Differentiation of Glomerular from Non-Glomerular Hematuria by Three Different Methods of Microscopic Examinations of Erythrocytes in Urine

A. Abolfathi, A. Hosaininasab, H. Argani

Abstract
Background: Morphological examinations of urinary erythrocytes can be of diagnostic value in initial evaluation of hematuria. Dysmorphic urinary red blood cells are known to indicate a glomerular origin of bleeding. We examined the clinical usefulness of this test in a population complained of hematuria by use of three different methods: light microscopy, phase contrast microscopy, and Wright staining and compared their sensitivity and specificity.

Methods: The study included 169 patients with hematuria (89 glomerular and 80 non-glomerular). The urine specimens were collected before invasive procedures such as biopsy and cystoscopy. In each urine sample, 100 urinary erythrocytes were examined. Statistical analysis was performed using Student's t test, correlation coefficient, and \( \chi^2 \). Reliability parameters including sensitivity, specificity and predictive values of negative and positive tests were also evaluated.

Results: Dysmorphic red cells were recorded as acanthocytes, doughnut-like cells, yeast like cells with more than one blebs and ghost forms. Isomorphic erythrocytes had uniform size and shape. Significant difference was found in the number of urinary dysmorphic red cells between the two groups of patients. Statistical analysis showed that by using percentage of glomerular type erythrocytes and setting the cut-off at 20-25%, the specificity for three procedures was almost the same (≈ 97.5%). But sensitivity for light microscopy, phase contrast microscopy, and Wright staining was in different ranges as 70.7%, 89.8%, and 86.5% respectively.

Conclusion: It was concluded that with some limitations, these simple, non-invasive techniques were useful in identifying the source of bleeding in the work up of hematuria by considering that sensitivity of the methods were in the order of phase contrast microscopy, Wright staining, and light microscopy.


Keywords • Red blood cell • morphology • hematuria • microscopy
Introduction

Evaluation of the cause of hematuria is a common diagnostic problem in clinical practice. Standard invasive investigations of these patients are expensive and potentially harmful.

Morphological examinations of urinary erythrocytes were suggested by Fairly and Birch, in an attempt to find out methods to be cost effective and low risk for initial evaluation of hematuria. In various studies, morphological examination of urinary erythrocytes by phase contrast microscopy, electron microscopy, or light microscopy with and without staining, has established differences between erythrocytes from glomerular and non-glomerular sources.

Erythrocytes of glomerular origin are usually small, and fragmented with variations in size and shape (dysmorphic). Erythrocytes of non-glomerular origin are largely uniform, resembling normal peripheral red cells (isomorphic), although a variable number may lose their hemoglobin (ghost cells) and few may be crenated.

Hematology autoanalyzers have been also used to provide a more objective assessment of urinary erythrocyte morphology to differentiate glomerular from non-glomerular hematuria. Because sensitivity and specificity of various methods used in different studies are contradictory, we aimed to compare the three different microscopic methods: light microscopy (LM) with and without Wright staining and phase contrast microscopy (PCM) to find out the most sensitive and specific method for differentiation of glomerular from non-glomerular hematuria.

Materials and Methods

Patients’ selection and specimens’ collection

The study included 169 patients referred to nephrology and urology wards of Imam Khomeini hospital affiliated to Tabriz University of Medical Sciences since October 2001 to September 2003 with the complaint of hematuria. Glomerular and non-glomerular renal disease were detected by intravenous pyelography (IVP), ultrasonography, cystoscopy, or renal biopsy. The urine microscopy was performed by two observers who were blinded to the final diagnosis.

The urine specimens were collected before invasive procedures such as biopsy and cystoscopy. Thirty ml of fresh morning midstream urine was obtained. Urine samples were analyzed by dipstick for albumin, pH, sugar, and blood. The specific gravity of urine samples was controlled by ERMA refractometer. The specimens with pH more than 8 and less than 5 and specific gravity less than 1010 were excluded. Serum blood urea nitrogen (BUN), creatinine, and glucose and complete blood count (CBC) of patients were also checked. The patients with abnormal morphology of red blood cells in peripheral blood and MCV less than 75 fl or end stage renal failure were excluded.

RBC morphology study

Ten ml of urine was centrifuged at 2000 rpm (400g) for 10 min and sediment was resuspended in 0.5 ml of supernatant. One drop of sediment was then examined by light microscopy (×400) and phase contrast microscopy (×400) by a Zeiss microscope with phase contrast attachment. In each sample of urine, 100 urinary erythrocytes were examined.

After adding one drop of 5% human serum albumin into urinary sediment and mixing it, a drop of suspension was placed on the glass slide and air-dried smear was prepared. The smears were then stained with 0.3g/100 Wright stain according to the method used by Chang. The slides were covered with cover glass. One hundred red cells were examined by high power (×1000) and checked for the percentage of dysmorphic, isomorphic, hypochromic and normochromic RBCs.

