



RESEARCH ARTICLE

Bone Marrow Examination in Pancytopenia: Patterns, Causes, and Clinical Correlation in Indian Patients

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ABSTRACT

Background: Pancytopenia is a hematological condition characterized by reduction in all three cellular components of blood and often indicates underlying bone marrow pathology. Bone marrow examination remains a cornerstone in diagnosis.

Objective: To evaluate bone marrow findings in pancytopenia and correlate with clinical and hematological parameters in Indian patients.

Methods: This retrospective study included 300 patients diagnosed with pancytopenia between October 2023 and September 2025 at the Department of Pathology, M.K.C.G Medical College, Berhampur, Odisha. Clinical data, hematological parameters, and bone marrow findings were analyzed. Statistical analysis was performed using SPSS.

Results: Megaloblastic anemia was the most common cause (36%), followed by aplastic anemia (22%), acute leukemia (14%), and hypersplenism (10%). Significant associations were observed between etiology and hemoglobin levels and mean corpuscular volume.

Conclusion: Bone marrow examination is essential for accurate diagnosis of pancytopenia. Nutritional causes remain predominant in Indian settings, highlighting the need for early intervention.

Keywords: Pancytopenia, Bone marrow aspiration, Megaloblastic anemia, Aplastic anemia, Hematological parameters.

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INTRODUCTION

Pancytopenia is defined as a reduction in hemoglobin concentration, leukocyte count, and platelet count below normal reference ranges and represents a manifestation of diverse underlying pathological processes rather than a disease entity itself (1). The etiological spectrum ranges from reversible nutritional deficiencies to life-threatening hematological malignancies (2).

In developing countries such as India, nutritional anemia—particularly vitamin B12 and folate deficiency—remains a leading cause, whereas in developed nations, bone marrow failure syndromes and malignancies are more prevalent (3,4). Bone marrow examination plays a pivotal role in establishing diagnosis, especially when peripheral smear findings are inconclusive (5).

The pathophysiology of pancytopenia involves decreased production, increased destruction, or sequestration of blood cells (6). Bone marrow aspiration and biopsy provide valuable insight into cellular morphology, marrow cellularity, and infiltration patterns (7).

Previous studies have demonstrated varied etiological patterns depending on geographic and socioeconomic factors (8–10). Understanding these patterns is essential for timely diagnosis and management.

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This study aims to analyze bone marrow findings in pancytopenia and correlate them with clinical and hematological parameters in an Indian tertiary care setting.

MATERIALS AND METHODS

Study Design

Retrospective observational study.

Study Setting

Department of Pathology, M.K.C.G Medical College and Hospital, Berhampur, Odisha.

Study Duration

October 2023 to September 2025.

Sample Size

300 cases.

Inclusion Criteria

- Patients diagnosed with pancytopenia
- All age groups

Exclusion Criteria

- Incomplete records
- Prior chemotherapy

STATISTICAL ANALYSIS

Data analyzed using SPSS v25. Chi-square test applied. $p < 0.05$ considered significant.

RESULTS

A total of 300 patients diagnosed with pancytopenia were included in the study.

Demographic Profile

The majority of cases were in the 21–40 years age group (40%). There was a slight male predominance in the study group.

Table 1 summarizes the age-wise distribution of cases.

As illustrated in Figure 1, the majority of patients belonged to the young adult population.

Etiological Distribution

Bone marrow examination revealed that megaloblastic anemia was the most common etiology (36%). The distribution of etiological causes is shown in Table 2.

Table 1: Age-wise distribution of pancytopenia cases (n = 300)

Age group (years)	Number of cases	Percentage (%)
<20	60	20%
21–40	120	40%
41–60	75	25%
>60	45	15%

Table 2: Etiological distribution of pancytopenia (n = 300)

Etiology	Number of cases	Percentage (%)
Megaloblastic anemia	108	36%
Aplastic anemia	66	22%
Acute leukemia	42	14%
Hypersplenism	30	10%
Others	54	18%

Table 3: Hematological parameters in pancytopenia patients

Parameter	Mean \pm SD
Hemoglobin (g/dL)	6.8 \pm 1.5
TLC ($\times 10^9/L$)	2.5 \pm 0.8
Platelet count ($\times 10^9/L$)	75 \pm 30
MCV (fL)	102 \pm 12

A graphical representation of etiological distribution is shown in Figure 2.

