Spot urine protein/creatinine ratio—A quick and accurate method for diagnosis of pre-eclampsia

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ABSTRACT

Objective(s): To assess the diagnostic accuracy of spot urine protein-creatinine (P/C) ratio and its comparison with 24-hour urine proteinuria for predicting eclampsia. Method(s): Spot urine P/C ratio was determined in a mid-stream urine sample, and the 24-hour urine protein was measured. The correlation between the spot P/C ratio and 24-hour urine protein amount was done. Logistic regression analysis and ROC curve analysis have been used to analyse data. Result(s): There was a strong correlation between the spot P/C ratio and 24-hour urine protein excretion (pearson’s correlation coefficient r = 0.71; P < 0.0001). The optimal spot P/C ratio cutoff point was 0.25, for 300 mg/24 h of protein excretion, with sensitivity and specificity of 69% and 75% respectively. Conclusion(s): Spot urine P/C ratio is a quick and reliable tool which can be used as an alternative method for evaluation of proteinuria for diagnosis of pre-eclampsia.

Keywords: Urine Protein-Creatinine Ratio; 24 Hour Urine Proteinuria; Pre-Eclampsia; Cutoff Point

1. INTRODUCTION

Pre-eclampsia is a multisystem disorder complicating 5% - 10% of pregnancy [1]. Pre-eclampsia is a major cause of maternal and fetal morbidity and mortality in India. This is a pregnancy specific syndrome of reduced organ perfusion secondary to vasospasm and endothelial activation [2]. The minimum criteria for the diagnosis of pre-eclampsia are hypertension plus proteinuria. Proteinuria is an important sign of pre-eclampsia. Significant proteinuria is described as 300 mg or more of urine protein per 24 hours period. The gold standard for diagnosis of significant proteinuria is 24 hours urine collections [3]. However the collection of urine for 24 hours is time-consuming and needs strong co-operation; requires hospitalization, inconvenient, costly and inaccurate due to incomplete collection and poor compliance. This can delay diagnosis and management of pre-eclampsia.

A more rapid test capable of accurately predicting end results of 24 hours urine is required. An alternative method for quantitative evaluation of proteinuria is the measurement of protein-creatinine ratio in spot urine sample, which avoids the influence of variation in urinary solute concentration and provides a more convenient and rapid method to assess protein excretion [4]. A rapid and accurate test may provide efficient in-patient and outpatient monitoring of proteinuria and may shorten the duration of hospitalization. This study was conducted to assess the diagnostic accuracy and determine an optimum cut-off point of spot urine protein-creatinine ratio for the prediction of significant proteinuria in patients with pre-eclampsia.

2. MATERIAL AND METHODS

The present study was planned as a prospective study which include 140 pregnant women of ≥20 weeks of gestation period and ≥140/90 mmHg Blood pressure, admitted in Upper India Sugar Exchange, Dept. of Gynaecology, G.S.V.M. Medical College, Kanpur over a period of 18 months during January, 2009 to June, 2010.

A detailed history with thorough clinical examination and routine investigations, liver function test, renal function test was done to select women. Women with chronic hypertension, intrinsic renal disease, liver disease, co-existing urinary tract infection, Gestational diabetes mellitus, Inadequate specimen, Heavy exercise—more than 1 hour, Bacteremia, Strict bed rest longer than 24 hours were excluded from the study.
All women included in the study irrespective of the severity of disease were asked to provide spot mid stream urine sample and collect all the urine subsequently for 24 hours period. The urine protein-creatinine ratio was determined on spot urine specimen. Total urine protein quantification was done by well established, Biuret calorimetric assay and urine-creatinine estimation was done by modified Jaffe’s method. The urine P/C ratio was obtained by dividing the total urine protein concentration by the urine creatinine concentration.

The relationship between protein-creatinine ratio and 24-hour protein excretion was assessed by Pearson correlation coefficient. Sensitivity, specificity, positive predictive value and negative predictive value of random urine protein creatinine ratio at various cut offs for prediction of significant proteinuria were estimated with a 95% confidence interval (95% CI), using the results of 24-hour urine collection as gold standard.

The received operating characteristic (ROC) curve analysis was used to determine the best discriminator values of the spot urine protein-creatinine ratios for preeclampsia (proteinuria ≥300 mg/24 hr). A P value of 0.05 was considered to be significant. The SPSS software was used for the statistical analyses.

3. RESULTS

In this study a total of 126 pregnant women with preeclampsia were selected according to the inclusion and exclusion criteria. 14 were excluded from the study, 7 were delivered before completion of 24 hour urine collection, 3 were diagnosed with chronic hypertension and 4 were having urinary tract infection.

