Primary Care Approach to Proteinuria

Amir Said Alizadeh Naderi, MD, and Robert F. Reilly, MD

Proteinuria is a common finding in primary care practice. Most adolescents who are diagnosed with proteinuria through screening urinalysis do not have renal disease, and the proteinuria will usually resolve on repeat testing. In contrast, proteinuria is suggestive of kidney disease in patients with diabetes mellitus, hypertension, primary renal disease, or other systemic illnesses. Quantification of proteinuria can be used longitudinally to monitor therapeutic effects of treatment of the underlying disease. Given the multitude of clinical settings in which proteinuria can occur, we suggest an algorithm that may help clinicians differentiate between benign and serious etiologies of proteinuria. (J Am Board Fam Med 2008;21:569–74.)

During a time when uroscopy was the only diagnostic tool for diagnosing renal diseases, Hippocrates described the association between “bubbles on the surface of the urine” and kidney disease. In the seventeenth century Frederick Dekkers described how urine samples became caseous after exposure to heat and acetic acid, but it took another 200 years until Richard Bright associated proteinuria with kidney disease. Nowadays, the presence of proteinuria is routinely and easily assessed in clinical practice. This article provides an overview of the most common causes of proteinuria in adults, describes ways to assess proteinuria, and offers an algorithmic approach to differentiate benign from serious etiologies of proteinuria.

Proteinuria

Proteinuria is a common clinical finding in primary care practice. Healthy kidneys excrete less than 150 mg of protein/day, of which approximately 20 mg is albumin. Tamm-Horsfall mucoproteins, which is secreted by tubular cells and immunoglobulins, are responsible for the remaining part of physiologic proteinuria. Microalbuminuria is defined as daily excretion of 30 to 300 mg of albumin/day and is an early and sensitive marker of diabetic nephropathy. Microalbuminuria is associated with an increased risk of cardiovascular disease in patients with and without diabetes and/or hypertension. Proteinuria has been shown to be an independent risk factor for the progression of kidney disease rather than simply being a marker for glomerular dysfunction. The daily excretion of more than 3.5 g of protein is called nephrotic range proteinuria. The discussion of pathophysiologic mechanisms of proteinuria is beyond the scope of this article (for more information, read a recent review article). Table 1 summarizes the different types of proteinuria.

Classification of Proteinuria

Proteinuria can be classified into 3 broad categories: (1) overflow proteinuria, (2) tubular proteinuria, and (3) glomerular proteinuria.

Overflow proteinuria

Low-molecular weight proteins filtered by the glomerulus are almost entirely reabsorbed in the proximal tubule. During states of increased low-molecular weight protein production and subsequent filtration, the amount of filtered protein exceeds tubular reabsorptive capacity, leading to proteinuria. In multiple myeloma, excessive amounts of immunoglobulin light chains are produced, which can be detected in the urine because of the limited reabsorptive capacity of the proximal tubule.

Tubular proteinuria

Tubulointerstitial diseases (eg, Fanconi syndrome) can lead to decreased reabsorptive capacity of the proximal tubule and up to 2 g of proteinuria per day.
because of impaired tubular absorption of filtered albumin, as well as loss of tubular proteins (β2 microglobulin).

**Glomerular proteinuria**

Glomerular proteinuria is a sensitive marker for glomerular disease. It develops as a result of increased filtration of macromolecules (eg, albumin) across the glomerular capillary wall (Table 2).

### Quantification of Proteinuria

Proteinuria can be quantified by different means. Urine dipstick testing, spot urine specimen analysis, timed urine collection, and the sulfosalicylic acid test are the most commonly employed tests.

**Dipstick testing**

Urinary dipstick testing is routinely performed in clinical practice. It is an inexpensive test that is used for screening of different disorders, including urinary, gastrointestinal, metabolic, and hematologic diseases. Besides quantification of proteinuria, urinary dipstick testing provides valuable information about other parameters, including urinary blood, glucose, ketones, pH, specific gravity, leukocyte esterase, and nitrite.

The standard dipstick measures albumin concentration in the urine via a colorimetric reaction between albumin and tetrabromophenol blue, causing different color shading depending on the albumin concentration in the sample. If the urine is dilute (low specific gravity), urine albumin concentration is decreased and is occasionally not detected by dipstick testing. The most conservative approach would be to repeat the value and, if positive, to obtain a protein/creatinine or albumin-to-creatinine ratio. Importantly, as dipsticks only measure albuminuria, other proteins like renally excreted immunoglobulin light chains in multiple myeloma are not detected. A positive urinary dipstick is a useful screening test for proteinuria. Its limitations are quantification of proteinuria and its inadequacy to detect immunoglobulin light chains.

**Spot urine analysis**

Spot urine analysis for protein-to-creatinine ratio (UPCR) and albumin-to-creatinine ratio (UACR) are currently preferred to the 24-hour urine collection for quantification of daily proteinuria given the greater convenience of spot urine sampling. Measurement of UACR was shown to be as good as a 24-hour urine collection for the quantification of proteinuria (strength of evidence A).16-18 The UACR is used to assess for microalbuminuria, whereas the UPCR can be used when proteinuria of more than 300 mg/day is suspected. First morning specimens are preferable.

