Altered Composition of Urinary Heparan Sulfate in Patients with COPD

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In patients with emphysema the integrity of the extracellular matrix (connective tissue skeleton) is compromised. In this study we analyzed glycosaminoglycans, which are main constituents of this matrix, in urines from patients with chronic obstructive pulmonary disease (COPD)/emphysema. Glycosaminoglycans (GAGs) were purified by anion exchange chromatography and quantified using the 1,9-dimethylmethylene blue assay. Heparan sulfate (HS) was assayed using three different chemical methods: digestion with heparitinase or with nitrous acid and by use of an adapted 1,9-dimethylmethylene blue assay. A specific epitope on the HS molecule, defined by the monoclonal antibody JM403, was determined using an inhibition enzyme immunoassay. In patients with COPD total urinary glycosaminoglycan and HS content were not altered. The JM403 epitope of HS, however, was greatly decreased in patients (0.6 versus 4.1 units/mg creatinine for control subjects, \( p < 0.0001 \)). A similar pattern was observed when patients with bronchial carcinoma with and without emphysema were compared (0.4 versus 2.4 units/mg creatinine respectively, \( p < 0.0005 \)). Patients with sarcoidosis did not show a decreased epitope content. These results indicate a structural change or an altered processing of the HS molecule in patients with emphysema. Taking into consideration the importance of HS for the stability of the alveolar extracellular matrix, this change may be associated with the pathogenesis of emphysema.


Airspace enlargement, destruction of alveoli, and loss of elasticity are the hallmarks of pulmonary emphysema. A loss of the integrity of the extracellular matrix is probably underlying the disease. The vast majority of research has focused on elastin. Breakdown of elastin, however, has not been unequivocally demonstrated in human emphysematous lung tissue (1), nor is there consensus about an increase in elastin-derived peptides in serum or urine of patients. Elevated levels of elastin-derived peptides have (2, 3), and have not (4-6) been observed. In laboratory animals, emphysema can be induced with and without destruction of elastin (1).

Other alveolar extracellular matrix molecules like collagens, proteoglycans, laminin, and fibronectin may also be of importance in the pathogenesis of emphysema. Proteoglycans deserve special attention because they play a key role in the integrity of the alveolar wall. They are involved in the fibrillogenesis and mechanical characteristics of elastin and collagen (7-9), they bind and modulate growth factors and cytokines (10), and they are powerful inhibitors of proteases, including human leukocyte elastase (11-13).

Proteoglycans are macromolecules consisting of a core protein to which negatively charged glycosaminoglycans (GAGs) are attached. GAGs are a class of long unbranched polysaccharides, consisting of repeating disaccharides of which one is an amino sugar residue and the other an uronic acid residue. In the alveolar wall, different types of GAGs are present (14, 15), including heparan sulfate (HS) which is located in basement membranes and on cell surfaces. Alterations in lung GAGs have been described in experimental as well as human emphysema (16-21). The results of these studies, however, are contradictory, likely due to differences in experimental setup. In human lung, for instance, an increase (16) and a decrease (17) in the percentage of hyaluronic acid have been found; also in human lung an increase (18) and no increase (19) in the glucosamine/galactosamine ratio has been demonstrated.

In this study we have analyzed the urinary excretion of GAGs from patients with chronic obstructive pulmonary disease (COPD) emphysema, with special emphasis on HS. For comparison, we also studied patients with bronchial carcinoma with and without emphysema, and patients with sarcoidosis. It was found that the presence of a specific epitope of HS is considerably reduced in urine of emphysematous patients, despite an unchanged total HS content.
METHODS

Patients and Control Subjects

Three groups of patients and control subjects were studied: Group 1, patients with COPD and matched control subjects; Group 2, patients with bronchial carcinoma with and without emphysema; and Group 3, patients with sarcoidosis and matched control subjects.

Physiologic data for individual patients are given in Tables 1-3. In all groups subjects with cardiovascular, alcohol-related, or renal diseases were excluded. Patients with α1-antitrypsin deficiency were also excluded.

Control subjects did not show any sign of pulmonary disease, nor any other disease. Patients and control subjects were matched for sex, age, and smoking habit. All subjects were studied according to guidelines of the Committee of Medical Ethics of the University Hospital Nijmegen and gave informed consent.

Static and dynamic lung function tests were performed with a wet spirometer and with a closed-circuit helium-dilution method (Pulmonet III; Sensormedics, Bilthoven, the Netherlands). Diffusion capacity (Kco) was measured with the single breath-holding carbon monoxide method (Sensormedics 2450) and was corrected for actual hemoglobin. Measurements were performed at least 12 h after smoking. Predicted spirometric values were derived from the European Community for Steel and Coal (E.C.S.C.) standards (22).

