

Prognosis of Chronic Myeloid Lymphoma with Red Cell Distribution Width

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Received date: June 15, 2018; Accepted date: June 22, 2018; Published date: June 27, 2018

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Citation: Khani MA, Karimi Z (2018) Prognosis of Chronic Myeloid Lymphoma with Red Cell Distribution Width. J Blood Res Vol.1 No. 1:5.

Editorial

Cancer is one of the leading causes of mortality in the world and is increasing every year, as of 2017, 600920 deaths due to cancer were reported in the United States [1,2]. So, it is necessary to prognosticate these diseases.

Some simple parameters, automatically reported by laboratory blood analyzers, can have multiple clinical applications; for example, Red cell distribution width (RDW) is regularly evaluated as part of a complete blood count (CBC) for collecting information from circulatory evaluations. RDW represents the abnormal survival of red blood cells and it has a high negative predictive value (NPV) to diagnose all types of disorders; it may be useful to evaluate the short-term and long-term prognosis. Recently, RDW has been widely recognized to have an important role in carcinogenesis, tumor progression, and prognosis [3,4]. RDW has been reported as an inflammatory marker for systemic inflammatory response in many diseases. This parameter has been studied as a prognostic factor and an indicator of disease activity in many malignancies; these studies are mainly conducted in cardiovascular diseases and have been studied in very low hematological diseases.

In hematological malignancies such as chronic myelogenous leukemia (CML) who some patients have unfavorable prognosis after treatment and it is extremely variable. Many prognostic factors are used to predict clinical outcome to date treatment response indicated by cytogenic or molecular evaluation.

There have been few but valuable studies in this regard. Zack et al. concluded in their study that RDW increases in hairy cell leukemia (HCL), which is related to the activity of the disease and becomes normal after treatment. In a study, examined the significance of RDW in patients with CML that an important role in the classification of patients to predict the responses and treatment outcomes; In fact, the RDW value in CML patients is higher than normal in most cases, it may predict the treatment response, this type of classification facilitates the planning of treatment [5]. Along with the studies that confirmed a significant association between RDW and an

increased risk of cancer and hematology, a study showed that RDW is stable and does not represent time-prognosis [6].

Though in previous studies relationship between elevated RDW and mortality has been reported in several contexts but in a study by Rafsanjani et al., there was a different outcome in which there was no relationship between the RDW level and the rate of mortality and relapse in pediatric patients [7]. On the other hand, Increasing RDW by age is a true biomarker of inflammatory and age-related diseases and is a major predictor of mortality [8]. That could be the reason for this difference; this connection should be further investigated in the future.

Investigating the relationship between RDW and mutations, changes in telomere length and prognosis of patients may provide a new insight into the treatment of CML. It's suggested that in the future, the findings should be considered in larger groups, and in different treatment categories. It seems to be helpful enter RDW parameter in the routine checkup tests list for the physician in the diagnosis and treatment of disease.

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