Megakaryocytic Clustering in Chronic Myeloid Leukemia: Can it be a Predictor of Clinical Outcome?

Zunairah Mughal¹, Hira Babar², Sobia Ashraf¹, Ambareen Hamid¹, Abbas Khokhar³ and Samina Qamar¹

¹Department of Pathology, King Edward Medical University, Lahore, Pakistan

²Department of Haematology, King Edward Medical University, Lahore, Pakistan

³Department of Oncology, King Edward Medical University, Lahore, Pakistan

ABSTRACT

Objective: To compare clinical outcome in patients of chronic myeloid leukemia (CML) with and without megakaryocytic clustering.

Study Design: Cross-sectional comparative study.

Place and Duration of Study: Pathology Department and CML Clinic Oncology Department, King Edward Medical University from March 2018 to March 2019.

Methodology: Ninety-four patients diagnosed with chronic phase of CML were included. Complete record of complete blood count, splenomegaly, findings of bone marrow aspirate and trephine biopsy was noted. Bone marrow trephine biopsy was reviewed for megakaryocytic clustering. Sokal scoring was done; and follow-up data for clinical outcome, i.e complete hematological response (CHR) at 3 months and major molecular response (MMR) at 6 months and 1 year (as per Institute's protocol) was obtained. All the data were analysed using SPSS version 25.

Results: Megakaryocytic clustering was present in 57 (60.6%) patients and absent in 37 (39.4%). In patients with megakaryocytic clustering, CHR was absent in 12 (21.1%), MMR at 6 months was absent in 21 (36.8%) and MMR at 1 year was absent in 25 (43.9%) patients. In patients without megakaryocytic clustering, absent CHR, MMR at 6 months and MMR at 1 year were seen in 1 (2.7%), 2 (5.4%) and 2 (5.4%)x, respectively. The correlation of megakaryocytic clustering and high sokal score was found to be statistically significant with a p-value <0.001.

Conclusion: Patients with megakaryocytic clustering have poor clinical outcome as indicated by their sokal score, absent CHR, MMR at 6 months and 1 year.

Key Words: Chronic myeloid leukaemia, Megakaryocytic clustering, Complete haematological response, Major molecular response, Cytogenetic response.

How to cite this article: Mughal Z, Babar H, Ashraf S, Hamid A, Khokhar A, Qamar S. Megakaryocytic Clustering in Chronic Myeloid Leukemia: Can it be a Predictor of Clinical Outcome?. *J Coll Physicians Surg Pak* 2021; **31(01)**:34-38.

INTRODUCTION

Chronic myeloid leukemia (CML) is characterised by 5' promotor on chromosome 9 translocation to 3' exon on chromosome 22, resulting in constitutive activation of JAK-STAT pathway leading to uncontrolled proliferation of myeloid series.¹ The disease is more common in males aging 55-60 years. The prominent clinical features of disease include fatigue, anorexia, weight loss, massive spleen. Complete blood counts show abnormally high total leucocyte count with presence of myeloid series cells in all stages of maturation. It is often accompanied by anemia and/or thrombocytopenia in blood.²

The disease is characterised by 3 phases: chronic, accelerated, and blast. Majority of the patients present in the chronic phase, around 10% present in the accelerated phase, and another 10% in the blast phase.³ Bone marrow trephine findings range from hypercellular marrow with minimal fibrosis to advance fibrosis,⁴ with or without megakaryocytic clustering. These findings help in deciding about appropriate choice of treatment.

Megakaryocytic clustering is defined as a cluster of 3-5 megakaryocytes in trephine biopsy. They are pleomorphic with topographic disorganisation. The clusters may be present in sheets.⁵

Several scores are used for staging of this disease. These include Hashford, EUTOS and sokal score.

Sokal score is widely used in patients for risk stratification at the time of presentation. It takes into account four clinical variables: age, size of spleen, percentage of blast cells, and platelet count. The hazard ratio (Sokal score) is calculated by entering data in the following equation: $11 \text{ Exp } [0.116 (\text{age}-43.4)] + 0.0345 (\text{spleen size}-7.51) + 0.188 [(\text{platelets}/700) 2 - 0.563] + 0.0887$

Correspondence to: Dr. Sobia Ashraf, Department of Pathology, King Edward Medical University, Lahore, Pakistan

E-mail: s.qayyum81@gmail.com

Received: September 04, 2020; Revised: November 12, 2020;