The National Kidney Foundation recommends annual screening for proteinuria in diabetics using the UACR in a spot urine sample. Because the amount of spot urine proteinuria might fluctuate, a positive UACR should be confirmed (in the absence of a urinary tract infection) with 2 additional first-void specimens collected over the next 3 to 6 months (strength of evidence B).19 It is worth mentioning that the UACR is a more sensitive screening test than the UPCR. UPCR is less expensive for a patient with established proteinuria and can be used longitudinally to evaluate therapeutic efforts to reduce proteinuria.

Clinicians should be aware that the accuracy of estimating proteinuria from a random urine specimen is diminished if creatinine excretion is substantially different from 1 g/day. Thus, the albumin-to-creatinine ratio will underestimate protein excretion in a muscular man with a high rate of creatinine excretion and will overestimate urinary protein excretion in a cachectic patient with low muscle mass and low creatinine excretion.

**24-hour urine collection**

Twenty-four-hour urine collection is the most accurate test for quantification of proteinuria and is also used for qualitative urinalysis.20,21 However, the test becomes inaccurate when difficulties with urine collection are encountered. To minimize collection and handling errors, patients should receive detailed instructions about 24-hour urine collection.

### Table 1. Terminology Used to Describe Proteinuria

<table>
<thead>
<tr>
<th></th>
<th>Albuminuria</th>
<th>Proteinuria</th>
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<tbody>
<tr>
<td>Physiologic range proteinuria</td>
<td>&lt;20 mg/day (15 μg/min)</td>
<td>&lt;150 mg/day</td>
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<tr>
<td>Microalbuminuria</td>
<td>30–300 mg/day (20–200 μg/min)</td>
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<tr>
<td>Macroalbuminuria</td>
<td>&gt;300 mg/day (&gt;200 μg/min)</td>
<td>&gt;3.5 g/day</td>
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<td>Nephrotic range proteinuria</td>
<td></td>
<td></td>
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<tr>
<td>Cause</td>
<td>Renal Manifestation</td>
<td>Clinical Presentation</td>
</tr>
<tr>
<td>--------------------------------------------</td>
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<tr>
<td>Diabetes mellitus type 1/type 2</td>
<td>Ranging from microalbuminuria to nephrotic range proteinuria; most common etiology of end-stage renal disease in the US</td>
<td>Polyuria, polydipsia, nocturia, weight loss</td>
</tr>
<tr>
<td>Hypertension</td>
<td>Protein excretion is usually less than 1 g/day¹¹</td>
<td>Often asymptomatic (excluding hypertensive crisis)</td>
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<tr>
<td>Pre-eclampsia</td>
<td>Nephrotic syndrome, a decreased GFR compared to healthy pregnant women¹⁴</td>
<td>Edema</td>
</tr>
<tr>
<td>Multiple myeloma</td>
<td>SCr is increased in 50% of patients¹³; urine protein is not detected on urine dipstick; sulfosalicylic acid test and UPCR detect proteinuria</td>
<td>Pallor, radiculopathy, peripheral neuropathy, bone pain in the back or chest is present in 60% of patients²⁷</td>
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<td>Light chain deposition disease</td>
<td>Most commonly associated with multiple myeloma; in rare cases it occurs with lymphoma</td>
<td>Related to renal, cardiac, or hepatic involvement</td>
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<td>Primary amyloidosis</td>
<td>Nephrotic syndrome with or without renal insufficiency</td>
<td>Peripheral neuropathy, macroglossia, hepatomegaly</td>
</tr>
<tr>
<td>HIV nephropathy</td>
<td>Nephritic syndrome, more common in African-American patients; nephritic syndrome</td>
<td>HIV positive, normal to big kidney size</td>
</tr>
<tr>
<td>Immunoglobulin A nephropathy</td>
<td>Nephritic syndrome</td>
<td>Hematuria</td>
</tr>
<tr>
<td>Minimal change disease and membranous glomerulonephritis</td>
<td>Nephritic syndrome</td>
<td>Edema, anasarca, foamy urine</td>
</tr>
<tr>
<td>Streptococcal glomerulonephritis</td>
<td>Ranging from microscopic hematuria to acute nephritic syndrome</td>
<td>Recurrent episodes of hematuria 1 to 3 days after URI</td>
</tr>
<tr>
<td>Rare forms of glomerulonephritis, other infectious or systemic diseases</td>
<td>Depending on underlying disease patients may have proteinuria with or without an active urinary sediment (red blood cell casts, dysmorphic erythrocytes)</td>
<td>Depending on underlying disease</td>
</tr>
</tbody>
</table>

*Nephrotic syndrome = more than 3.5 g proteinuria/day, hypoalbuminemia, edema, hyperlipidemia, lipiduria.

¹Nephritic syndrome = red to brown urine, proteinuria (usually <3.5 g/day, edema, hypertension, and acute renal failure).

