Repetitive Fragmentation Products of Albumin and α1-Antitrypsin in Glomerular Diseases Associated with Nephrotic Syndrome

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Even if nephrotic syndrome is characterized by massive urinary loss of major plasma proteins, a clear structural characterization based on proteomics has never been reported. Urine and plasma of 23 patients with different idiopathic nephrotic syndromes (10 steroid-sensitive minimal-change nephropathy, seven steroid-resistant FSGS, and six membranous glomerulonephritis) were analyzed with two-dimensional electrophoresis in soft gel, Western blot, and matrix-assisted laser desorption/ionization time of flight mass spectrometry; 72 urinary components corresponded to fragments of albumin and/or of α1-antitrypsin. Several repetitive fragmentation motives and a few differences among different pathologies were found. Several (21 of 72) urinary albumin fragments also were detected in plasma, although in lower concentration, suggesting a preferential excretion. The bulk of components with low molecular weight were detected only in urine, suggesting an in situ formation; zymograms with albumin as substrate showed the presence in urine of specific proteases. A final but not secondary point was the characterization of albumin adducts that harbor both the COOH and NH2 terminal parts of the protein, suggesting the formation of new covalent chemical groups. Altogether, these new findings reveal unexpected structural and functional aspects of proteinuria that may play a key role in pathogenesis. Characterization of urinary fragmentation patterns should be extended to other renal diseases.


Nephrotic syndrome is a clinical condition that is characterized by severe hypoproteinemia that is produced by protracted losses in urine of major plasma proteins that are not counterbalanced by resynthesis. The large spectrum of renal diseases that cause proteinuria and nephrotic syndrome ranges from primary diseases such as FSGS, minimal-change nephropathy (MCN), and membranous glomerulonephritis (MGN) to secondary conditions such as diabetes and amyloidosis (1). Although each renal disease that leads to proteinuria has specific pathogenic mechanisms (immunologic versus metabolic versus plasma factors), there is a common background that includes structural glomerular alterations, in particular of podocytes, with failure in control of protein trafficking (2–4). The identification of specific biomarkers in urine should greatly improve the diagnostic potential and also direct the clinical management. The great accessibility of urine makes this fluid an ideal target for the search of disease markers (5,6).

The definition of the mechanisms of glomerular filtration of proteins represented a historic research advance in the early 1980s, when it became clear that, in the presence of a normal slit diaphragm, sieving properties mainly are a function of both dimension and charge of any given molecule and that also conformation plays a role (7). It also was postulated that variations in protein structure could influence proteinuria in pathologic conditions, such as FSGS, but a clear demonstration of this hypothesis has never been reported so far (8–10). Few data are available on structural aspects of plasma proteins, in particular of albumin in human renal diseases that are associated with nephrotic syndrome. Advances in the field of proteomics allow today a definite structural approach, including a fine structure definition, and this new technology potential should prompt us to revise our knowledge on the structural aspects of albumin and of other major plasma proteins in nephrotic syndrome. Within the frame work of a collaborative program on proteomics of biologic fluids, we performed studies on the composition of plasma and urine in different pathologic conditions that cause proteinuria, such as primary nephrotic syndrome in children (MCN and FSGS) and MGN in adults. We describe here an
unexpected finding: The presence of fragmentation products of two major plasma proteins, albumin and α1-antitrypsin (α1AT). The finding of both repetitive fragmentation motives and specific patterns in different renal diseases suggests common fragmentation pathways as well as specific mechanisms. In the case of albumin, we also could describe the formation of adducts that suggest stable changes of fine protein structure. As a whole, these data suggest new, unexplored mechanisms underlying kidney damage in primary renal diseases.