Hematological Parameters

The mean hematological values of the study population are summarized in Table 3.

Patients with megaloblastic anemia showed significantly higher MCV values compared to other groups.

Bone Marrow Findings

Bone marrow examination revealed:

- Hypercellular marrow in megaloblastic anemia cases
- Hypocellular marrow in aplastic anemia
- Marrow infiltration/blasts in leukemia

These findings are summarized in Table 4.

Statistical Analysis

Statistical analysis showed significant associations between etiology and hematological parameters:

- Hemoglobin levels varied significantly across etiologies ($p < 0.001$)
- Mean corpuscular volume (MCV) showed strong association with megaloblastic anemia ($p < 0.001$)
- Platelet count differences were statistically significant ($p = 0.02$)

Chi-square analysis demonstrated a significant association between bone marrow diagnosis and MCV category ($\chi^2 = 32.6$, $df = 4$, $p < 0.001$).

DISCUSSION

The present study highlights megaloblastic anemia as the most common cause of pancytopenia, consistent with findings from other Indian studies (11–13). Nutritional deficiencies remain prevalent due to dietary habits and socioeconomic factors (14).

Table 4: Bone marrow cellularity patterns

Diagnosis	Cellularity pattern	Key morphological features
Megaloblastic anemia	Hypercellular	Megaloblastic erythropoiesis, nuclear-cytoplasmic asynchrony
Aplastic anemia	Hypocellular	Marked reduction in hematopoietic cells, fatty marrow
Acute leukemia	Hypercellular	Presence of blasts, suppressed normal hematopoiesis
Hypersplenism	Normocellular	Normal marrow with peripheral destruction

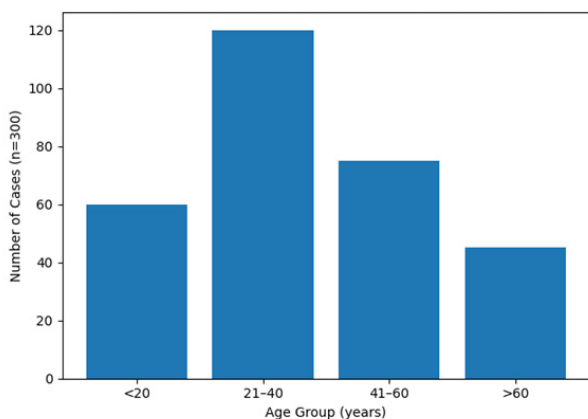


Figure 1: Age-wise distribution of pancytopenia cases

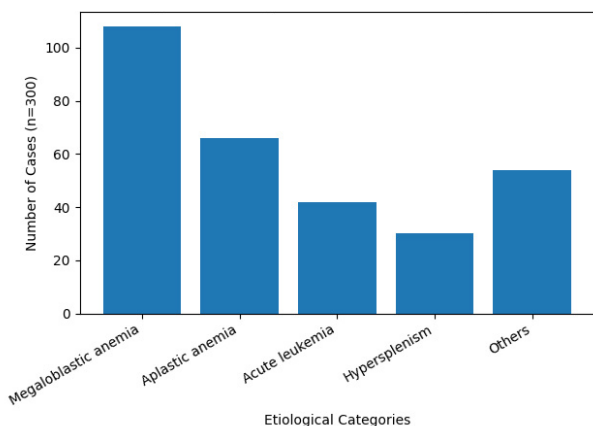


Figure 2: Etiological distribution of pancytopenia

Aplastic anemia was the second most common cause, aligning with previous reports (15,16). Bone marrow examination revealed hypocellularity in these cases, confirming diagnosis.

Acute leukemia constituted a significant proportion, emphasizing the importance of early bone marrow evaluation (17). Morphological analysis remains crucial in resource-limited settings (18).

Statistically significant correlation between MCV and megaloblastic anemia supports its diagnostic utility (19). Similarly, thrombocytopenia severity varied significantly across etiologies (20).

Comparisons with global studies reveal differences in etiological patterns, highlighting regional variability (21–23). Early diagnosis through bone marrow examination can significantly improve outcomes (24,25).

CONCLUSION

Bone marrow examination is indispensable in evaluating pancytopenia. Megaloblastic anemia remains the leading cause in Indian patients. Early diagnosis and intervention can reduce morbidity and mortality.