Arterial blood samples were taken after the patients had been sitting for at least 15 min. PaO2 and PaCO2 were measured with a Corning Ph 127 blood gas analyzer (Corning Glass, Medfield, MA).

TABLE 1

<table>
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<th>BMI (kg/m²)</th>
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Definition of abbreviations: BMI = body mass index (weight/length²); TLC = total lung capacity; RV = residual volume; FEV1 = forced expiratory volume in one second; S = salbutamol; Kco = diffusion capacity for carbon monoxide corrected for alveolar volume; % pred = percentage of the predicted value; ND = not determined.

TABLE 2

<table>
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<tr>
<th>Patient Code</th>
<th>Age (yr)</th>
<th>Sex</th>
<th>TLC (% pred)</th>
<th>RV (% pred)</th>
<th>FEV1 (% pred) before S</th>
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<th>Kco (kPa)</th>
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<th>PaCO2 (kPa)</th>
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</table>
| Patients with bronchial carcinoma and emphysema

| 15 | 64 | F | 113 | 119 | 44 | 47 | 53 | 8.1 | 5.6 | 40 |
| 67 | 60 | M | 94 | 114 | 61 | 58 | 67 | 10.6 | 5.3 | 35 |
| 72 | 75 | M | 136 | 120 | 64 | 68 | 78 | 10.6 | 5.1 | 10 |
| 44 | 58 | M | 116 | 157 | 64 | 67 | 68 | 8.3 | 4.6 | 20 |
| 71 | 66 | M | 112 | 105 | 42 | 39 | 46 | 8.0 | 5.5 | 40 |
| 45 | 70 | M | 96 | 100 | 75 | 74 | 71 | 8.9 | 5.4 | 50 |
| 58 | 77 | M | 109 | 95 | 80 | 78 | 52 | 10.3 | 5.4 | 15 |
| 18 | 78 | M | 132 | 89 | 70 | 71 | 54 | 9.5 | 4.9 | 20 |

Patients with bronchial carcinoma, without emphysema

| 21 | 58 | M | 79 | 65 | 76 | 76 | 120 | 13.5 | 5.2 | 20 |
| 65 | 68 | M | 102 | 106 | 75 | 75 | 86 | 7.0 | 5.48 | 20 |
| 37 | 72 | M | 94 | 106 | 104 | 104 | 123 | 9.85 | 4.8 | 10 |
| 41 | 96 | M | 93 | 103 | 75 | 75 | 90 | 7.7 | 5.0 | 25 |
| 3 | 68 | M | 85 | 78 | 71 | 75 | 89 | 10.6 | 4.5 | 30 |
| 8 | 81 | M | 103 | 81 | 110 | 115 | 83 | 11.2 | 4.9 | * |
| 56 | 64 | M | 66 | 65 | 47 | 50 | 93 | 9.2 | 5.6 | 40 |
| 39 | 63 | M | 98 | 110 | 71 | 75 | 104 | 8.2 | 5.3 | 30 |

For definition of abbreviations, see Table 1.

* Nonsmoker.
Group 1: Patients with COPD and control subjects. Physiologic data of all patients are shown in Table 1. Patients were characterized by severe airflow obstruction and (in most cases) hyperinflation, and a reduced Kco. No improvement of FEVi was observed after inhalation of 400 µg salbutamol. Patients were in a stable condition, without exacerbations during the preceding 3 mo. All subjects used inhaled β₂-sympathomimetimics and/or anticholinergics. Fourteen patients (70%) used inhaled corticosteroids in a mean dosage of 630 ± 240 µg daily. None used theophylline or oral corticosteroids. Lung function data and body measures of control subjects are given in Table 4.

Group 2: Patients with bronchial carcinoma with and without emphysema. Physiologic data of all individual patients are shown in Table 2. Patients were characterized by a recently detected and histologically proven bronchial carcinoma (in all cases squamous cell carcinoma). Patients with bronchial carcinoma without emphysema had no radiographic or histologic signs of emphysema, and total lung capacity and diffusion capacity were within predicted mean ± 1.64 SD.

Group 3: Patients with sarcoidosis and control subjects. Physiologic data of all individual patients are shown in Table 3. Patients had histologically proven sarcoidosis stage 2 (mediastinal and/or hilar lymphadenopathy and radiographic signs of interstitial involvement). Six of 15 patients used oral corticosteroids in a dosage of 5 to 25 mg/d, whereas the other patients did not receive any medication. Lung function data and body measures of control subjects are shown in Table 5.