²Active urine sediment = red blood cell casts or dysmorphic erythrocytes.

RBC, red blood cell; GFR, glomerular filtration rate; ANA, antinuclear antibodies; HIV, human immunodeficiency virus; LVH, left ventricular hypertrophy; SCr, serum creatinine; URI, upper respiratory tract infection; VDRL, Venereal Disease Research Laboratory.
Box 1. Patient Instructions for 24-hr Urine Collection

On the first day of collection, urinate into the toilet when you get up in the morning. Afterward, collect all urine in a special container for the next 24 hours. On the second day, urinate into the container when you get up in the morning.
- Cap the container.
- Keep the sample in the refrigerator during the collection period.
- Label the container with your name, the date, and the time of completion and return it to your healthcare provider.

A timed urine collection is often not collected properly. To determine the completeness of a 24-hour urine collection, the total daily urine creatinine is often used as a standard. In patients younger than 50 years, normal daily urinary creatinine range is 15 to 20 mg/kg lean body weight for women and 20 to 25 mg/kg lean body weight for men. In older people, the production of daily creatinine is reduced, primarily because of decreased muscle mass. In the latter case, calculating the ratio of creatinine clearance measured by 24-hour collection to the estimated creatinine clearance from a serum sample can help ensure proper collection. A ratio of 0.9:1.1 suggests a complete 24-hour urine collection because both estimated and measured creatinine clearances are roughly equal. Although in many cases the timed urine collection is incomplete, measurement of UACR and UPCR from this urine collection is still helpful and may be more accurate than a spot urine specimen.

Sulfosalicylic Acid Test
The sulfosalicylic acid test detects all types of urinary proteins. This characteristic makes the sulfosalicylic acid test useful in the diagnosis of multiple myeloma, which is characterized by urinary excretion of immunoglobulin light chains (these are not detected by urine dipstick testing). A sulfosalicylic acid test should not be used as a screening test because it cannot detect small quantities of monoclonal free light chains. In the latter case, examination of 24-hour urine samples using electrophoresis and immunofixation is more sensitive. Overall, the sulfosalicylic acid test is no longer routinely used in the assessment of proteinuria.

Box 2. 24-hr Urine Collection in the Diagnosis of Orthostatic Proteinuria

The test requires a 24-hour urine collection in 2 different jugs, which can be labeled as “day time urine” and “night time urine.” During the collection time the urine needs to be stored in a refrigerator. The first morning urine is discarded. For the rest of the day all urine is collected in the daytime urine jug. In the evening, the patient should lie down for approximately 2 hours before going to sleep. Just before sleeping, the patient should void for the last time during the day and add this urine to the daytime urine jug. By lying down before the nighttime collection contamination of the nighttime urine with urine formed during the day when the patient was upright is avoided. The following morning (8 hours after the last night time collection) the patient shall collect the first morning urine in the nighttime urine jug. Both urine jugs are taken to the laboratory thereafter.

Transient and Orthostatic Proteinuria
The first step in the assessment of proteinuria is to differentiate between benign and serious causes of proteinuria. In adolescents, proteinuria can be induced by strenuous exercise, fever, and extreme cold. In all of these circumstances, proteinuria resolves spontaneously after the cessation of the causal factor, so an extensive work-up is not needed. Up to 5% of adolescents may have orthostatic or postural proteinuria. This benign condition is characterized by an increase in protein excretion in the upright position, which normalizes when the patient is recumbent. As a rule of thumb, patients with orthostatic proteinuria excrete less than 1 g of protein over 24 hours in the upright position; the rate of protein excretion completely normalizes in a supine position (<50 mg per 8 hours). The long-term prognosis for adolescents with orthostatic proteinuria is excellent, even after decades of follow-up. Orthostatic proteinuria is uncommon over the age of 30. Young adults with proteinuria should be evaluated by a nephrologist if proteinuria persists on repeat testing, especially if they develop an increase in serum creatinine concentration or when there are associated abnormalities in the urinary sediment.
The gold standard for assessment of orthostatic proteinuria is the collection of a 24-hour urine sample.

Alternatively, a normal UPCR of <0.15 mg/mg on a first morning spot urine specimen and an elevated total protein to creatinine ratioof
more than 0.15 mg/mg on a second upright spot urine specimen is indicative of orthostatic proteinuria.

Conclusion

Proteinuria is a common diagnosis in primary care practice. Using an algorithm as suggested in this paper (Figure 1) can help the clinician to distinguish between benign and serious causes of proteinuria. Physicians need to be aware that although patients with diabetes and/or hypertension commonly develop proteinuria in the setting of their underlying disease, proteinuria might also develop as a consequence of a second disease. Thus, all adults should also have a urine sediment examination and urine protein electrophoresis to exclude Bence Jones proteinuria. The presence of unexplained proteinuria or active urine sediment suggestive of a nephritic syndrome requires nephrology consultation.

References

2. Bright R. Cases and observations, illustrative of renal disease accompanied with the secretion of albuminous urine. London Medical Gazette 1835;72–74.