Materials and Methods
Patients
Three groups of patients with nephrotic syndrome and different pathologic backgrounds were enrolled in the study (see Table 1); the diagnosis was based on clinical data and on renal histology. The general criteria for enrollment were (1) presence of nephrotic syndrome (proteinuria >40 mg/h per m²) with onset at <16 yr of age and, in cases of steroid dependence and/or resistance, availability of a renal biopsy; (2) absence of familial traits of nephrotic syndrome and/or relevant mutations of slit-diaphragm genes (NPHS2, CD2AP, and ACTN4) (11–14); and (3) presence of nephrotic syndrome at any age with MGN. Molecular screening was done as already described (CD2 and ACTN4) (12). Blood samples that were collected in EDTA and morning fresh urine that was collected in sterile conditions were immediately centrifuged at 3°C for 10 min, then stored at –80°C for <4 mo before processing. In all cases, plasma and urine were collected at the beginning of proteinuria (i.e., before the starting of any therapy). Clinical parameters included blood cell counts, serum creatinine and urea levels, cholesterol, triglycerides, liver function tests, antinuclear antibodies, antineutrophil and antibodies against native DNA, and antibodies against hepatitis virus B and C. All evaluations were performed according to international standardized procedures. Renal biopsies were processed by standard procedures that included hematoxylin-eosin, periodic acid-Schiff, and green Masson staining; immunofluorescence of frozen samples was carried out with a panel of anti-serum protein antibodies against IgA, IgM, IgG, C3, and C4. Biopsy samples were evaluated by a pathologist who discussed all relevant aspects with clinicians. Adult patients and parents (for patients who were younger than 18 yr) were requested to give their informed consent to DNA analysis. The following groups of patients were studied.

Steroid-Sensitive or Steroid-Dependent Primary Nephrotic Syndrome (MCN). Ten children with nephrotic syndrome had been treated with prednisolone (2 mg/kg starting dosages followed by tapering) (15,16) and presented a good initial response. Six patients were in stable remission at the end of the therapeutic cycle; another four presented steroid dependence and received the minimum amount of prednisolone required to achieve remission. Renal morphology was analyzed to define renal lesions before starting therapy with major immune suppressors. Renal histology showed MCN on the basis of absence of immune deposits in glomeruli and of histologic lesions. Electron microscopy was not required in these cases.

Steroid-Resistant Primary Nephrotic Syndrome (FSGS). Seven children had strict resistance to prednisolone given at the dosage above and received cyclosporine as associated therapy. Cyclosporine was given in all cases at 5 mg/kg starting dosage, followed by tapering to reach the minimum dosage required for maintaining cyclosporine serum levels between 50 and 100 ng/ml (17). All of these patients had a histologic diagnosis of FSGS on the basis of the presence of at least one segmental area of glomerulosclerosis in a histologic sample that contained at least six glomeruli.

MGN. Six patients had a histologic diagnosis of MGN performed at the onset of proteinuria; they were adults with the exception of a young girl aged 12 yr. After the diagnosis, all patients with MGN received steroids as single drug (1 to 2 mg/kg) or in combination with immune suppressors (18) or angiotensin-converting enzyme inhibitors (ramipril 5 to 8 mg/m²). MGN was considered a primary form on the basis of negativity of relevant immunologic parameters (antinuclear antibody, nDNA, C3, and ANCA) and viral tests

Normal Control Subjects. The control group consisted in 10 normal donors of the laboratory staff (five women and five men; mean age 28 yr; range 22 to 34 yr) and their children (n = 6; four girls and two boys; mean age 4 yr; range 2 to 6 yr). Plasma and morning fresh urine were collected in sterile conditions and immediately frozen at –80°C as above.

Table 1. Clinical and biochemical parameters in 23 patients with idiopathic nephrotic syndrome according to renal pathology (10 steroid-sensitive MCN, seven steroid-resistant FSGS, and six MGN)a

