Is there any relationship between RDW levels and atrial fibrillation in hypertensive patient?

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Abstract

Background: Atrial fibrillation (AF) is the most common cardiac arrhythmia and increases the risk of stroke and death. Patients with hypertension have an increased risk of developing atrial fibrillation. RDW (Red blood cell distribution width) levels are elevated in cardiovascular disorders including heart failure, stable coronary disease, acute coronary syndrome, slow coronary flow and stroke.

Objective: We aimed to investigate the relation between RDW and AF in patients with hypertensive

Method: We retrospectively examined 126 consecutive hypertensive patients (63 hypertensive patients with AF and 63 hypertensive patients without AF matched with age and sex

Results: The mean age of the study population was 71.09 ± 8.50 (af group) and 70.97 ± 8.24 (non-af group) years. RDW level was different among patients with atrial fibrillation and without atrial fibrillation. (15.13 ± 1.58 and 14.05 ± 1.15 p<001). Logistic regression analysis showed that RDW and left atrial dimension were only independently risk factor associated with atrial fibrillation. (Rdw odds ratio:1.846 CI; 1.221-2.793 p<0.05). Roc curve analyses were applied to determine the cut-off point. Cut-off point was at 14,195 and Sensitive, specificity was %71,4, %56 respectively.

Conclusion: RDW levels were higher in hypertensive patients with atrial fibrillation. An increased RDW level in the patient with hypertension may alert physician on developing or presence of atrial fibrillation.

Keywords: atrial fibrillation, red blood cell distribution width, hypertension.

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Introduction

Atrial fibrillation (AF) is the most common cardiac arrhythmia and increases the risk of stroke and death. Several atherosclerotic risk factors, such as obesity, diabetes and hypertension, are associated with development of AF. (1-3) Many studies have shown that individuals with hypertension have an increased risk of developing atrial fibrillation compared with normotensive individuals. (4-6) Several inflammatory markers such as C-reactive protein (CRP), tumor necrosis factor-α (TNF-α), interleukin (IL)-2, IL-6, IL-8, and monocyte chemo attractant protein (MCP)-1 and mediators have been shown associated with the presence or the outcome of AF. (7) A growing body of literature supports a role for the immune system and chronic inflammation in the development of hypertension. (8; 9) Red blood cell distribution width (RDW) is a measure of the variability in size of circulating erythrocytes. RDW is obtained routinely in standard complete blood cell counts (CBCs). Increased RDW indicates the presence of anisocytosis, which is related to impaired erythropoiesis and erythrocyte degradation, reflecting chronic inflammation and a high level of oxidative stress. (10-12) Recent studies have reported that RDW levels are elevated in cardiovascular disorders including heart failure, stable coronary disease, acute coronary syndrome, slow coronary flow and in non-cardiovascular diseases, stroke. (13-17)

However, to the best of our knowledge, to evaluate the relation between RDW and AF in patients with hypertensive has not been reported in the literature previously. In this study, we aimed to investigate the relation between RDW and AF in patients with hypertensive
Method

Patient selection
We retrospectively examined 126 consecutive hypertensive patients (63 hypertensive patients with AF and 63 hypertensive patients without AF matched with age and sex) at university of Bozok and GOP. All patients’ records of history and physical examinations were examined in detail. No patient had a recent history of an acute infection or an inflammatory disease. Patients with renal failure, concomitant valvular disease, cardiomyopathy, and previous cardiac surgery and seconder hypertension were excluded.

Echocardiographic assessment
Transthoracic 2-dimensional and Doppler echocardiographic assessment was performed. (Aloca, Japon). Measurements of the left atrium, left ventricle and right ventricle was obtained from parasternal long axis view according to standard criteria. Left ventricular ejection fraction (LVEF) was calculated using the modified Simpson’s rule in the 2- and 4-chamber apical views.

Measurement of laboratory parameters
On admission, venous bloods were obtained from the antecubital vein from all patients after fasting for at least 8 h. Fasting plasma glucose (FBG), creatinine, plasma levels of total cholesterol (TC), low-density lipoprotein (LDL), high-density lipoprotein (HDL) cholesterol and triglyceride (TG) were determined by a hospital auto analyzer. CBC analysis including RDW was performed with Beckman Coulter LH 750 automated analyzer within 2 h.

Statistic
The statistical analyses were performed using software (SPSS 18.0). Parametric values were given as mean±standard deviation and non-parametric values were given as a percentage. To compare parametric continuous variables, Student’s t-test was used; to compare nonparametric continuous variables, the Mann–Whitney U-test was used. Categorical data were compared by Chi-square distribution. Receiver operator characteristic (ROC) curve analysis was performed to identify the optimal cut-off point of RDW (at which sensitivity and specificity would be maximal) for the prediction of AF. Areas under the curve (AUC) were calculated as measures of the accuracy of the tests. We compared the AUC with the use of the Z test. Variables found to be statistically significant in univariate analyses were entered into multivariate logistic regression analysis. Multivariate logistic regression models were created to identify independent predictors of AF. Two-tailed P-values of less than 0.05 were considered to indicate statistical significance.