Accepted: December 30, 2020

DOI: <https://doi.org/10.29271/jcpsp.2021.01.34>

(blast % > 2.10). Based upon score, patients are divided in three groups: low risk group (sokal score < 0.8), intermediate group (sokal score 0.8-1.2), and high risk group (sokal score > 1.2).⁶

Patients are followed up for complete hematological response (CHR) and major molecular response (MMR). CHR is defined as total leucocyte count of less than $10 \times 10^9/L$, platelet count of $450 \times 10^9/L$, absence of immature cells (blasts, promyelocytes, or myelocytes) in the peripheral blood, and disappearance of all clinical signs and symptoms including splenomegaly. MMR is achieved when BCR-ABL fusion gene cannot be demonstrated in the cases.^{7,8}

Tyrosine kinase inhibitors (TKI's) are a novel approach in management of CML.⁹ There are three generations of TKI's, Imatinib is first line drug, given to patients in chronic phase.¹⁰ It is given in a dose of 600 mg. It cause complete hematological response in 89% patients and complete cytogenetic response in 10%. Median survival rate is 7 months.¹¹

According to WHO guidelines, megakaryocytic clustering and fibrosis are taken as presumptive evidence of accelerated phase in the absence of any other criteria of accelerated phase.¹²

The objective of this study was to determine frequency of megakaryocytic clustering in patients of CML in chronic phase at the time of presentation and comparison of response to treatment with Imatinib in patients with or without megakaryocytic clustering.

METHODOLOGY

It was a retrospective study conducted at Mayo Hospital, Lahore in collaboration with CML Clinic Mayo Hospital and Hematology Section of Pathology Department, King Edward Medical University, from March 2018 to March 2019. All the diagnosed patients of CML in chronic phase with positive BCR/ABL fusion gene were included in the study regardless of the gender and age, and who were taking treatment with first line drugs and were compliant with a regular follow-up. All those diagnosed patients of CML with some additional chromosomal mutation or with JAK2 positivity, were excluded from the study.

After obtaining permission from University Ethical Review Board, all the data regarding personal information, medical history, physical examination, and investigations of included patients were recorded. The complete blood count for hemoglobin levels, total leucocyte count, differential leucocytic count, and platelet count done at the time of presentation, were reviewed. Findings of abdominal ultrasound/CT abdomen, for the presence of splenomegaly and size of spleen as per patient's record were also recorded after keeping full confidentiality.

Bone marrow trephine biopsy stained with hematoxylin and eosin done for diagnosis of disease, were re-examined for the presence of megakaryocytic clustering by two independent observers to minimise inter-observer bias. Megakaryocytic clustering is said to be present, if three or more clusters of five or more megakaryocytes are seen in an area of 1 cm.

Treatment and follow-up response of all the patients were retrieved. The initial therapy in all patients was tablet Imatinib 400mg. After three months, the patients were followed up for CHR. As per institutional policy, assessment for CHR is done by complete blood count, peripheral smear, and ultrasound/CT abdomen. Those who failed to achieve CHR were prescribed an increased dose of 800 mg of Imatinib. At 6 months and one year, all the patients were assessed for major molecular response. Ideally, cytogenetic analysis is to be done at 6 months and molecular analysis at 1 year as per the guidelines; but due to economic constraints, only molecular analysis is done both at 6 months and 1 year as per institutional policy.

The results were recorded and data were analysed using SPSS V 25. Continuous variable like age, Hb, TLC, spleen size, were described in terms of minimum and maximum values with mean $\pm$ S.D and categorical variables like gender, were presented as frequencies.

Chi-square test was applied to determine correlation among qualitative variables and a p-value of < 0.05 was considered statistically significant.

RESULTS

A total of 94 patients of CML in CP were identified during the defined study period of 6 months. There were 51 (54.3%) male patients and 43 (45.7%) female patients with a mean age of 47.02 ± 9.3 years. Other characteristic variables are shown in Table I.

The bone marrow trephine biopsy slides of all the patients were evaluated for megakaryocytic clustering. It was present in 57 (60.6%) of the patients. Bone Marrow trephine biopsy figures without megakaryocytic clustering and with megakaryocytic clustering are shown below (Fig 1a and 1b, respectively).

The sokal score of each patient was calculated according to the formula mentioned above. A total of 18 (19.1%) patients had high sokal score, 33 (35.1%) had intermediate sokal score and 43 (45.7%) had low sokal score. The correlation of megakaryocytic clustering and high sokal score was found to be statistically significant ($p < 0.001$, Table II).