<table>
<thead>
<tr>
<th></th>
<th>Gender (M/F)</th>
<th>Age (yr)</th>
<th>Age at Onset (yr)</th>
<th>Proteinuria (g/24 h)</th>
<th>Serum Creatinine (mg/dl)</th>
<th>Serum Cholesterol (mg/dl)</th>
<th>Serum Albumin (g/dl)</th>
<th>Therapy (St/CsA/ACEi)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Steroid-sensitive MCN</td>
<td>10</td>
<td>7/3</td>
<td>8.2 (3 to 16)</td>
<td>3.0 (1.5 to 6)</td>
<td>6.5 (2 to 7)</td>
<td>0.5 (0.3 to 0.8)</td>
<td>348 (230 to 670)</td>
<td>12 (0.9 to 1.4)</td>
</tr>
<tr>
<td>Steroid-resistant FSGS</td>
<td>7</td>
<td>5/2</td>
<td>8.5 (5 to 16)</td>
<td>3.1 (1 to 7)</td>
<td>4.8 (2.7 to 7)</td>
<td>0.6 (0.4 to 1.0)</td>
<td>300 (200 to 460)</td>
<td>1.6 (1.1 to 2.2)</td>
</tr>
<tr>
<td>MGN</td>
<td>6</td>
<td>4/2</td>
<td>39 (12 to 55)</td>
<td>36 (10 to 53)</td>
<td>6 (1 to 8)</td>
<td>1.1 (0.5 to 1.4)</td>
<td>280 (220 to 500)</td>
<td>1.5 (1.1 to 2)</td>
</tr>
</tbody>
</table>

aIn all cases, plasma and urine were collected before any therapy was started and before a correct diagnosis was made on the basis of clinical and pathologic findings. All patients who were classified as steroid-sensitive MCN presented a good sensitivity to steroids, although with some relapses of proteinuria. In these cases, MCN was diagnosed on the basis of the absence of Ig deposition in glomeruli and of any histologic alteration. Electron microscopy was not required in these cases. Results are given as mean range of variation. NS, nephrotic syndrome; MCN, minimal-change nephropathy; St, steroids; CsA, cyclosporine; ACEi, angiotensin-converting enzyme inhibitors.
ampholyte cocktail) that contained 60% of the pH 3.5 to 10 and 40% of the pH 4 to 8 intervals. Before isoelectric focusing, samples were incubated in this solution for 3 h to allow proper reduction and alkylation. For prevention of overalkylation during the isoelectric focusing step, excess of iodoacetamide was destroyed by adding a molar amount of dithiothreitol. The first dimension strips that were used for 2D maps were 18-cm-long, soft, homemade, immobilized-pH-gradient gels. In the second dimension, proteins were separated on the basis of their size in 6 to 16% T gradient polyacrylamide gel slabs that were 180 × 160 × 1.5 mm.

For Western blot, proteins were transblotted to nitrocellulose membranes Protean BA (Schleicher & Schuell, Dassel, Germany) with a Novablot semidry system using a continuous buffer system with 2-amino 2-idoxymethyl 1,3-propanediol tris 38 mM, glycine 39 mM, SDS 0.035%, and methanol 20%. The transfer was achieved at 1.55 mA/cm² for 1.5 h.

### Gel/Membrane Staining and Image Analysis

After separation in SDS-PAGE gels, the proteins were visualized by a double-staining procedure: First the methyl-trichloroacetate negative staining (20) followed by the silver staining or Blue silver colloidal Coomassie (21) for preparative mass spectrometry analysis. Images of stained gels were digitized using a GS800 photometer and were acquired using a Versa DOC 400. All images were analyzed with the PD Quest software (Bio-Rad, Hercules, CA). Chemiluminescence was used to detect proteins after Western blot. Hybridization was preceded by an overnight incubation at room temperature with a blocking solution of 3% BSA in TBS Tween 0.15% (vol/vol; TBST). Incubation with primary anti-human albumin and anti-human a1AT antibodies (see the Validation of Antibody Specificity section) diluted 1:40,000 (vol/vol) was performed in 3% (wt/vol) BSA in TBST for 2 h at room temperature. The membrane then was washed with TBST four times, 15 min each, before the incubation with peroxidase-conjugated goat anti-rabbit IgG diluted 1:40,000 (vol/vol) BSA in TBST for 2 h at room temperature. The membrane then was washed four times, 15 min each, before developing the immune blot with the Super-Signal West Pico chemiluminescent Substrate (Pierce, Rockford, IL). The detection of fluorescence signals was acquired with Versa Doc 4000 (Bio-Rad).