Statistical analysis was performed using Student’s t test, correlation coefficient, x², and cross tabulation using SPSS software (version 11-0). All the values are reported as means ± SD. Reliability parameters including sensitivity, specificity and predictive values of negative and positive tests were also evaluated.

Results

The diagnosis of the 169 patients with hematuria on the basis of the clinical picture and results of the appropriate investigations in radiology, cystoscopy, pathology and immunohistopathology were established. Of these 169 patients, hematuria was considered glomerular in 89 (53%) and non-glomerular in 80 (47%) patients. The mean age of the patients with glomerular hematuria was 35±17 and with non-glomerular hematuria was 46±20 years.

Statistical analysis (x² test) did not show significant difference in the number of red blood cells in the urine samples between the two groups of patients (p> 0.5).

Morphology of urinary erythrocytes as isomorphic and dysmorphic with different microscopic methods including W.S, PCM and LM were studied. Dysmorphic red cells were recorded as acanthocytes, doughnut–like cells, yeast like cells, spherical cells with more than one blebs and ghost forms of erythrocytes. In contrast to these, isomorphic erythrocytes had uniform size and shape (figures 1, 2). As noted in
Table 1 there was significant difference ($p<0.0001$) in the number of urinary dysmorphic red cells between the two groups of patients.

The two groups of patients were compared based on the presence of $\geq 20\%-25\%$ of dysmorphic erythrocytes in urine specimens and the results listed in Table 2. If the cut off point for dysmorphic erythrocytes be considered as $\geq 20\%$ of whole red cells count, 75 patients studied by light microscopy, 86 patients by phase contrast microscopy, and 84 patients by Wright staining out of the 89 patients will be diagnosed as having glomerular hematuria. In 80 patients with non-glomerular hematuria, 5
patients studied by light microscopy and 2 patients studied by phase contrast microscopy and Wright staining had 20% or more dysmorphic RBCs respectively. If the cut off point for dysmorphic RBCs be considered ≥ 25%, these figures for patients with glomerular hematuria will be 63, 80 and 77 respectively.

Table 1: Source of hematuria determined by conventional evaluation

<table>
<thead>
<tr>
<th>Disease</th>
<th>Number of patients</th>
</tr>
</thead>
<tbody>
<tr>
<td>Glomerular disease</td>
<td>89</td>
</tr>
<tr>
<td>Crescentic glomerulonephritis (CGN)</td>
<td>6</td>
</tr>
<tr>
<td>Membranoproliferative glomerulonephritis</td>
<td>24</td>
</tr>
<tr>
<td>(MPGN)</td>
<td></td>
</tr>
<tr>
<td>Mesangioproliferative glomerulonephritis</td>
<td>4</td>
</tr>
<tr>
<td>(MEGN)</td>
<td></td>
</tr>
<tr>
<td>Focal and segmental glomerulonephritis</td>
<td>15</td>
</tr>
<tr>
<td>(F&amp;SGS)</td>
<td></td>
</tr>
<tr>
<td>Minimal change nephrotic syndrome (MCNS)</td>
<td>2</td>
</tr>
<tr>
<td>Post streptococcal glomerulonephritis</td>
<td>16</td>
</tr>
<tr>
<td>Post infection glomerulonephriti (PIGN)</td>
<td>2</td>
</tr>
<tr>
<td>IgA nephropathy (IgAN)</td>
<td>2</td>
</tr>
<tr>
<td>Diabetic nephropathy (DN)</td>
<td>6</td>
</tr>
<tr>
<td>(AM)</td>
<td>1</td>
</tr>
<tr>
<td>Systemic lupus erythematos</td>
<td>9</td>
</tr>
<tr>
<td>Goodpasture’s disease (GP)</td>
<td>2</td>
</tr>
<tr>
<td>Urolological disease</td>
<td>80</td>
</tr>
<tr>
<td>Renal calculi</td>
<td>40</td>
</tr>
<tr>
<td>Bladder cancer</td>
<td>15</td>
</tr>
<tr>
<td>Urethral cancer</td>
<td>2</td>
</tr>
<tr>
<td>Prostate cancer</td>
<td>1</td>
</tr>
<tr>
<td>Benign prostate hypertrophy</td>
<td>10</td>
</tr>
<tr>
<td>Post-transurethral resection</td>
<td>6</td>
</tr>
</tbody>
</table>

Table 2: Comparison of mean value of dysmorphic urinary RBCs in two groups of G and NG patients by three methods.