Purification of Glycosaminoglycans from Urine

Urines were collected and stored at −70°C. After thawing, 20-ml samples of urine were diluted with an equal volume of 7 M urea/10 mM Tris-HCl (pH 6.8) and applied to an ion-exchange column of 5 × 0.5 cm containing 1 ml diethylaminoethyl (DEAE)-Sepharose Fast Flow (Pharmacia, Upppsala, Sweden). After an initial wash with 3 ml 0.2 M NaCl/10 mM Tris-HCl (pH 6.8), GAGs were eluted with 1 ml 2 M NaCl/10 mM Tris-HCl (pH 6.8) followed by 3 ml 10 mM Tris-HCl (pH 6.8). The recovery of the GAGs was monitored by adding 15,000 dpm [35S]GAGs (specific activity 106 dpm/µg GAG) to the urine. Salts were removed by ethanol precipitation of the GAGs. Five volumes of ethanol were added (final ethanol concentration 83%) and GAGs were allowed to precipitate for 16 h at −20°C. After centrifugation for 30 min at 10,000 × g (−10°C), the precipitated GAGs were dried and dissolved in 2 ml demineralized water. Recoveries were between 85 and 100%. Two cycles of freeze-thawing of urine did not influence the results. A third cycle occasionally gave erroneous data.

Quantification of Glycosaminoglycans

Sulfated GAGs were quantified according to the method of Farndale and coworkers (23). This method is based on a metachromatic shift in absorption maximum of 1,9-dimethylmethylene blue (DMMB) (Aldrich Chemical Co., Bornem, Belgium). To 100 µl GAG sample, 2.5 ml of DMMB reagent was added and the absorbance at 525 nm was measured immediately. The DMMB reagent contains 46 µM DMMB (initially 46 µmol DMMB was dissolved in 5 ml 96% ethanol), 40 mM glycine, and 42 mM NaCl, adjusted to pH 3.0 with 1 M HCl. Chondroitin 4-sulfate (Sigma, St. Louis, MO) was taken as a standard and included within each series of assays.

Quantification of Total Heparan Sulfate

Three methods for the determination of HS were used. These methods are based on different aspects of the HS structure. Bovine kidney HS (Seikagaku, Tokyo, Japan) was used as a standard.

Method 1 is a modified version of the 1,9-dimethylmethylene blue assay and is based on the backbone structure of heparan sulfate (24). Addition of bovine serum albumin and adaptation of pH and salt concentration of the DMMB reagent results in a specific elimination of HS-based absorbance. To 100 µl GAG sample, 100 µl 3% bovine serum albumin (BSA) (in demineralized water) and 2.4 ml of the modified DMMB...
reagent was added. This reagent contains 48 \mu M DMBB (initially 48 
\mu mol DMBB was dissolved in 5 ml 95% ethanol, 42 mM glycine, and
63 mM NaCl, adjusted to pH 2.75 with 1 M HCl. After 30 min at 22° C
the absorbance was measured at 525 nm. By subtracting the value thus
obtained (representing other GAGs) from the total GAGs, the content
of HS was calculated (24).

Method 2, digestion of HS by heparitinase, is based on cleavage of
glycosidic linkages in HS involving glucuronic acid residues. To lyophy-
lized GAG samples, 25 \mu l heparitinase (Seikagaku, Tokyo, Japan) (0.01
IU/ml containing 0.05 M sodium acetate and 0.05 M calcium acetate
[pH 7.0]) was added. Incubation was for 16 h at 43° C. The remaining
GAGs were quantified using the 1,9-dimethylmethylene blue assay (see
quantification of GAGs). By subtracting this value from the total GAGs
the content of HS was determined.

Method 3, digestion with nitrous acid, is based on cleavage of glyco-
sidic linkages involving hexosamine residues in which the amino groups
are either N-sulfated or unsubstituted. Nitrous acid digests HS and hepa-
rin, but not other GAGs (25). Purified GAGs were treated for 90 min
at 22° C with nitrous acid prepared by mixing equal volumes of 5%
(wt/vol) \( \text{NaNO}_2 \) and 1.5 M \( \text{HCl} \). The residual GAGs were quantified
using the 1,9-dimethylmethylene blue assay (see quantification of GAGs).
By subtracting this value from the total GAGs the content of HS and
heparin was obtained. The contribution of heparin is negligible.