The percentage of patients with megakaryocytic clustering and absent CHR, MMR at 6 months and 1 year is illustrated in Figure 2.

All the patients were followed up to assess complete hematological response at the end of three months and MMR after 6 months and 1 year. The cytogenetic response is not done owing to institutional policy. It was observed that CHR was achieved in 81 (86.2%) and absent in 13 (13.8%) patients; out of which, 12 (92.3%) patients had megakaryocytic clustering (p -value = 0.012). Similarly, MMR at 6 months was absent in 23 (24.5%) patients. Out of them, megakaryocytic clustering was observed in 21 (91.3%, p -value = 0.001). At 12 months, MMR was absent in 27 (28.7%) patients, out of which 25 (92.6%) (p -value < 0.001) had megakaryocytic clustering (Table II).

Table I: Demographic and hematological variables of CML-CP patients.

Variables	Mean $\pm$ S.D.	Minimum	Maximum
Age (years)	47.02 $\pm$ 9.3	28	67
Hb (g/dl)	9.97 $\pm$ 1.8	5.5	13.1
TLC (10^3 uL)	126.34 $\pm$ 42.1	34	212
Platelets (10^9 uL)	386.95 $\pm$ 161.7	112	890
Blast count (%)	3.80 $\pm$ 4.48	1	9
Spleen size (cm)	15.22 $\pm$ 1.8	9	23

Table II: Relationship of megakaryocytic clustering with gender, sokal score, CHR, MMR at 6 months and 1 year.

		Gender		Sokal			CHR		MMR at 6 months		MMR at 12 months	
		Female N = 43	Male N = 51	High N = 18	Intermediate N = 33	Low N = 43	Absent N = 13	Present N = 81	Absent N = 23	Present N = 71	Absent N = 27	Present N = 67
Megakaryocytic clusters	Absent	19 (44.2%)	18 (35.3%)	1 (5.6%)	9 (27.3%)	27 (62.8%)	1 (7.7%)	36 (44.4%)	2 (8.7%)	35 (49.3%)	2 (7.4%)	35 (52.2%)
	Present	24 (55.8%)	33 (64.7%)	17 (94.4%)	24 (72.7%)	16 (37.2%)	12 (92.3%)	45 (55.6%)	21 (91.3%)	36 (50.7%)	25 (92.6%)	32 (47.8%)
	p-values	0.379		<0.001			0.012		0.001		<0.001

Figure 1a: CML chronic phase without bone marrow clustering. **Figure 1b: CML chronic phase with bone marrow clustering.**

DISCUSSION

CML involves malignant transformation of multipotent progenitor cells including granulocytes and megakaryocytes, erythroid cells and macrophages at a variable extent. In this study, 94 patients of CML in chronic phase were included. Mean age of the patient was 47.02 $\pm$ 9.3 years with male predominance. Average hemoglobin level was found to be 9.9, and anemia was seen in 47 patients, mean platelet count was 386,000/uL, and was found to be normal or slightly on the lower side. Mean WBC count was 126,000/uL. These findings are in accordance with study by Hamid *et al.*¹³

Average spleen size was 15 cm and splenomegaly was seen in 23 patients. In a study by Kumar, no significant splenomegaly was found among the patients and they also found platelet count to be slightly higher in the patients having CML.¹⁴

Sokal Index for CML predicts survival based on clinical and laboratory investigation, done before the commencement of treatment. Sokal score was high in 18 (19.1%) patients, intermediate in 33 (35.1%), and low in 43 (45.7%) patients. Similar results were found in the study conducted by Laskshmaiah in India.¹⁵

Megakaryocytes were the main focus of this study. The megakaryocytic number and clustering was closely observed in the trephine biopsy slides. The megakaryocytic number was raised in 57 (60.6%) of cases observed. Megakaryocytic clustering was seen in all these patients; whereas, dwarf forms were seen in 20% of the cases.

Figure 2: Relationship of poor treatment response with megakaryocytic clustering.

Arun *et al.* has studied the number and morphologic features of megakaryocytes in CML patients. The number was raised in 67% patients and 32% patients had megakaryocytic clustering. They divided patients according to classification of CML by Burkhardt into CML-GM (granulocytic megakaryocytic) and CML-G (granulocytic). Increased number and clustering was seen exclusively in CML-GM group.¹⁶

All patients were followed for at least 1 year and treatment response was noted. Good response in the form of complete CHR, MMR1 and MMR2 was seen in 57 patients and bad response in the form of failure of either one or all CHR, MMR 1 and MMR 2 was seen in 37 patients.