<table>
<thead>
<tr>
<th></th>
<th>NG=80 Mean±SD</th>
<th>G=89 Mean±SD</th>
<th>P value</th>
</tr>
</thead>
<tbody>
<tr>
<td>LM</td>
<td>7.8±5.7</td>
<td>36.2±15.2</td>
<td>&lt;0.0001</td>
</tr>
<tr>
<td>PCM</td>
<td>8.3±6.6</td>
<td>48.7±18.4</td>
<td>&lt;0.0001</td>
</tr>
<tr>
<td>W.S</td>
<td>7.9±5.1</td>
<td>46.6±18.1</td>
<td>&lt;0.0001</td>
</tr>
</tbody>
</table>

To find the most sensitive method among the three methods, statistical analysis showed that both phase contrast microscopy and Wright staining were more sensitive than light microscopy to differentiate dysmorphic from isomorphic erythrocytes. There was no significant difference between phase contrast microscopy and Wright staining in diagnosis of dysmorphic RBCs. Correlation coefficients of 0.93, 0.94 and 0.99 were obtained respectively when phase contrast microscopy with light microscopy, Wright staining with light microscopy and phase contrast microscopy with Wright staining were compared.

**Discussion**

Hematuria is the most prevalent clinical finding of kidney and urinary tract disease. With respect to its origin it could be divided to glomerular and non-glomerular. The cause of red cell deformity which is used for differential diagnosis is still speculative. Different factors including, mechanical damage caused by passage of red blood cells through the glomerular basement membrane, and biochemical changes (e.g osmolality, pH, urinary protein), are assumed to be associated with dysmorphic transformation of urinary erythrocytes. Examination of urinary erythrocyte morphology has been proposed to differentiate glomerular from non-glomerular hematuria since 1979. After nearly 25 years of this approach, reported
diagnostic rates are variable. The morphological definition of a dysmorphic erythrocyte, and absolute number of dysmorphic erythrocytes necessary to make diagnosis of glomerular bleeding, are widely variable. The clinical use of examination is also controversial.

In attempt to suggest definite criteria for glomerular bleeding from non-glomerular in this study, two parameters were found to be more important: glomerular and non-glomerular shape of urinary erythrocytes including size and form and percentage. The first report of variation in size and shape of urinary red blood cells in more detail was described by Birch and Fairley, who correlated changes with clinical information. They recorded crenate and uniform as normal and anisocytes with abnormal shapes as dysmorphic. While Fassett and his colleagues, reported echinocytes as abnormal but Kohler and Eveline, described echinocytes and stomatocytes as normal red blood cells in urine. Tomati and his colleagues classified urinary erythrocytes into 10 shapes by differential interference microscopy (DIM). In their classification; acanthocytes, doughnut-like cells with blebs, and yeast-like cells were the most distinguished among dysmorphic cells. In recent studies the detection of acanthocytes, doughnut shaped RBCs, and G1 cells in urine are suggested as a useful tool for the diagnosis of glomerular bleeding. In addition to dysmorphic changes, urinary red blood cells from patients with glomerular disease were often hypochromic. So that color of red blood cells could have diagnostic value.

In the present study, we tried to make definition of dysmorphic more accurately. In our study only restricted shapes (acanthocytes, doughnut-like cells, yeast-like cells and smaller erythrocytes with deformity) were counted and other dysmorphic shapes were ignored. Our cut-off to differentiate glomerular from non-glomerular hematuria was 20-25% dysmorphic red blood cells in urine. Our cut-off point was similar to Dekemerchou, who diagnosed glomerular hematuria in the presence of 20% dysmorphic RBC. But other cut-off points that were defined for glomerular bleeding were variable from 10 to 80% dysmorphic RBC. This variance could be reflected from different definitions of dysmorphic RBC. According to our criteria the dysmorphic RBCs in urine of patients with non-glomerular hematuria were rare and so 20-25% cut-off was an accurate point to specify glomerular bleeding.

In contrast to Mittenyi and his colleagues, who reported a relatively low specificity (63.6%) and sensitivity (84%) of dysmorphism for the diagnosis of glomerular disease, the reported specificity (96-98%) and sensitivity (87-90%) in this study is in agreement with some other reports.

Although according to the above-mentioned criteria, we found that phase contrast microscopy and Wright staining methods were more sensitive than light microscopy, but it still can be useful in the presence of skilled cytopathologist or expert nephrologists to investigate morphology of urine erythrocytes properly.

Our study confirms that examination of erythrocytes in the urine with these methods is simple, inexpensive, non-invasive and repeatable that permits an accurate distinction between glomerular and non-glomerular hematuria. These methods do not lead to a definite diagnosis, but do enable the selection of the most appropriate test and thus avoid unnecessary, often invasive diagnostic procedures.

References


Differentiation of glomerular from non-glomerular hematuria by examinations of erythrocytes in urine