REFERENCES

1. Young NS. Aplastic anemia. *N Engl J Med.* 2018;379(17):1643–1656.
2. Tefferi A. Pancytopenia: a clinical approach. *Mayo Clin Proc.* 2003;78(10):1274–1280.
3. Khunger JM, Arculselvi S, Sharma U, Ranga S, Talib VH. Pancytopenia—a clinico-hematological study of 200 cases. *J Assoc Physicians India.* 2002;50:518–522.
4. Tilak V, Jain R. Pancytopenia—a clinico-hematologic analysis of 77 cases. *Indian J Pathol Microbiol.* 1999;42(4):399–404.
5. Bain BJ. Bone marrow aspiration. *J Clin Pathol.* 2001;54(9):657–663.
6. McKenzie SB, Williams JL. *Clinical Laboratory Hematology.* 3rd ed. Pearson; 2015.
7. Lewis SM, Bain BJ, Bates I. Dacie and Lewis Practical Haematology. 12th ed. Elsevier; 2017.
8. Varma N, Dash S. A reappraisal of underlying pathology in adult pancytopenia. *Trop Geogr Med.* 1992;44(4):322–327.
9. Kumar R, Kalra SP, Kumar H, Anand AC, Madan H. Pancytopenia—a six year study. *J Assoc Physicians India.* 2001;49:1078–1081.
10. Jain A, Naniwadekar M. An etiological reappraisal of pancytopenia—largest series reported to date from a single tertiary care teaching hospital. *BMC Hematol.* 2013;13(1):10.
11. Gayathri BN, Rao KS. Pancytopenia: a clinico-hematological study. *J Lab Physicians.* 2011;3(1):15–20.
12. Niazi M, Raziq F. The incidence of underlying pathology in pancytopenia—an experience of 89 cases. *J Postgrad Med Inst.* 2004;18(1):76–79.

13. Jha A, Sayami G, Adhikari RC, Panta AD, Jha R. Bone marrow examination in cases of pancytopenia. *J Nepal Med Assoc.* 2008;47(169):12–17.
14. O’Leary F, Samman S. Vitamin B12 in health and disease. *Nutrients.* 2010;2(3):299–316.
15. Marsh JCW, Ball SE, Cavenagh J, Darbyshire P, Dokal I, Gordon-Smith EC. Guidelines for the diagnosis and management of aplastic anemia. *Br J Haematol.* 2009;147(1):43–70.
16. Camitta BM, Thomas ED, Nathan DG, Gale RP, Kopecky KJ, Rapoport JM. A prospective study of androgens and bone marrow transplantation for treatment of severe aplastic anemia. *Blood.* 1979;53(3):504–514.
17. Hoffbrand AV, Moss PAH. *Essential Haematology.* 7th ed. Wiley-Blackwell; 2016.
18. Greer JP, Arber DA, Glader B, List AF, Means RT, Paraskevas F. *Wintrobe’s Clinical Hematology.* 14th ed. Wolters Kluwer; 2018.
19. Hoffbrand AV, Higgs DR, Keeling DM, Mehta AB. *Postgraduate Haematology.* 7th ed. Wiley-Blackwell; 2016.
20. Lee GR, Foerster J, Lukens J, Paraskevas F, Greer JP, Rodgers GM. *Wintrobe’s Clinical Hematology.* 10th ed. Lippincott Williams & Wilkins; 1999.
21. Kar M, Ghosh A. Pancytopenia: correlation of clinical findings with bone marrow morphology. *J Pathol Nepal.* 2012;2(4):265–269.
22. Kumar DB, Raghupathi AR. Clinico-hematologic analysis of pancytopenia: study in a tertiary care center. *Int J Health Sci Res.* 2014;4(9):15–20.
23. Dasgupta S, Mandal PK, Chakrabarti S. Etiological profile of pancytopenia in adults. *Int J Med Sci Public Health.* 2015;4(9):1309–1313.
24. Lichtman MA, Kaushansky K, Prchal JT, Levi MM, Burns LJ, Linch DC. *Williams Hematology.* 9th ed. McGraw-Hill; 2016.
25. Kaushansky K, Lichtman MA, Prchal JT, Levi MM, Press OW, Burns LJ. *Williams Hematology.* 10th ed. McGraw-Hill; 2021.