### Validation of Antibody Specificity

For Western blot, the following two antibodies were used: (1) Polyclonal rabbit anti-human albumin (A0001; Dako, Glostrup, Denmark) and (2) polyclonal rabbit anti-human a1AT (A0012; Dako). The second antibody was a peroxidase-conjugated goat anti-rabbit IgG from Vector Laboratories (Burlingame, CA). Anti-actin antibodies from the same antibody was a peroxidase-conjugated goat anti-rabbit IgG from Vector Laboratories (Burlingame, CA). Anti-actin antibodies from the same

### Tryptic Digestion and Matrix-Assisted Laser Desorption/Ionization Time of Flight Protein Identifications

Protein spots were excised from 2D PAGE gel, and cysteines were reduced with dithiothreitol and alkylated with iodoacetamide. After derivatization, the samples were incubated at 37°C for 16 to 18 h with 20 μl of a buffer solution, 50 mM ammonium bicarbonate, that contained 10 ng/μl sequence-grade trypsin (Sigma, St. Louis, MO). The reaction was stopped by adding a final concentration of 0.1% trifluoroacetic acid and by freezing gel plugs at −20°C. After digestion, samples were used for protein identification by mass fingerprint matrix-assisted laser desorption/ionization time of flight (MALDI-TOF) analysis.

For the mass spectrometric analysis, the tryptic digests were extracted by ZipTip C18 reverse phase according to the manufacturer (Millipore, Bedford, CA). Elution and spotting were obtained with a 50% acetonitrile/ water solution saturated with α-cyano-4-hydroxycinnamic acid (Sigma) and deposited onto individual spots on the target MTP Ground Steel 384 (Bruker Daltonics, Billerica, CA). MALDI-TOF spectra were obtained by Reflex IV time-of-flight instrument (Bruker Daltonics) equipped with a 337-nm nitrogen laser. Mass spectra were acquired in reflectron-positive ion mode at a voltage of 20.00, 16.80, and 8.90 kV for ion source 1, 2, and lens, respectively, and 23.00 kV for Reflection in the 700- to 4500-m/z range. A 50-pmol/μl peptide-mix solution of angiotensin I (1296.68 Da), ACTH 18-39 (2465.19 Da), bradykinin (1060.57 Da), renin substrate (1758.93 Da), and Glu-Fibrinopep (1570.67 Da) was used as external calibration standard. Monoisotopic peaks were selected for peptide fingerprinting using the FlexAnalysis 2.0 software (Bruker Daltonics), and BioTools Version 2.2 was used to submit monoisotopic peptide masses for protein identification. Mass spectrometry (MS) data inference was compared with the National Center for Biotechnology Information nonredundant database data by the MASCOT software (Matrix Science, http://www.matrixscience.com). Protein identification was considered significant at P < 0.05 with respect to a random matching event. The query was restricted to human proteins, the maximum tolerance for masses was 100 ppm after an internal calibration using autolytic trypsin product, at most one miss-cleavage for tryptic peptides was allowed, and the modifications accepted were carboxymethylation of cysteines and possible oxidation of methionines. In accordance with the Proposed Publication Guidelines for the Analysis and Documentation of Peptide and Protein Identifications (Paris consensus), we included in Table 2 the complete figures of merit regarding the MS data. In particular, these include (1) matched peptides (the number of tryptic peptides actually aligned on the identified protein sequence), (2) sequence coverage (the percentage of the identified protein sequence covered by the MS data), and (3) identification score (a probabilistic measure of the goodness of the MS protein identification). All of the included protein identifications were significant at P < 0.05.

### Zymograms

Human serum albumin (Baxter) was reduced in 8 M urea, 65 mM dithioerythritol, 100 mM Tris(hydroxymethyl)aminomethane (TRIS), and 0.1 mM EDTA for 1.5 h at room temperature; 260 mM iodoacetamide then was added for 1 h at room temperature. Overalkylation was stopped by addition of 260 mM of dithioerythritol. Albumin finally was dialyzed three times against 10 L of 10 mM TRIS-HCl (pH 8.8).

