

Introduction

- Developed countries, where guidelines on the treatment of PJP have been created, have dramatically lower rates of latent and active TB than developing countries.
- PJP may delay the diagnosis and treatment of TB.
- Furthermore, treatment of PJP with corticosteroids may be detrimental to the course of TB.
- The objective of this study was to examine the frequency and the clinical characteristics of the co-infection of PJP and TB in HIV.

Methods

- The clinical details of all HIV patients being treated at Amrita Hospital in South India have been prospectively collected in an electronic database since 2006.
- We compiled the data from 2006 to 2018 and further examined the clinical and laboratory results from electronic charts of patients admitted with PJP.
- Statistical analysis of the data was performed with descriptive analysis.

Results

- A total of 21 of the 576 HIV patients had admissions for PJP.
- Of these 9/21 (43%) were co-infected with PJP and TB.
- In all cases PJP presentation was the event leading to the diagnosis of HIV.
- The mean age and gender ratios were similar in our total population and our PJP subgroup however, the CD-4 count of the PJP group was significantly lower (50 vs 237 cells/ul p=0.004). All of our PJP cases were below the classical 200/ul threshold for developing PJP.
- Seven of the nine PJP-TB co-infected patients had been started on antitubercular therapy in the community prior to diagnosis of HIV.
- There were no significant differences between the PJP only and the PJP-TB group.

Table 1. Comparison of PJP group to all HIV patients

Parameter (SD)	All HIV patients	PJP group
Age	43 years (10)	43 years (5.7)
Gender	64% Male 36%	71% Male 29%
	Female	Female
CD-4 count at	237 cells/ul (265)	50 cells/ul (40)
presentation	Range (2-1874)	Range (2-135)
Number	576	21

Pneumocystis-Tuberculosis Co-infection in the HIV Positive Host

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The occurrence and co-occurance of opportunistic infections in HIV positive patients is dependent on numerous factors. In a South Indian population we demonstrate a higher than previously reported rate of TB in patients presenting with pneumocystis pneumonia. Further study is needed of this entity, particularly the effect of steroids in this population.

Table 2. Comparison of TB-PJP group to PJP only

Parameter	TB Positive	TB Negative
	Mean (SD) N	Mean (SD) N
Age	40 years (5.1) 9	45 years (5.3) 12
Male:Female	8:1	7:5
CD-4 Count	49.7 (40.5) 9	50.9 (40.9) 12
Length of Stay	16.5 Days (17.5) 9	11.8 Days (6.2) 11
ICU Admission Rate	33% 9	36% 11
Hypoxia (SpO2 =88%)Rate</th <th>88% 8</th> <th>55% 11</th>	88% 8	55% 11
Expired	0/9 9	3/12 11
WBC count/ul	8116 (3494) 9	7004 (3685) 11
Lymphocyte count/ul	1017(689) 9	1131 (402) 11
Serum Sodium mEq/L	128.6 (3.8) 8	130.7 (4.6) 10
Albumin:Globulin Ratio	0.67 (0.18) 9	0.67 (0.17) 11



Table 3. Previously reported rates of TB-PJP co-infection in patients presenting with PJP pneumonia in different populations. Study

Barnes 1 (Los Ang Castro 2 (Miami) **Orlovic** 2 (South A

Udaiwad 2009 (Mu **Rani 201** (Chenna) **Our Data**

Abbreviations; HIV, Human Immunodeficiency Virus; PJP, Pneumocystis Jiroveci Pneumonia; TB, Tuberculosis



Results

	TB-PJP prevalence	National HIV prevalence	National TB prevalence
992 (eles)	4%	0.6%	10/100,000
000	10%	0.4%	5/100,000
2007 frica)	9%	0.7%	20/100,000
ia ımbai)	10%	0.3%	40/100,000
0 i)	40%	0.3%	40/100,000
a	43%	0.26%	40/100,000

Conclusion

Our report highlights the need to increase awareness of occurrence of this dual infection in HIV infected patients, as both infections can mimic each other clinically and radiologically

In addition we report a much higher rate of underlying TB in our patients with PJP than what has been generally reported in the literature

Further research is needed to determine the risk benefit ratio of steroids in these populations.

We conclude that higher clinical suspicion for this entity is warranted, particularly in countries with a high prevalence of TB.

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