Results
There were no significant differences among patients with or without atrial fibrillation in terms of gender, age, coronary artery disease, diabetes Mellitus. The mean age of the study population was 71,09± 8,50 (af group) and 70,97±8,24 (non-af group) years. LVEF and left atrium dimensions (LAD) TG,TC,and RDW levels were different between the two groups (Table 1).

Table 2. Multivariate regression analyses

<table>
<thead>
<tr>
<th></th>
<th>Odds ratio</th>
<th>CI</th>
<th>P value</th>
</tr>
</thead>
<tbody>
<tr>
<td>RDW</td>
<td>1,846</td>
<td>1,221-2,793</td>
<td>0,004</td>
</tr>
<tr>
<td>EF</td>
<td>0,914</td>
<td>0,835-1,004</td>
<td>0,06</td>
</tr>
<tr>
<td>LAD</td>
<td>6,985</td>
<td>2,451-19,904</td>
<td>0,001</td>
</tr>
<tr>
<td>TC</td>
<td>0,992</td>
<td>0,982-1,002</td>
<td>0,137</td>
</tr>
<tr>
<td>TG</td>
<td>1,000</td>
<td>0,993-1,006</td>
<td>0,904</td>
</tr>
</tbody>
</table>

### Table 1. Baseline characteristics of patients

<table>
<thead>
<tr>
<th>Groups</th>
<th>HT(+) AF(+) N:63</th>
<th>HT(+) AF(-) N:63</th>
<th>P value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age</td>
<td>71.09±8.50</td>
<td>70.97±8.24</td>
<td>0.944</td>
</tr>
<tr>
<td>Sex(woman)%</td>
<td>52.2</td>
<td>47.8</td>
<td>0.360</td>
</tr>
<tr>
<td>DM%</td>
<td>36.5</td>
<td>39.5</td>
<td>0.283</td>
</tr>
<tr>
<td>CAD%</td>
<td>22.2</td>
<td>33.3</td>
<td>0.232</td>
</tr>
<tr>
<td>Glucose</td>
<td>108.178±29.354</td>
<td>112.05±33.80</td>
<td>0.855</td>
</tr>
<tr>
<td>Creatinin</td>
<td>1.16±1.21</td>
<td>0.90±0.23</td>
<td>0.314</td>
</tr>
<tr>
<td>TG</td>
<td>115.03±70.97</td>
<td>154.05±105.68</td>
<td>0.023</td>
</tr>
<tr>
<td>TC</td>
<td>170.26±46.13</td>
<td>214.81±80.86</td>
<td>0.002</td>
</tr>
<tr>
<td>HDL</td>
<td>42.85±8.26</td>
<td>44.83±16.75</td>
<td>0.140</td>
</tr>
<tr>
<td>LDL</td>
<td>106.11±31.91</td>
<td>123.94±43.28</td>
<td>0.28</td>
</tr>
<tr>
<td>RDW</td>
<td>15.13±1.58</td>
<td>14.05±1.157</td>
<td>0.001</td>
</tr>
<tr>
<td>HB</td>
<td>13.74±1.38</td>
<td>13.88±1.62</td>
<td>0.323</td>
</tr>
<tr>
<td>EF%</td>
<td>52.68±8.20</td>
<td>57.98±4.84</td>
<td>0.001</td>
</tr>
<tr>
<td>LAD</td>
<td>4.43±0.78</td>
<td>3.8±0.27</td>
<td>0.001</td>
</tr>
</tbody>
</table>

Dm: diabetes mellitus  
CAD: coronary artery disease  
TG: Triglyceride  
TC: Total cholesterol  
HDL: High Density Cholesterol  
LDL: Low Density Cholesterol  
RDW: Red Blood Cell Distribution Wide  
Hb: Hemoglobin  
EF: Ejection fraction  
LAD: Left Atrial Diameter

The patient with AF had less EF (%52.68±8.20 and 57.98±4.84 p<0.05) and LAD (4.43±0.78 and 3.8±0.27 p<0.05) than non-atrial fibrillation group. TC (170.26±46.13 and 214.81±80.86 p<0.05), TG (115.03±70.97 and 154.05±105.68 p<0.05) and Hb (13.74±1.38 and 13.88±1.62 p<0.05) were different between patients with or without atrial fibrillation respectively. (Table 1) RDW level was different among patients with atrial fibrillation and without atrial fibrillation. (15.13±1.58 and 14.05±1.15 p<0.01) (table 1). Variables found to be statistically significant in univariate analyses were entered into multivariate logistic regression analysis. Multivariate Logistic regression analysis showed that RDW and left atrial dimension were only independently risk factor associated with atrial fibrillation. (Rdw odds ratio:1.846 CI; 1.221-2.793 p<0.05) (table 2).
Roc curve analyses were applied to determine the cut-off point. (Figure 1) Cut-off point was at 14,195 and Sensitive, specificity was %71, 4 and %56 respectively.

