

# Assessment of Hypercoagulability (Platelet Count and Indices) in Sudanese Patients with Acute Myeloblastic Leukemia

Hana Salah Taha<sup>1</sup>, Tarig Altayeb Mohammed<sup>2</sup>, Ahmed El-basher Abdlegader<sup>3</sup>, Amar B Elhussein<sup>4</sup>, Mohand Gafar Nabag<sup>5</sup>, IhabHamed Nourein<sup>6</sup>

<sup>1,2</sup>AL-Yarmouk College, Department of Hematology and Immunohematology, Sudan

<sup>3</sup>Department of Hematology, Collage of Applied Medical Sciences, Najran University, KSA

<sup>4</sup>Department of Biochemistry, Collage of Applied Medical Sciences, Najran University, KSA

<sup>5</sup>Faculty of Public Health & Health informatics, Umm Al-Qura University, KSA

<sup>6</sup>Department of Histopathology, Collage of Medical Laboratory Sciences, University of Science and Technology, Sudan

**Abstract:** This study was done in patients with acute myelogenous leukemia in Radiation and Isotopes Centre Khartoum, Khartoum state, Sudan, during the period from November 2015 to March 2016. **Aim:** This study aimed to measure the Platelet count and indices in patients with acute myelogenous leukemia. **Methods:** Venous blood (2.5ml) were collected in Ethylenediamine Tetra Acetic acid container from patients with acute myelogenous leukemia and measurement by automation technique (Sysmex KX-21N for complete blood count). **Results:** The result showed that the mean of platelets count was  $(164.82 \pm 182.18 \times 10^3 / \mu l \text{ SD})$  with p-value (0.004) while the mean of PDW was  $(10.2 \pm 8.8 \text{ fL})$  with p-value (0.08) and the mean of the MPV was  $(6.1 \pm 4.3 \text{ fL})$  with p-value (0.00). **Conclusion:** Insignificant change in platelet count and indices of patients with acute myeloid leukemia when correlated with age, duration, and treatment. Also significant difference of mean PL.C and MPV, as well as insignificant mean of PDW was observed when it correlated with healthy control.

**Keyword:** Hypercoagulability, Acute myeloblastic leukemia, PL.C, MPV, PDW

## 1. Introduction

The leukemia is a group of disorders characterized by the accumulation of malignant white cells in the bone marrow and blood, and lead to (i) bone marrow failure (i.e. anemia, neutropenia, and thrombocytopenia) and (ii) infiltration of organs. "Leukemia" is the general term for four different types of blood cancer called: Acute lymphocytic (lymphoblastic) leukemia (ALL), Acute myelogenous (myeloid) leukemia (AML), Chronic lymphocytic leukemia (CLL) and Chronic myelogenous leukemia (CML)<sup>(1, 2)</sup>.

Acute myeloid leukemia (AML) is a malignant disease characterized by an accumulation of immature myeloid blast cells in the bone marrow and most often in the peripheral blood. AML can also be present in other tissues such as in the skin<sup>(3, 4)</sup>.

Globally, during the period from 2009-2013 the number of new cases of acute myeloid leukemia and the number of death were 4.1/100.000 and 2.8/100.000 of population per year respectively, in contrast the number of new cases in 2016 was 19.950/100.000 populations and the number of death was 10.430/100.000 populations of all new cancer cases<sup>(5)</sup>.

In Sudan the Leukemia is one of the most significant current health problems, it associated with large population resulting of high mortality among Sudanese population<sup>(6)</sup>.

Despite of this high mortality there is no enough data reflect the relation between coagulation factors and AML Patient's. Therefore the aim of this study was to assess the hypercoagulability (platelet count and indices) in Sudanese patients with acute myeloblastic leukemia.

## 2. Materials and Methods

This study was case control, hospital base study, conducted at Radiation and Isotopes center Khartoum (RICK), Sudan.

In study 80 blood samples were included, 40 of these samples were collected from acute myeloid leukemia patient and the remaining 40 samples were collected from healthy persons and used as control.

### Samples Collection

Blood samples were collected in possible aseptic condition from mid vein of the forearm (2.5ml). Collection was done using standard venipuncture technique disposal plastic syringe and put it in Ethylene Di-amine Tetra-acetic Acid (EDTA) container, each container was numbered and every number was related to the questioner used in data collection, and then each container placed on electric mixer for 3 minutes. Then they immediately took to the lab to be analyzed using Sysmex (KX-21N), for complete blood count (CBC). Inside the machine the WBC count, RBC count, mean corpuscular volume (MCV), mean corpuscular hemoglobin concentration (MCHC) and platelets were measured by using the Direct Current detection method. The

HGB detector block measured the hemoglobin concentration using the non-cyanide hemoglobin method. After analyzing of each sample a result paper obtained from the machine containing the measurement of each parameter.