Albumin zymography was performed on 8 to 16% polyacrylamide gels (160 × 140 × 0.75 mm) that contained 2 mg/ml carboxymethylated albumin under Laemmli conditions (22). After electrophoresis (12 mA, constant current at 12°C), gels were washed twice in 250 ml of 2.5% Triton X-100 (30 min each at 4°C) and incubated in 50 mM TRIS-HCl and 5 mM CaCl₂ (pH 7.8; 48 h, 37°C). Gels were fixed in 40% ethanol and 10% acetic acid and stained with 0.2% Coomassie brilliant blue G-250 (40% ethanol, 10% acetic acid) and destained appropriately (40% ethanol, 10% acetic acid). Gelatinase activity was evident as cleared regions.

### Results

Fresh urine was obtained from normal children and adults and from patients with nephrotic syndrome at the start of proteinuria. The study cohorts consisted of two groups of children with pri-
mary nephrotic syndrome and different response to steroids and an additional group that consisted mainly of adults with MGN. After the therapeutic approach, nephrotic children were subdi-
vided in responsive and resistant to steroids, and in case of de-
pendence or resistance, renal histology was evaluated to exclude
other primary and secondary pathologies.

Normal urine from adults and children was analyzed re-
petitively by a combination of 2D electrophoresis and MS,
and the same pattern was always observed. Owing to the
very low protein level in normal urine, gels were overloaded
with 2 ml of normal urine (containing 200 μg of total pro-
tein); a major band of uromodulin (acid protein with 80 kD
molecular weight) was evident, together with a few other
acid proteins between 50 and 20 kD and a low amount of
intact albumin (Figure 1, A and B). Therefore, the constant
pattern of normal urine in both children and adults was
characterized by the absence of albumin fragments. The com-
position of nephrotic urine was remarkably different. Nu-
merous fragments of albumin and of α1-antitrypsin (α1AT) in fact were
detected in urine of all patients with a few patterns common

Table 2. Urinary and plasma proteins from patients with nephrotic syndrome were characterized by two-
dimensional electrophoresis-Western blot and by MALDI-TOF

<table>
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<tr>
<th>Spot</th>
<th>Source</th>
<th>Identification Technique</th>
<th>Protein Name</th>
<th>NCBI</th>
<th>SC%</th>
<th>Score</th>
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<td>21</td>
<td>88</td>
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<td>84</td>
</tr>
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<td>305</td>
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<td>78</td>
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*A total of 72 spots were identified as fragments of albumin and/or of α1-antitrypsin (α1AT). Most (n = 72) were recovered from urine, and only some (n = 21) were also detected in plasma. According to the Proposed Publication Guidelines for the Analysis and Documentation of Peptide and Protein Identifications (Paris consensus), we included in this table the complete figures of merit regarding the mass spectrometry (MS) data. In particular, these include sequence coverage (SC%), indicating the percentage of the identified protein sequence covered by the MS data, and identification score (Score), indicating a probabilistic measure of the goodness of the MS protein identification. All included protein identifications were significant at P < 0.05. MALDI-TOF, matrix-assisted laser desorption/ionization time of flight; NCBI, National Center for Biotechnology Information; WB, Western blot.*
Intact albumin was the major protein in nephrotic urine. In parallel, numerous albumin fragments that covered the entire range of molecular weight from 64 to 20 kD were detected. An example is given in Figure 1E (Western blot with anti-albumin antibodies) and shows the pattern of a morning fresh urine sample that was obtained from a child with steroid resistance and FSGS; by comparison, a urine sample that was obtained from a child with steroid-sensitive nephrotic syndrome is...
shown in Figure 1, G and H. In both cases, the same amount of protein analyzed in normal urine (1 μl, containing 200 μg of protein) was loaded on gels. The presence of albumin fragments was confirmed by Western blot analysis of several urine samples from patients both with steroid-resistant FSGS (Figures 1, D through F, and 2, tracks 10 through 16) and with steroid-sensitive MCN (Figures 1, G and H, and 2, tracks 4 through 9 and tracks 17 through 20). Figure 2 shows the pattern of urinary albumin in both reduced (+β mercaptoethanol) and nonreduced (−β mercaptoethanol) conditions, indicating that some albumin fragments react with each other and form aggregates through disulphide bridges that are destroyed by β mercaptoethanol. Finally, albumin fragments also were detected with some differences in patients with MGN (Figure 1, L and M). These patients lacked, in fact, the isoforms with the lowest molecular weight. A comprehensive characterization of the differences between FSGS/MCN and MGN by dual labeling is in progress and will be presented in the near future.