Quantification of the HS JM403 Epitope

The HS JM403 epitope, defined by monoclonal antibody (mAb) JM403
(26, 27), was quantitated using an inhibition enzyme immunoassay (Figure
1). In the wells of polystyrene microtiter plates (Greiner GmbH, Frick-
enhausen, Germany; catalog number 655061), 100 \mu l of ascites contain-
ing mAb JM403 (diluted 1:40,000 in Tris-buffered saline containing 0.1%
(wt/vol) NaNOz and 1.5 M HC1. The residual GAGs were quantified
using the 1,9-dimethylmethylene blue assay (see quantification of GAGs).
By subtracting this value from the total GAG the content of HS was
determined.

Figure 1. Standard curve for the inhibition immunoassay for the hepa-
ran sulfate epitope JM403. A volume of 100 \mu l containing various
amounts of bovine kidney HS was incubated for 16 h at 4° C with
50 \mu l of JM403 antibody. Next, 100 \mu l of this mixture was trans-
ferred to a well previously coated with 1 \mu g HS. After incubation for 2 h
at 22° C, the antibodies bound to the well were detected by addition
of alkaline phosphatase-conjugated secondary antibodies. Af-
her addition of substrate the color developed was spectrophotometri-
cally measured at 405 nm. Data are mean ± SD (n = 3).

RESULTS

Pilot experiments showed that the urinary content of HS JM403
epitope and of HS on base of creatinine remained constant during
the day, and from day to day. This also holds for GAGs, which
is in agreement with data from the literature (28). There was no
correlation between age and urinary content of HS and GAG.
There was, however, a correlation between age and content of
HS JM403 epitope (r = 0.26, p < 0.03).

In patients with COPD the content of urinary GAGs was not
altered compared with controls (Figure 2, Table 6). The urinary
content of HS, measured by three different methods, was also
not changed (Figure 3, Table 6). The urinary content of the HS
JM403 epitope was significantly decreased in patients with COPD
(Figure 4, Table 6).

To examine the specificity of this finding, we studied two ad-
ditional groups of patients: patients with bronchial carcinoma
with and without emphysema, and patients with sarcoidosis. The

Figure 2. Glycosaminoglycan content in the urine of individual con-
trol subjects and patients with COPD. Open squares: control; closed
squares: patient with COPD.
The specificity of the decreased HS JM403 value in COPD patients was confirmed by data of two additional groups of patients (patients with bronchial carcinoma with and without emphysema, and patients with sarcoidosis). Because all but one patient in the group of bronchial carcinoma are smokers, a major effect of smoking is not likely. Likewise, a major effect of medication is also unlikely because the group of patients with sarcoidosis who received oral corticosteroids had no decrease of urinary HS JM403 epitope. The content of HS determined with three different assays, based on different aspects of the HS molecule, showed no significant differences between patients and control subjects. Differences in urinary GAG and HS content, however, may be obscured by the relatively small contribution of GAG and HS derived from the lung. The amount of lung-derived HS JM403 epitope may, however, be quite substantial. Lung parenchyma is rich in basement membranes and the HS JM403 epitope is basement membrane specific (26). In lung the proportion of JM403 epitope per milligram of HS is quite high, e.g., in rat lung 35-fold higher than in kidney and heart (Van Den Born and coworkers, unpublished data).

**DISCUSSION**

In the urine of patients with COPD we found a decreased content of the HS JM403 epitope together with a normal content of HS. The major group of patients studied (Group 1) was identified on a clinical basis, including lung function tests, and had a strong reduction of predicted FEV1. Although we had no histologic evidence, the presence of emphysema is likely in the lungs of patients with COPD (29). In addition, the group of patients with bronchial carcinoma with histologically proven emphysema showed a reduced content of the HS JM403 epitope.

All values were normalized to the creatinine content, but COPD patients often have a reduced concentration of creatinine in the urine because of reduced muscle mass (29). Correcting for this would result in a downward adjustment of the values for COPD patients. This would make the difference between the HS JM403/mg creatinine values of control subjects and COPD patients even greater. A decreased HS JM403 value as a result of blocking of the epitope by a component present in the urine is unlikely, since prior to analysis, urine samples were first diluted with an equal volume of 7 M urea/10 mM Tris-HCl. Urea is a chaotropic agent which disrupts noncovalent bonds. GAGs were subsequently purified using ion-exchange chromatography.