Control blood (EIGHTCHECK-3WP) using X control or L-J control program was used before starting sample analysis, to optimize the results.

The ethical clearance of this study was approved by department of hematology collage of medical laboratory science. Data were collected with verbal informed consent from patients, and the questionnaires were filled.

### 3. Results

The objective of this study is to determine the mean of Platelets count and indices in patients with AML, as well as to correlate the possible risk factor (age, duration, treatment) with platelets count and indices.

In this study of 40 studied patients 21 (52.5%) were male and 19 (47.5%) were female with age range from less than 9 years old up to 69 years old.

The result showed that the mean of platelets count was  $(164.82 \pm 182.18 \times 10^3 / \mu\text{l})$  SD) with p-value (0.004) while the mean of PDW was  $(10.2 \pm 8.8 \text{ fL})$  with p-value (0.08) and the mean of the MPV was  $(6.1 \pm 4.3 \text{ fL})$  with p-value (0.00) when compared to control group **table (1)**.

As illustrated in **figure (1)** the age of study populations was classified in to different group started from (0-9), (10-19), (20-29), (30-39), (40-49), (50-59) and (60-69). These age groups were subsequently correlated with PL.C, PDW and MPV, the result showed that, the maximum PL.C was observed at age group (0-9)  $(292 \text{ cell/mm}^3)$ , the maximum PDW and MPV at 50-59  $(15.7 \text{ fL}$  and  $6.6 \text{ fL})$  respectively. Statistically no significant association were observed

between age group and PL.C, PDW and MPV the *p. value* was  $> 0.05$  as showed in **table (2)**.

Also the study populations were subsequently classified in to three groups according to the duration of disease (0-6 months, 7-12 month and 13-48 months), accordingly, the PL.C, PDW and MPV were  $(178.42 \text{ cell/mm}^3, 116.83 \text{ cell/mm}^3$  and  $20.33 \text{ cell/mm}^3)$ ,  $(9.8 \text{ fL}, 11.3 \text{ fL}$  and  $13.3 \text{ fL})$  and  $(6.2 \text{ fL}, 5.8 \text{ fL}$  and  $5.6 \text{ fL})$  respectively as showed in **figure (2)**. Statistically no significant association were observed between the PL.C, PDW and MPV of patients and duration of the disease *p. value* ( $> 0.05$ ) as described in **table (2)**.

In this study the PL.C, PDW and MPV of patients were also correlated with the treatment (ATRA (treinoin) and (Daunorub) **figure (3)**. The mean of PL.C was normal in those that used ATRA (treinoin) and (Daunorub) drugs, while the PDW and MPV within normal range in those that used (Daunorub) treatment and insignificant decreased in those that used ATRA (treinoin). Statistically no significant association were observed between (PL.C, PDW and MPV) and treatment that used *p. value* ( $> 0.05$ ), **table (2)**.

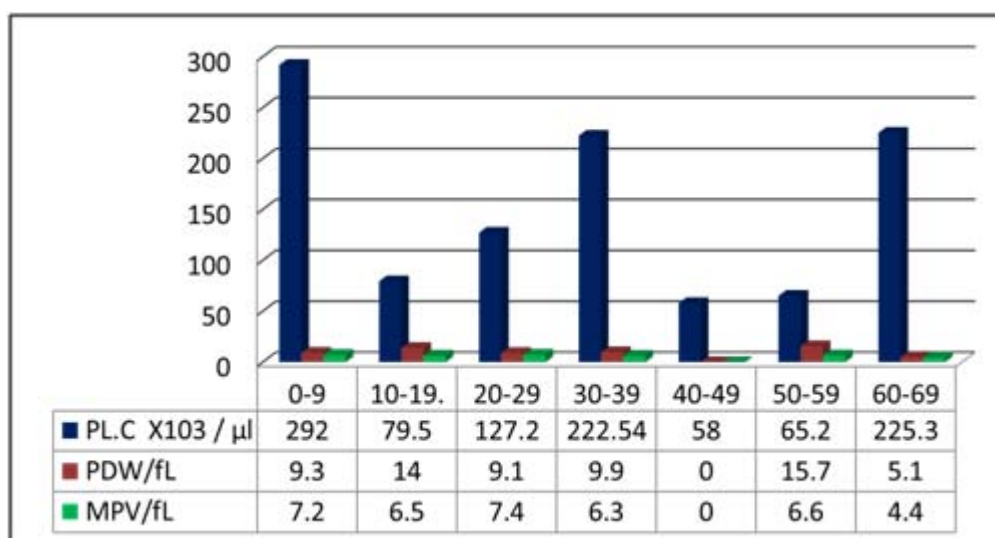
**Table 1:** Mean Differences of PL.C, PDW and MPV in AML patients and healthy control.