**Table 1** shows age group 21 to 30 years contributed the highest percentage (87.92%). Majority of subjects were nulliparous (46.03%), followed by primipara (19.04%). Majority of subjects having high blood pressure were belong to group 28 to 32 week gestational age (42.06%), followed by group 33 to 36 weeks gestational age (31.74%). Mean systolic blood pressure was 160.79 ± 15.158 and diastolic blood pressure was 104.37 ± 13.653 at admission.

The mean urinary protein excretion in 24 hours was 1067.8 ± 78 mg/dl (range, 200 mg/dl to 3000 mg/dl). The mean P/c ratio was 0.338 ± 0.15 (range 0.143 to 0.72). The correlation coefficient for the P/C ratio against the 24 hour urine protein excretion was 0.71 (R² = 0.71).

The predictor model of regression equation was Y = −269.19 + 3715.15 X, (P < 0.0001) where X = spot protein creatinine ratio and Y = 24 hour urine protein.

**Figure 1** shows the relationship between 24 hours protein excretion and spot protein-creatinine (P/C) ratio. The curve indicates very good linear relationship between the two variables which is also indicated by strong correlation coefficient (r = 0.71, P = 0.0001).

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**Table 1.** Distribution of cases according to clinical parameters.

<table>
<thead>
<tr>
<th>Range Age (years)</th>
<th>No. of cases (n = 126)</th>
<th>Percentage</th>
<th>Mean</th>
<th>SD</th>
</tr>
</thead>
<tbody>
<tr>
<td>&lt;20</td>
<td>9</td>
<td>7.14</td>
<td>25.151</td>
<td>3.769</td>
</tr>
<tr>
<td>21 - 30</td>
<td>107</td>
<td>87.92</td>
<td></td>
<td></td>
</tr>
<tr>
<td>31 - 40</td>
<td>10</td>
<td>7.9</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Parity</td>
<td></td>
<td></td>
<td>1.262</td>
<td>1.465</td>
</tr>
<tr>
<td>P0</td>
<td>58</td>
<td>46.03</td>
<td></td>
<td></td>
</tr>
<tr>
<td>P1</td>
<td>24</td>
<td>19.04</td>
<td></td>
<td></td>
</tr>
<tr>
<td>P2</td>
<td>16</td>
<td>12.7</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Gestational age (in weeks)</td>
<td>No. of cases</td>
<td>Percentage</td>
<td>Mean</td>
<td>SD</td>
</tr>
<tr>
<td>28 - 32</td>
<td>53</td>
<td>42.06</td>
<td></td>
<td></td>
</tr>
<tr>
<td>33 - 36</td>
<td>40</td>
<td>31.74</td>
<td></td>
<td></td>
</tr>
<tr>
<td>37 - 40</td>
<td>31</td>
<td>24.60</td>
<td></td>
<td></td>
</tr>
<tr>
<td>&gt;40</td>
<td>2</td>
<td>1.58</td>
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<td></td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Systolic Blood pressure (mm·Hg)</th>
<th>No. of cases</th>
<th>Percentage</th>
<th>Mean</th>
<th>SD</th>
</tr>
</thead>
<tbody>
<tr>
<td>&lt;160</td>
<td>50</td>
<td>39.68</td>
<td></td>
<td></td>
</tr>
<tr>
<td>≥160</td>
<td>76</td>
<td>60.32</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Diastolic blood pressure (mm·Hg)</td>
<td>No. of cases</td>
<td>Percentage</td>
<td>Mean</td>
<td>SD</td>
</tr>
<tr>
<td>&lt;110</td>
<td>74</td>
<td>58.73</td>
<td></td>
<td></td>
</tr>
<tr>
<td>&gt;110</td>
<td>52</td>
<td>41.27</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

**Table 2.** Area under the curve. Test result variable(s): VAR 0001.

<table>
<thead>
<tr>
<th>Area</th>
<th>Std. Error</th>
<th>Asymptotic Sig</th>
<th>Asymptotic 95% Confidence Interval</th>
</tr>
</thead>
<tbody>
<tr>
<td>0.793</td>
<td>0.068</td>
<td>0.001</td>
<td>0.659</td>
</tr>
</tbody>
</table>

*Null hypothesis: true area = 0.5.*
protein creatinine ratio provides excellent discrimination. Figure 2 shows ROC curve, area under the curve was 0.793 (with 95% CI, P < 0.001).

In Figure 3 graph is showing sensitivity and specificity at different level of spot urine protein creatinine ratio. This graph shows that with increasing sensitivity the specificity of spot protein creatinine ratio were decreased at various cut off points. At 0.14 cut off value, sensitivity was highest 99.1% and specificity 50%. At cut off value 0.37, the sensitivity 31% and specificity 100%. At cut off p/c ratio 0.25 was identified as the best threshold to detect 24 hrs urine protein excretion of 300 mg/24 hr with sensitivity and specificity 69% and 75%. The positive predictive value was 92.3% and negative predictive value was 90.4%.