The cytogenetic response is not done owing to institutional criteria. However for the correlation of responses as shown by Jabbour *et al.* in updates of CML, instead of repeated bone marrow biopsy, a complete CCyR is equal to negative FISH test and BCR-ABL <1% and partial CCyR is equal to BCR-ABL <10%.¹⁷ It was observed that CHR was achieved in 81 (86.2%) and absent in 13 (13.8%) patients, out of which 12 (92.3%) patients has megakaryocytic clustering ($p = 0.012$). In a study by Hagop Kartarijan, CHR was observed in 89% of the patients.¹⁸

Similarly, MMR at 6 and 12 months was absent in 23 (24.5%) and 27 (28.7%) patients in which megakaryocytic clustering was observed in 21 (91.3%, $p = 0.001$) and 25 (96.2%, $p < 0.001$) patients.

The correlation of sokal score with megakaryocytic clustering was done, which showed that nearly all the patients with high sokal score had megakaryocytic clustering seen in their trephine biopsy. Those with intermediate sokal score, 72% of those had clustering, showing a significant correlation. Similar results were seen in a study by Theil *et al.*, in which increased CD 61 positive megakaryocytic cells were associated with increased tumor burden and advanced stage of disease.¹⁹

Similar correlation was seen in a study by Forrest *et al.*, in which CHR and MMR correlated with the sokal score.²⁰ Nachi *et al.* has conducted study on effect of gender on treatment

outcome. In that study, females were found to have far better outcome (MMR 80 vs. 45%). Similar results were found in our study as males had more megakaryocytic clustering than females. Bad response was seen in 33 males and 24 females and good response was seen in 18 males and 19 females.²¹

CONCLUSION

Patients with megakaryocytic clustering have poor clinical outcome as indicated by their sokal score, absent complete hematological response (CHR), molecular response at 3 months and 1 year. Trephine biopsy of all the patients with chronic CML should be assessed for megakaryocytic clustering, as it is presumptive of poor prognosis in CML patients.

ETHICAL APPROVAL:

Ethical approval of this study was obtained from King Edward Medical University Ethics Committee prior to initiation of the research work.

PATIENTS' CONSENT:

Informed consents were obtained from all patients to publish the data concerning this study.

CONFLICT OF INTEREST:

Authors declared no conflict of interest.

AUTHORS' CONTRIBUTION:

ZM, HB: Data acquisition, analysis and drafting of the study, agreement to be accountable for all aspects of the work in ensuring that questions related to the accuracy or integrity of any part of the work are appropriately investigated and resolved.

SA, AH, AK: Conception and design of the study, review and final approval of the draft, agreement to be accountable for all aspects of the work in ensuring that questions related to the accuracy or integrity of any part of the work are appropriately investigated and resolved.

SQ: Drafting of the study, data analysis and interpretation, agreement to be accountable for all aspects of the work in ensuring that questions related to the accuracy or integrity of any part of the work are appropriately investigated and resolved.

REFERENCES

1. Zhang B, Li L, Ho Y, Li M, Marcucci G, Tong W, *et al.* Heterogeneity of leukemia-initiating capacity of chronic myelogenous leukemia stem cells. *J clinical Investigation* 2016; **126**(3):975-91. doi: 10.1172/JCI79196.
2. Kansal R. Acute myeloid leukemia. Tumors and cancers: Endocrine glands-blood-marrow-lymph. 2017.
3. Karkuzhali P, Shanthi V, Usha T. A case of chronic myeloid leukemia presenting as megakaryocytic blast crisis (AML M7). *Ecanermedicalscience* 2013; **7**:375. doi: 10.3332/ecanecr.2013.375.