Overall, 63 fragments of albumin were detected clearly in nephrotic urine (Figure 3); they were recovered from the gel and analyzed by MS that unequivocally confirmed their identity. The molecular weight of albumin fragments was calculated both from the human serum albumin (HSA) sequence that was obtained from the mass fingerprint experiments and from the SDS-PAGE. Both molecular weight values showed a good correlation, demonstrating that most of the HSA sequence was investigated in the MALDI-TOF-MS experiments. Moreover, few protein spots (Figure 4) showed consistently a peculiar distribution of the fingerprint peptide pattern that contained both the N-terminal region and the C-terminal domain with a complete lack of any MS signals for the internal sequence region. The reconstructed joint domains were calculated from mass fingerprint data that defined the HSA sequence regions that consistently were matched by experimentally measured tryptic peptides. In addition, the molecular weights of albumin spots estimated from 2D PAGE were not compatible with an albumin isoform that spanned the complete continuous sequence. Such tryptic peptide distribution could not be explained directly by the presence of the 16 disulfide bridges that were reported for the HSA structure given the strong reducing conditions that were used in sample preparation and separation. Therefore, we might speculate on the presence of covalent bound intramolecular cross-links that would be resistant to the strong denaturing conditions that were used in protein separation.

To exclude that the pattern of albumin fragmentation that was observed in nephrotic urine was due, in fact, to proteases in the tubule and/or in urine, we repeated the same analysis with Western blot in the companion plasma, where a large part of fragmentation products were recognized. The results are shown in Figure 5 (a 2D analysis and Western blot of plasma
albumin in one patient per group) and are summarized in Table 2. Twenty-one of the spots that were characterized as albumin/α1AT in urine also were present in plasma even at a lower concentration, suggesting preferential excretion of low molecular components. However, a lot of low molecular weight isoforms that were observed in urine were not present in plasma (see Table 2), suggesting an in situ fragmentation process that implies the presence of proteases in urine. In general, the components that were detected only in urine presented a low molecular weight and migrated to an area of gels indicated by a square in gels of Figure 1. On this basis, the presence of urinary proteases was postulated and experiments were planned to demonstrate their existence (see the Urinary Proteases Specific for Albumin section).

α1AT

α1AT is practically absent in normal urine (Figure 1, A and C). Intact and fragmented α1AT instead is well evident in urine of all different cohorts of patients (Figure 1, F, I, and N). MS confirmed identity of α1AT (Table 2). As for albumin also for α1AT, experiments with Western blot were carried out in plasma, where the presence of a part of α1AT fragments could be confirmed (Figure 5). Again, comparison of plasma and urine α1AT composition showed that a few isoforms with low molecular weight are produced in situ as an effect of proteases.

Urinary Proteases Specific for Albumin

To explain the formation of albumin fragments in urine, we performed a zymogram with albumin as substrate that indicated the presence of specific proteases. Comparison of normal and nephrotic urine demonstrated the presence of two major bands with protease activity that were upregulated in nephrotic patients (Figure 6). This finding indirectly supports the existence of a specific mechanism for in situ digestion of albumin in urine.

Discussion

This study was aimed at defining protein composition of urine and plasma in nephrotic patients and at reevaluating relevant structural aspects of major proteins such as albumin. For this purpose, we took advantage of the progress that has been made in the vast area of proteomics. Different clinical and pathologic conditions that are determined by different mechanisms such as primary nephrotic syndrome in children (permeability factors?) and MGN (immune complex) (23,24) were considered. The following relevant findings reveal unexpected
elements that suggest pathogenic implications and warrant wide discussion. The first regards the composition of nephrotic urine that practically contains only intact and fragmentation products of albumin and of $\alpha_1$AT. With the exception of a few low molecular weight urinary isoforms, some albumin and $\alpha_1$AT fragments also were detected in plasma, suggesting that fragmentation takes place at this site. Few isoforms instead are produced in urine, and we could demonstrate the presence in urine of specific proteases. A parallel aspect is how fragmentation influences renal handling of both proteins, which is the reason that nephrotic urine is enriched in fragments. The final point is about albumin adducts and their significance.