The specificity of the decreased HS JM403 value in COPD patients was confirmed by data of two additional groups of patients (patients with bronchial carcinoma with and without emphysema, and patients with sarcoidosis). Because all but one patient in the group of bronchial carcinoma are smokers, a major effect of smoking is not likely. Likewise, a major effect of medication is also unlikely because the group of patients with sarcoidosis who received oral corticosteroids had no decrease of urinary HS JM403 epitope. The content of HS determined with three different assays, based on different aspects of the HS molecule, showed no significant differences between patients and control subjects. Differences in urinary GAG and HS content, however, may be obscured by the relatively small contribution of GAG and HS derived from the lung. The amount of lung-derived HS JM403 epitope may, however, be quite substantial. Lung parenchyma is rich in basement membranes and the HS JM403 epitope is basement membrane specific (26). In lung the proportion of JM403 epitope per milligram of HS is quite high, e.g., in rat lung 35-fold higher than in kidney and heart (Van Den Born and coworkers, unpublished data).
The normal content of HS and the decreased content of the HS JM403 epitope, suggest a structural alteration in or an altered processing of the HS molecule in the lungs of emphysematous patients. At present, we do not know the exact nature of the epitope recognized by the JM403 antibody. Studies on chemically and enzymatically modified HS indicate that the epitope contains one or more N-unsubstituted glucosamine and D-glucuronic acid units, and is located in a region of the heparan sulfate chain composed of mixed N-sulfated and N-acetylated disaccharide units (30). The biologic significance of N-unsubstituted glucosamines has recently been demonstrated in the binding of HS to L-selectin (31). Blocking of the epitope in the glomerular basement membrane by injection of JM403 antibody leads to proteinuria in rats (27). Interestingly, microalbuminuria has recently been described in a small group of patients with COPD (32).

The etiology of pulmonary emphysema is not known. Considering that the development of (clinically evident) emphysema is a process of several decades, the basic alteration may be quantitatively small in nature. The decreased content of a specific epitope of HS, together with a normal content of total HS, in the urines of emphysema patients hints at a small, but significant alteration. Alterations in the HS molecule would have consequences for the protecting HS-proteoglycan barrier of the alveolus (14). In view of the functions of HS (antiprotease activity, binding and modulation of growth factors and cytokines, involvement in the fibrillogenesis of collagen and elastin), this could lead to a destabilization of the extracellular matrix, ultimately resulting in emphysematous lesions. This concept is supported by the findings in rats that specific inhibition of proteoglycan synthesis induces emphysematous lesions (33) and that HS proteoglycans are very vulnerable toward elastase degradation (34). In a genetic model for emphysema, the tight-skin mouse, antibodies to HS have been detected (35).

### TABLE 7

**Urinary content of glycosaminoglycans, heparan sulfate, and heparan sulfate JM403 epitope of patients with bronchial carcinoma with and without emphysema**

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<td>Glycosaminoglycan, µg per mg creatinine</td>
<td>16.7 ± 2.4</td>
<td>16.5 ± 2.2</td>
<td>NS</td>
</tr>
<tr>
<td>Heparan sulfate</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Modified DMMB assay, µg</td>
<td>6.4 ± 0.7</td>
<td>6.6 ± 0.7</td>
<td>NS</td>
</tr>
<tr>
<td>Epitope JM403, U</td>
<td>2.4 ± 0.5</td>
<td>0.4 ± 0.1</td>
<td>p &lt; 0.0005</td>
</tr>
</tbody>
</table>

NS = not significant.
* Values are mean ± SEM. Characteristics of patients are given in Table 2.

### TABLE 8

**Urinary content of glycosaminoglycans, heparan sulfate, and heparan sulfate JM403 epitope of control subjects and patients with sarcoidosis**

<table>
<thead>
<tr>
<th>Content</th>
<th>Controls (n = 41)</th>
<th>Patients (n = 13)</th>
<th>Significance</th>
</tr>
</thead>
<tbody>
<tr>
<td>Glycosaminoglycan, µg</td>
<td>14.3 ± 0.5</td>
<td>12.8 ± 0.9</td>
<td>NS</td>
</tr>
<tr>
<td>Heparan sulfate</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Modified DMMB assay, µg</td>
<td>4.3 ± 0.3</td>
<td>3.0 ± 1.0</td>
<td>NS</td>
</tr>
<tr>
<td>Epitope JM403, U</td>
<td>2.4 ± 0.4</td>
<td>1.7 ± 0.2</td>
<td>NS</td>
</tr>
</tbody>
</table>

NS = not significant.
* Values are mean ± SEM. Characteristics of control subjects are given in Table 5, of patients in Table 3.
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References