Table 2 shows area under the curve with 95% confidence interval.

Figure 2 shows ROC curve, the area under the curve has been calculated, which came out to be 0.79.

In Figure 3, graph showing sensitivity and specificity at different levels of spot urine P/C ratio.

4. DISCUSSION

Pre-eclampsia is the 3rd leading cause of maternal mor-
tality and morbidity [1]. The present study included 126 women with pregnancy induced hypertension. The highest incidence of PIH was found in the age group 21 - 30 yrs (87.92%) in our study with the mean age of total patients being 25.15 ± 3.77 yrs. Majority of patients of PIH in our study were nulliparous (46.03%). Majority of subjects having high blood pressure were belonged in group 28 - 32 weeks gestational age (42.06%). In our study majority of cases had systolic blood pressure ≥ 160 mm·Hg (60.32%) and diastolic blood pressure < 110 mm·Hg (58.73%). This is supported by the findings of J. O. Eigbefoh et al. [5] who has also reported the maximum incidence in age group 20 - 29 years (75.6%) followed by age group 32 - 39 years and majority of subject were nullipara (54.7%) followed by primipara (15.15%).

In our study sensitivity, specificity, positive predictive value, and negative predictive value calculated at various cut off points of P/C ratio. Maximum sensitivity 99.1% was found with protein creatinine ratio cut off point 0.14 and maximum specificity 100% was found with protein creatinine ratio cut off point 0.37. A very good linear relationship exists between 24 hr protein excretion and spot protein creatinine ratio with strong correlation coefficient (r = 0.71, P = 0.0001).

By the Receiver Operating Characteristic (ROC) curve analysis, the area under the curve has been calculated, which came out to be 0.79 which indicates that the spot urine P/C ratio provides excellent discrimination (Table 2). The P/C ratio of 0.25 was identified as the best cut-off point to detect urine protein excretion of >300 mg/24 hr. The positive predictive value was 92.3% and negative predictive value was 90.4%.

Attempts for establishing spot P/C ratio as a marker for pre-eclampsia have been done past also. Durnwald and Mercer [6] determined lower correlation coefficients of 0.56 and 0.41 and cautioned against substituting spot P/C ratio for the 24 hour urine protein collection. Al and colleagues [4] found significant association between 24-hour protein and random protein-creatinine ratio (r = 0.56, P < 0.01). With a cut off protein creatinine ratio greater than 0.19 as a predictor of significant proteinuria, sensitivity and specificity were 85% and 73%. Positive and negative predictive values were 46% and 95% respectively. Wheeler and colleagues [7] found in their study, that the P/C ratio of 0.46, 0.82 and 3.0 represented 1000 mg/ 24 h 2000 mg/ 24 hr and 3000 mg/24 hr, respectively. All of these were characterized by excellent accuracy. The urine P/C ratio of 0.21 corresponded with a protein excretion rate of 300 mg /24 hr. The area under curve was 0.8% indicating good accuracy which supports our findings. Nahid Shahbazian et al. [8] found the strong correlation between the spot P/C ratio and 24 hr urine protein excretion (r = 0.84, P < 0.001). The optimal spot P/C ratio cut off point was 0.2 for 300 mg/ 24 hr of
protein excretion, with a sensitivity, specificity, positive predictive value and negative predictive value of 91.2%, 87.8%, 94.4% and 96.8% respectively.

Our study suggests that the spot urine protein-creatinine ratio provides excellent discrimination between insignificant and significant proteinuria in preeclampsia. This test can be used for prompt clinical decision in spite of waiting for 24 hr urine collection. Drawback of our study is large number of false negative cases (30%). Further studies should be done large sample size and utility of this test in ambulatory management of preeclamptic patients.

5. CONCLUSION

There is a strong correlation between spot urine P/C ratio and the 24 hour urine protein excretion. Spot urine protein-creatinine ratio can be used to predict the amount of 24 hours urine protein excretion with high accuracy. But the 24 hr urine collection remains the gold standard for evaluation for pre-eclampsia. So, spot urine protein-creatinine ratio can be used for detection of significant proteinuria in pregnant women with suspected pre-eclampsia with high accuracy, which is more rapid than 24 hour urine protein excretion. Thus this quick method with high accuracy is very useful to prevent feto-maternal morbidity and mortality in India.

REFERENCES


ABBREVIATIONS

P/C ratio-protein/creatinine ratio
ROC curve-received operating characteristics curve
CI-coefficient interval