**Figure 1. ROC curve analysis of RDW**

![ROC Curve](image)

**Discussion**

Our study showed that an increased RDW level was independently associated with atrial fibrillation in the patient with hypertension. The cut of point of 14,195 levels was a strong predictor of atrial fibrillation in the patient with hypertension. The sensitive, specificity was %71,4 and %56 respectively.

Many studies have shown that individuals with hypertension have an increased risk of developing atrial fibrillation compared with normotensive individuals. (4-6) Elevated systolic blood pressure is associated with increases in left atrial fibrosis,(18; 19) which in turn is related to prevalent atrial fibrillation. (20) Some studies suggest that left ventricular hypertrophy and increases in left atrial size may also mediate the relationship between blood pressure and incident atrial fibrillation.(21; 22)

In our study, LAD was higher in patient with hypertensive patients with AF that those without AF. Anatomic evidence of inflammation, such as myocyte necrosis, inflammatory infiltrates and fibrosis have been shown in the atrial wall in biopsies of patient with persistent. (23) Several inflammatory markers such as CRP, TNF-α, IL-2, IL-6, IL-8, and MCP-1 and mediators have been shown associated with the presence or the outcome of AF. (7) Patients with high baseline CRP levels are at higher risk of having postoperative atrial fibrillation in both on-pump and off-pump surgery.(24)

RDW is a numerical measure of the size variability of circulating erythrocytes and is routinely reported as a component of complete blood count in the differential diagnosis of anemia. Studies recently have shown that RDW was strong and independent predictor of poor outcome in the general population.(25) RDW has been recently reported that being predictive of poor outcome in several cardiovascular disease including heart failure, stable coronary artery disease and acute myocardial infarction.(7; 15; 26; 27) The mechanism of elevated RDW values are unknown that why it is linked with adverse outcomes. Some studies have been suggested that RDW may reflect prognostic markers in heart failure, such as inflammatory cytokines that may affect bone marrow function and iron metabolism.(28; 29) Agarwal and at all. have demonstrated that subjects with low cardiorespiratory fitness have elevated RDW by virtue of chronic inflammation and high oxidative stress, which is likely to promote atherosclerosis and lead to cardiovascular disease.(19)

A growing body of literature supports a role for the immune system and chronic inflammation in the development of hypertension.(8; 9) Higher RDW values were strongly correlated with higher systolic and diastolic blood pressures.(30) RDW was significantly increased in patients with non-dipper hypertension compared with the dipper hypertension. Inflammatory activity
was closely related to RDW in non-dipper hypertensive patient. In present study, Increased RDW, as a surrogate marker of inflammation, was associated with atrial fibrillation in the patient with hypertensive.

Inflammatory activity is increased in the patient with hypertension. Also, inflammatory activity plays a major role in the presence and developing of atrial fibrillation. RDW is a new inflammatory marker. Therefore, developing of atrial fibrillation in the patient with hypertension may not only associated with diastolic dysfunction also may be related to inflammation but why RDW is increased in the patient with atrial fibrillation with hypertensive is not clear. However, best of our knowledge to evaluate the relation between RDW and AF in patients with hypertensive has not been reported in the literature previously. The present study revealed that RDW levels were higher in hypertensive patients with AF than that those without AF. Incidence of hypertension is high in the general population. Developing atrial fibrillation in this patient may poorly affect prognosis of the patient. An increased RDW level in the patient with hypertension may alert physician on developing or presence of atrial fibrillation.

Conclusion

RDW levels were higher in hypertensive patients with AF than that those without AF. An increased RDW level in the patient with hypertension may alert physician on developing or presence of atrial fibrillation.

Limitations

The main limitation of our study was the small sample size. A small sample size can result in a low statistical power for equivalency testing, leading to false-negative results. Second, because of the retrospective nature of data collection, echocardiographic parameters were not obtained concomitantly with blood sampling. Also, we were unable to evaluate some factors, such as Vitamin B12, ferritin that have been effected to the RDW, because data on these variables were not collected and we cannot apply our results for general population due to the broad exclusion criteria.

Author contributions

Sarıkaya S conceived and designed the study and was responsible for the acquisition of data. Sahin S conceived and designed the study and was responsible for the analysis and interpretation of data, manuscript draft and critical revision of the manuscript. Akyol L and Börekci E were responsible for the acquisition of data. Yılmaz YK conceived and designed the study and was responsible for the acquisition of data. Altunkas F conceived and designed the study and was responsible for the analysis and interpretation of data and critical revision of the manuscript. Karaman K was responsible for manuscript draft and critical revision of the manuscript.

References