	Mean $\pm$ SD		P. Value
	Patients	Healthy control	
PL.C $\times 10^3 / \mu\text{l}$	$164.82 \pm 182.18 \text{SD}$	$258.10 \pm 81.0 \text{SD}$	0.004**
PDW/fL	$10.2 \pm 8.8 \text{SD}$	$12.7 \pm 1.0 \text{SD}$	0.08
MPV/fL	$6.1 \pm 4.3 \text{SD}$	$9.6 \pm 0.9 \text{SD}$	0.00**

**Table 2:** Correlation of PL.C, MPV and PDW with duration, Age and Treatment in AML

\*\*Correlation is significant if the *p-value* is less than 0.05

Risk factor	P. Value		
	PL.C $\times 10^3 / \mu\text{l}$	PDW/fL	MPV/fL
Duration	0.692	0.46	0.76
Age	0.258	0.85	0.33
Treatment	0.95	0.21	0.50



**Figure 1:** Correlation of mean PL.C, MPV and PDW of patients with age group

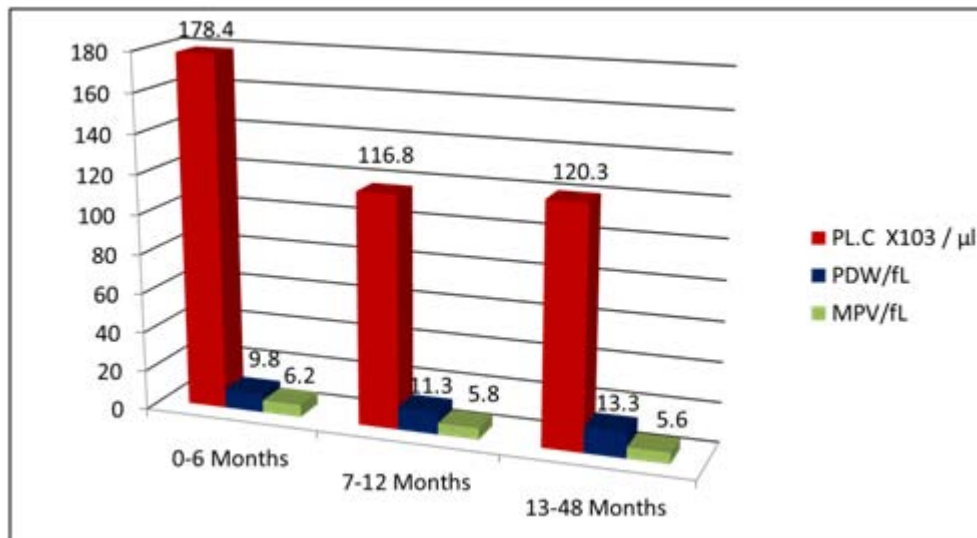


Figure 2: Correlation of mean PL.C, MPV and PDW of patients with Duration group

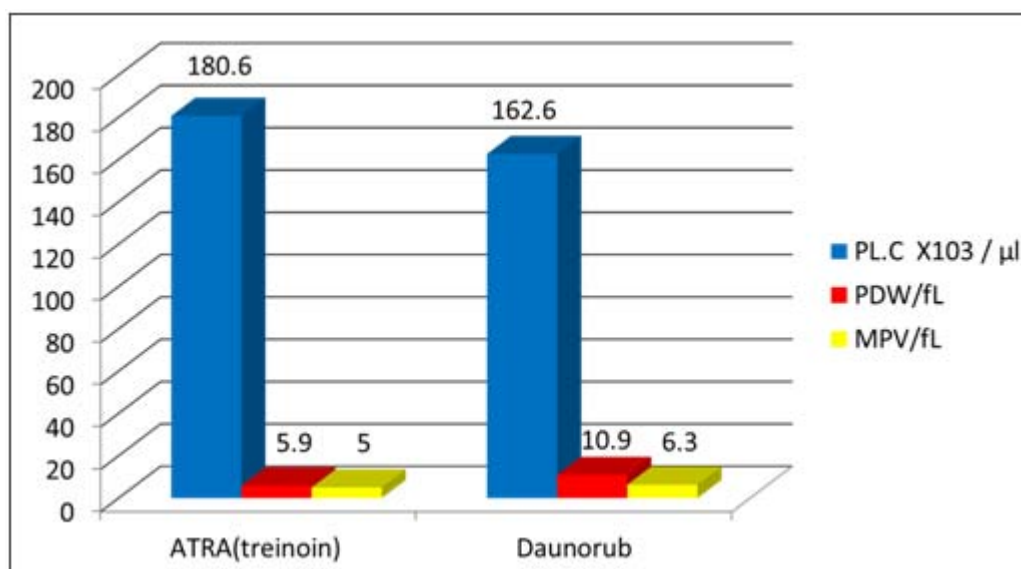


Figure 3: Correlation of mean PL.C, MPV and PDW of patients with treatment

#### 4. Discussion

This study included 80 persons, 40 patients with AML male and female were selected for this study, and 40 normal healthy individual were selected as control group. This study was measured for them platelet count and indices.

