Comparison of Bone Marrow Abnormalities of Two HIV/AIDS Patient Populations

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Background

•Bone marrow aspirate and biopsy from human immunodeficiency virus (HIV)/acquired immune deficiency syndrome (AIDS) patients have been known to exhibit various abnormalities including trilineage dysplasia and plasmacytosis. However, quantitative data on the degree of abnormalities is Further, no studies have sparse. reported on the variation observed among different HIV/AIDS populations.

Design

•Under IRB approval, a retrospective chart review was conducted on all marrows performed on bone HIV/AIDS patients from 2009 to 2011 at two hospitals. Hospital 'A' serves a population without ready access to care in an HIV-prevalent area while Hospital 'B' serves a population opposite to this. Data collected include demographics, CD4 T-cell counts, and bone marrow findings including erythrocyte and dysplasia, megakaryocyte and plasma cell count. Plasma cell percentage of 3 or more was Exclusion considered increased. criteria were patients with active hematologic neoplasms. primary The Fisher's exact test and Mann-Whitney U test were employed for statistical analysis.

	Hospital A patients (n=21)	Hospital B patients (n=19)		
Age (mean, in years)	32.3	38.9		
Gender				
Male	12	13		
Female	9	6		
Race				
Caucasian	1	7		
African American	16	10		
Hispanic	4	1		
Other	0	1		
CD4 count (mean absolute counts)	72.7	77.9		
Biopsy findings				
Plasma cell count [*] (mean %, % patients with increase)	8.6%, 76.2%	4%, 52.6%		
Erythrocyte dysplasia (number of patients, %)	10 (47.8%)	11 (57.9%)		
Megakaryocyte dysplasia (number of patients, %)	11 (52.4%)	11 (57.9%)		

*The only statistically significant difference observed between the two hospitals is the degree of plasmacytosis

There is a statistically significant difference in t populations

•There are no significant differences in:

- Number of patients with erythrocyte and megakaryocyte dysplasia
- Age of patients
- CD4 counts
- Number of patients with increased plasma cell numbers

•The difference in plasmacytosis at Hospital 'A' versus Hospital 'B' without corresponding differences in cellular dysplasia suggests that these bone marrow abnormalities may be unrelated in pathogenesis. Indeed, the plasmacytosis associated with HIV could be related to an extrinsic factor such as lack of medication compliance or other factors related to inaccessibility to medical care. Further studies would be needed to consider the degree of plasmacytosis as a marker in monitoring HIV patients.

Results

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Conclusions



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