It is relevant that most of the proteins that are detected in urine of children with nephrotic syndrome correspond to intact albumin and $\alpha_1$AT or their fragments as it clearly emerges from the comparison of the 2D gels of the same urinary sample stained by Coomassie and Western blot reported in Figure 1. MS confirmed this conclusion.

To our knowledge, the finding of protein fragments in serum and urine of patients with renal diseases is new and suggests different fragmentation mechanisms. The high concentration of albumin fragments in nephrotic urine suggests a preferential urinary excretion and reflects the partial maintenance of size selectivity properties of the kidney. Albumin fragments account, in fact, for 25 to 30% of total albumin in urine of nephrotic patients, whereas their proportion in serum does not exceed 5%. Considering that in nephrotic syndrome at least 2 to 3 g of albumin is lost every day in urine and that up to 1 g (30%) is represented by fragments, it can be concluded that almost 30 g of fragments is cleared monthly from plasma, accounting for 30% of the albumin pool in a patient who weighs 30 kg. Our conclusion is that preferential urinary excretion of albumin fragments in nephrotic patients represents an efficient depurative mechanism that restores normal albumin composition in plasma, and the same considerations apply to $\alpha_1$AT. It is critical to this conclusion that also normal plasma contains albumin fragments, which suggests that in nephrotic syndrome the fragmentation process is amplified. This seems even more interesting for $\alpha_1$AT, which acts as a major circulating defense against exogenous proteolysis enzymes of viral or bacterial derivation and also controls the proteolysis response that is triggered by normal immunologic process. It is a suicide protein in the sense that it attracts proteases at a bait region and is degraded after binding by the proteosome (25,26). To our knowledge, nothing like this has been described before in human pathology, and

\begin{figure}
\centering
\includegraphics[width=\textwidth]{image}
\caption{2D electrophoresis and Western blot with anti-albumin (A, C, E, and G) and anti-$\alpha_1$AT (B, D, F, and H) polyclonal antibodies of plasma from normal volunteers (A and B) and from patients with nephrotic syndrome caused by different renal pathologies such as FSGS (C and D), MCN (E and F), and MGN (G and H).}
\end{figure}
experimental and in vitro models are required to investigate this aspect further. It seems relevant that inactivation of α1AT by leukocytes has been reported extensively in the literature, this mechanism involving the production of free radicals from activated neutrophils and activation of specific proteases (27–29). This observation reinforces the hypothesis of a role of free radicals in nephrotic syndrome that has been proposed in the past but never received experimental support.

A final but not secondary point is the description of albumin adducts. They were characterized by MS as peptides that harbor both the COOH and NH2 terminal parts of the protein. Such evidence cannot be explained simply by the presence of disulfide bridges or the adventitious lack of MS signals given the 2D electrophoresis separation conditions and the calculated molecular weight. These data might be suggestive of the presence of a covalent chemical adduct that links the two otherwise separated parts of albumin (30). Such cross-links have been described as end products of Amadori compounds that are generated from the glycation of lysine residues. Nevertheless, the large number of chemical structures that are formed during nonglycated glycation does not allow us to draw any conclusion on the new molecules described here (31). The presence of albumin adducts in FSGS is new and obviously will receive further attention in the near future for a definite chemical characterization.

Conclusion

Albumin and α1AT fragments represent the major proteins in urine of patients with nephrotic syndrome, where at least 50 isoforms of these proteins were characterized by proteomics analysis. Most of these isoforms derive from plasma, but a few were formed in situ by specific proteolysis. Albumin adducts that harbor both the COOH and NH2 terminal parts of the protein also were detected, suggesting the formation of covalent chemical adducts. These new findings reveal unexpected structural and functional aspects of proteinuria that may play a key role in pathogenesis.

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