The results showed significant difference of mean platelet count between the patients and healthy control groups. And showed significantly decreased of mean MPV of patients in correlation with healthy group, and showed insignificant difference of mean PDW of patients in correlation with healthy control group, these findings is in an agreement with study carried by Kim, *et al* (2014)<sup>(7)</sup>, when they compare the platelet parameters in thrombocytopenic patients associated with AML and primary immune thrombocytopenia. In contrast our study reject the findings of Amar R, *et al* (2013)<sup>(8)</sup>, when they reported significant decreased of platelet count in study aimed to assess the role of platelet parameters in diagnosing various clinical conditions. As well as the present study disagreed with Bumbea, *et al* (2011)<sup>(9)</sup> when they assess the platelet dysfunction in acute myeloid leukemia's and myelodysplastic syndromes, they

reported significant decreased in platelet in patients AML *p.value* (0.001).

Also the result of PL.C, MPV and PDW was insignificant when correlated to age group, duration and treatment. This disagreed with Vijaya B, *et al* (1990)<sup>(10)</sup> when they reported significant decreased PL.C, MPV and PDW in patients with AML when correlated to age and sex.

#### 5. Conclusion

The present study was concluded that, insignificant change in platelet count and indices of patients with acute myeloid leukemia when correlated with age, duration, and treatment. Also significant difference of mean PL.C and MPV, as well as insignificant mean of PDW was observed when it correlated with healthy control.

## References

- [1] Hoffbrand A, Moss P, and Pettit J, "Essential Haematology" 5<sup>th</sup>ed by black well publishing, Australia 2006.
- [2] Swerdlow S, "WHO classification of tumours of haematopoietic and lymphoid tissues," in World Health Organization classification of tumours, IARC Press: Lyon. pp. 110-123, 2008.
- [3] Sharma, SK, "Leukemia cutis: an unusual presentation," *Indian J Hematol Blood Transfus.* Vol. 28, no (3), pp. 175-7, 2012.
- [4] Rao AG and Danturty I, "Leukemia cutis," *Indian J Dermatol.* Vol.57, no (6). pp. 504, 2012.
- [5] SEER Stat Fact Sheets: Leukemia. National Cancer institute  
<http://seer.cancer.gov/statfacts/html/amyl.html>  
[national cancer institute 2016.](http://seer.cancer.gov/statfacts/html/amyl.html)
- [6] Intisar ES, Hsin YW, Kamal HM and Sulma IM, "Cancer incidence in Khartoum, Sudan: first results from the Cancer Registry, 2009–2010," *Cancer Medicine*, vol.3 no (4), pp. 1075–1084, 2014. Doi: [10.1002/cam4.254](https://doi.org/10.1002/cam4.254).
- [7] Kim MJ, Park PW, Seo YH, Kim KH, Seo JY, Jeong JH, Park MJ and Ahn JY, "Comparison of platelet parameters in thrombocytopenic patients associated with acute myeloid leukemia and primary immune thrombocytopenia," *Blood Coagul Fibrinolysis*, vol. Apr, 25, no (3), pp. 221-5, 2014.
- [8] Doi: [10.1097/MBC.0000000000000027](https://doi.org/10.1097/MBC.0000000000000027).
- [9] Amar RS, Sanjay NC, Menka HS, "Role of Platelet Parameters in Diagnosing Various Clinical Conditions," *Natl J Med Res*, vol.3, no (2), pp. 162-165, 2013.
- [10] Bumbea, Minidora O, "Platelet Dysfunction in Acute Leukemias and Myelodysplastic Syndromes," *Rom. J. Intern. Med*, vol. 49, no (1), pp. 93–96, 2011.
- [11] Vijaya B, Reddy MD, Areta KV, Debra AH, Arvind K, Jeanine MW, Jawed F, Schumacher HR, "Global and Molecular Hemostatic Markers in Acute Myeloid Leukemia," *American Journal of Clinical Pathology*, vol. 94, no(4), pp.397-403, 1990.
- [12] DOI: <http://dx.doi.org/10.1093/ajcp/94.4.397>.