase treatment, suggests that many sequences terminating in b-galactose were probably masked by sialic acid and the intense staining by SNA and MAA, which bind to sialic acid in α2,6-(SNA) and α2,3-linkage (MAA), would confirm this. In contrast, neuraminidase pre-treatment of sections either slightly diminished or had no effect on subsequent staining with SNA suggesting the presence of both NeuNaca2,6Gal and GalNac or b-galactose, whereas the residual staining after neuraminidase treatment probably arose from its binding to galactose and/or 2-deoxy,2-acetamido-galactose only.

High expression of GalNAca1- in a1,3 linkage (HPA) appeared in the capillary endothelial cells in UIP, while the staining of the endothelium with MPA may be due to the presence of 2-deoxy,2-acetamido-a-D-galactose, b-linked galactose or the disaccharide GalNAcb1,3GalNAca1-. The results obtained with MPA, were similar to those of Sarker, et al, who described strong staining of endothelial cells also in human tissue. The staining of endothelial cells with PTL-I, but not BSA-IB4, in all cases, indicated the general presence of Gala1-, but not in type II chains.14

The negative results obtained with UEA-I in normal capillary endothelium both here and in our earlier study contrast with those of Holthofer, et al,22 Honda, et al,12 and Alroy, et al,20 who described UEA-I as a specific marker for vascular endothelium in human tissue, regardless of vessel size or ABO blood group. The expression of the fucosylated UEA-I binding glycoprotein on the endothelial cells in UIP is a distinctive finding in comparison with normal ones, while LTA does not bind in either case and AAA only binds to endothelial cells of normal capillaries. These data imply that L-fucosylated derivatives of type I glycan chains bound by AAA are present in normal endothelium, but not the simpler derivatives of type II forms shown by UEA-1 binding. In contrast, loss of AAA but the presence of UEA-I binding in UIP lung shows abolition of L-fucosylated derivatives of type I glycan chains, but an appearance of type II ones.15

In conclusion, in comparison to the glycosylation pattern of normal lung capillary endothelial cells, those taken from patients with UIP generally showed an enhanced expression of GalNAca1- in a1,3 linkage (HPA), Gala1- (MPA), and an appearance of bi/tri-antennary bisected complex N-linked sequences (ePHA). This was accompanied by a loss of type I chains (AAA) and fucosylation of type II chains (UEA-I) which were absent or inaccessible in normal controls. The alteration of glycan structure seen in UIP clearly reflects the phenotypic alterations during fibrosis. The results may also indicate the possibility of two distinct populations of endothel-
lial cells or a pattern of differentiation and angiogenesis, which is not detectable morphologically. However, the selective and high affinity of UEA-I lectin to capillary endothelium in UIP is a distinctive finding and may be therefore used as a reliable and specific marker for detection of fibrosis.

**Conflicts of Interest:** None declared.

**References**

1. Katzenstein A-LA, Myers JL. Idiopathic pulmonary fibrosis clinical relevance of pathologic classification. *Am J Respir Crit Care Med* 1998;157:1301-15.

2. Coultas DB, Zumwalt RE, Black WC, Sobonya RE. The epidemiology of interstitial lung diseases. *Am J Respir Crit Care Med* 1994;150:967-72.

3. Johnston ID, Prescott RJ, Chalmers JC, Rudd RM. British British Thoracic Society study of cryptogenic fibrosing alveolitis: current presentation and initial management. Fibrosing Alveolitis Subcommittee of the Research Committee of the British Thoracic Society. *Thorax* 1997;52:38-44.

4. Britton J, Hubbard R. Recent advances in the aetiology of cryptogenic fibrosing alveolitis. *Histopathology* 2000;37:387-92.

5. Travis WD, King TE, Bateman ED, et al. ATS/ERS international multidisciplinary consensus classification of the idiopathic interstitial pneumonias. *Am J Respir Crit Care Med* 2002;165:277-304.

6. Bjoraker JA, Ryu JH, Edwin MK, et al. Prognostic significance of histopathologic subsets in idiopathic pulmonary fibrosis. *Am J Respir Crit Care Med* 1998;157:199-203.

7. Katzenstein AL, Myers JL, Prophet WD, et al. Bronchiolitis obliterans and usual interstitial pneumonia: a comparative clinicopathologic study. *Am J Surg Pathol* 1986;10:373-81.

8. Carrington CB, Gaensler EA, Coutu RE, et al. Natural history and treated course of usual and desquamative interstitial pneumonia. *N Engl J Med* 1978;298:801-9.

9. Guerry-Force ML, Muller NL, Wright JL, et al. A comparison of bronchiolitis obliterans with organizing pneumonia, usual interstitial pneumonia, and small airways disease. *Am Rev Respir Dis* 1987;135:705-12.

10. Hubbard R, Johnston I, Coultas DB, Britton J. Mortality rates from cryptogenic fibrosing alveolitis in seven countries. *Thorax* 1996;51:711-6.

11. Johnston I, Britton J, Kinnear W, Logan R. Rising mortality from cryptogenic fibrosing alveolitis. *BMJ* 1990;301:1017.

12. Honda T, Ono K, Katsuyama T, Nakayama J. Mucosubstance histochemistry of the normal mucosa and epithelial neoplasms of the lung. *Pathol Int* 1986;36:665-80.

13. Honda T, Schulte BA, Spicer SS. Comparison of glycoconjugates at the surface of developing type II pneumocytes and Clara cells. *Histochem J* 1989;21:241-7.

14. Sarker AB, Koirala TR, Heij JH, Murakami I. Lectin histochemistry of normal lung and pulmonary carcinoma. *Indian J Pathol Microbiol* 1994;37:29.

15. Barkhordari A, Stoddart RW, McClure SF, McClure J. Lectin histochemistry of normal human lung. *J Mol Histol* 2004;35:147-56.

16. Jones CJP, Stoddart RW. A post-embedding avidinbiotin peroxidase system to demonstrate the light and electron microscopic localization of lectin binding sites in rat kidney tubules. *Histochem J* 1986;18:371-9.

17. Mazzuca M, Roche AC, Lhermitte M, Roussel P. Limulus polyphemus lectin sites in human bronchial mucosa. *J Histochem Cytochem* 1977;25:470-3.

18. McBride S, Tatrai E, Blundell R, et al. Characterisation of lectin binding patterns of mouse bronchiolar and rat alveolar epithelial cells in culture. *Histochem J* 2000;32:33-40.

19. Stoddart RW, Jones CJP. Lectin Histochemistry and Cytochemistry Light Microscopy. Lectin Methods and Protocols: Springer, 1998: pp 21-39.

20. Alroy J, Goyal V, Skutelsky E. Lectin histochemistry of mammalian endothelium. *Histochemistry* 1987;86:603-7.

21. Holthofer H, Virtanen I, Kariniemi AL, et al. Ulex europaeus I lectin as a marker for vascular endothelium in human tissues. *Lab Invest* 1982;47:60-6.

22. Toppila S, Paavonen T, Laitinen A, et al. Endothelial sulfated sialyl Lewis x glycans, putative L-selectin ligands, are preferentially expressed in bronchial asthma but not in other chronic inflammatory lung diseases. *Am J Respir Cell Mol Biol* 2000;23:492.