4. Kumar S, Gupta VK, Bharti A, Meena LP, Gupta V, Shukla J. A study to determine the clinical, hematological, cytogenetic, and molecular profile in CML patient in and around Eastern UP, India. *J family Medicine Primary Care* 2019; **8(7)**:2450. doi: 10.4103/jfmpc.jfmpc_307_19.
5. Khonglah Y, Basu D, Dutta TK. Bone marrow trephine biopsy findings in chronic myeloid leukemia. *Malaysian J Pathology* 2002; **24(1)**:37-43.
6. Usman M, Syed NN, Kakepoto GN, Adil SN, Khurshid M. Chronic phase chronic myeloid leukemia: response of Imatinib mesylate and significance of Sokal score, age and disease duration in predicting the hematological and cytogenetic response. *JAPI* 2007; **55**:103-07.
7. Baccarani M, Pane F, Saglio G. Monitoring treatment of chronic myeloid leukemia. *Haematologica* 2008; **93(2)**:161-9. doi: 10.3324/haematol.12588.
8. Saussele S, Hehlmann R, Fabarius A, Jeromin S, Proetel U, Rinaldetti S, et al. Defining therapy goals for major molecular remission in chronic myeloid leukemia: results of the randomized CML Study IV. *Leukemia* 2018; **32(5)**:1222-8.
9. Bhatia R. Novel approaches to therapy in CML. *Hematology Am Soc Hematol Educ Program* 2017; **2017(1)**:115-20. doi: 10.1182/asheducation-2017.1.115.
10. Goldman JM. Treatment strategies for CML. *Best Practice & Research Clinical Haematol* 2009; **22(3)**:303-13.
11. Rosti G, Martinelli G, Bassi S, Amabile M, Trabacchi E, Gianini B, et al. Molecular response to imatinib in late chronic-phase chronic myeloid leukemia. *Blood* 2004; **103(6)**:2284-90.
12. Shaffer LG. Editors. An International System for Human Cytogenetic Nomenclature. 2013.
13. Hamid A, Ashraf S, Qamar S, Naveed MA, Hameed A, Farooq MA. Myelofibrosis in patients of chronic myeloid leukemia in chronic phase at presentation. *J Coll Phys Surg Pakistan* 2019; **29(11)**:1096-100. doi: 10.29271/jcpsp.2019.11.1096.
14. Arunachalam AK, Jain M, Kumar A, Kushwaha R, Singh US, Tripathi AK. Megakaryocytes in chronic phase of chronic myeloid leukemia: A descriptive case series. *Annals Pathol Labor Med* 2016; **3(3)**:A176-182.
15. Kuntegowdanahalli LC, Kanakasetty GB, Thanky AH, Dasappa L, Jacob LA, Mallekavu SB, et al. Prognostic and predictive implications of Sokal, Euro and EUTOS scores in chronic myeloid leukaemia in the imatinib era—experience from a tertiary oncology centre in Southern India. *Ecan-cermedicalscience* 2016; **10**.
16. Burkhardt R, Bartl R, Jäger K, Frisch B, Kettner G, Mahl G, et al. Working classification of chronic myeloproliferative disorders based on histological, haematological, and clinical findings. *J Clin Pathol* 1986; **39(3)**:237-52. doi: 10.1136/jcp.39.3.237.
17. Jabbour E, Kantarjian H. Chronic myeloid leukemia: 2018 update on diagnosis, therapy and monitoring. *Am J Hematol* 2018; **93(3)**:442-59. doi: 10.1002/ajh.25011.
18. Cortes J, Talpaz M, O'Brien S, Jones D, Luthra R, Shan J, et al. Molecular responses in patients with chronic myelogenous leukemia in chronic phase treated with imatinib mesylate. *Clin Cancer Res* 2005; **11(9)**:3425-32. doi: 10.1158/1078-0432.CCR-04-2139.
19. Thiele J, Kvasnicka HM, Fischer R, Diehl V. Clinicopathological impact of the interaction between megakaryocytes and myeloid stroma in chronic myeloproliferative disorders: a concise update. *Leukemia & Lymphoma* 1997; **24(5-6)**:463-81. doi: 10.3109/10428199709055584.
20. Forrest DL, Trainor S, Brinkman RR, Barnett MJ, Hogge DE, Nevill TJ, et al. Cytogenetic and molecular responses to standard-dose imatinib in chronic myeloid leukemia are correlated with Sokal risk scores and duration of therapy but not trough imatinib plasma levels. *Leukemia research* 2009; **33(2)**:271-5. doi: 10.1016/j.leukres.2008.07.015.
21. Nachi M, Kihel I, Guella D, Dali-Ali A, Abed A, Boukhatmi Y. Sex and major molecular response to imatinib treatment for patients with chronic myeloid leukemia. *Biochem Pharmacol (Los Angel)* 2019; **8(263)**:2167-0501. doi: 10.4172/2167-0501.1000263.

• • • • •