INTRODUCTION

Government of India launched the free National ART program on 1 st April 2004. As on March 2013, there are around 18.13 lakhs people living with HIV (PLWHA) in India. Currently near 6.5 lakhs are on first-line ART. India is estimated to have around 1.16 lakhs annual new HIV infections among adults. The high prevalence states account for only 31% of new infections, while the ten low prevalence states of Odisha, Jharkand, Bihar, Uttar Pradesh, West Bengal, Gujarat, Chattisgarh, Rajasthan, Punjab & Uttarakhand together account for 57% of new infections. The high prevalence states in South India (Andhra Pradesh, Karnataka, Maharashtra and Tamil Nadu) account for 53% of all HIV infected population in the country [1].

The introduction of HAART (Highly Active Anti-Retroviral Therapy) has led to an increase in survival among HIV-infected patients, decreased HIV-associated mortality, and improved quality of life among HIV patients. In India, under the National Aids control program (NACO), the generic fixed drug combination of zidovudine (or) Stavudine, Lamivudine and nevirapine (or) efavirenz is being used. Despite the reduction in morbidity and mortality, a considerable proportion of patients fail to achieve a sustained virological response to therapy [2].

In the resource limited setting like India, where the cost of treatment is very high and where routine virological monitoring and genotyping resistance is not done to start the therapy and see the response to therapy, there is a need for parameters to stratify the patients to different stages of risk of failure, which will be useful for the clinicians. In routine clinical practice, where monitoring of ART adherence, investigations to assess the failure, and clinical follow-up are generally less rigorous, failure rates are often higher [2].

India has the second largest burden in the world after South Africa. There are few data on the incidence and risk factors for treatment failure associated with generic HAART regimens in India. Treatment failure is an increasing concern [3].
Since 2004 with extension of ART program to all over India, increased number of HIV patients have got access to treatment. With increased duration of treatment years as well as quantum of patients, we expect some failure rates to first-line ART. Failure rates to first-line ART may vary region wise and need not be uniform all over the country due to various identified reasons.

To our knowledge, ours is the largest retrospective cross-sectional study in India on the prevalence of failure of first-line ART, over one decade, covering six districts of combined Andhra Pradesh. Our objectives were a) to study the prevalence of failure of first-line ART and b) to study the statistical significance of association of first-line ART failure with duration of ART, gender, and base line CD4 count.

**MATERIALS AND METHODS**

Our study was a retrospective cross-sectional study from April 2004 – March 2014, a period of 10 years. This study was done at ART-Plus centre, established in Government Siddhartha Medical College and Government General Hospital, a tertiary level teaching hospital at Vijayawada, Andhra Pradesh, India. This study was approved by institutional ethics committee. This ART-Plus centre is a referral centre and serves six districts, namely Krishna, Guntur, West Godavari, Nellore, Prakasam and Khammam districts of Andhra Pradesh (before state bifurcation), in relation to confirmation of first-line ART failure and provision of second-line ART (Fig.1). This centre follows national guidelines for treatment of HIV patients.

**Study population** included PLWHA, registered for first-line ART in the 16 nodal ART centres, from 6 districts. Study duration was a decade spanning from April 2004-March 2014. Among study population males were 27,797 and females were 29,877.

**Inclusion criteria**
- Subjects of > 15 years of age, on first-line ART
- Subjects on first-line ART, with at least one follow-up visit at 6 months
- Patients switched over to second-line as per NACO guidelines.

**Exclusion criteria**
- Children of <15 years age, who are on ART
- Pregnant and lactating women
- Subjects on alternate first-line ART

We calculated the prevalence of failure of first-line ART. We also assessed the statistical significance of association of first-line ART failure with a) duration of ART, b) gender and c) base line CD4 count. The data was collected retrospectively from the electronic health record (EHR) of the ART-Plus centre, Vijayawada. The EHR contains complete information of each patient, including demographics, HIV risk behaviors, adherence to ART, medications, which are entered as a part of routine clinical documentation and which is updated every day. Patients, whose complete record was not available and patients who could not be traced, were excluded from the study.

**Treatment Failure**

Treatment failure was assessed using definitions of ART failure given by WHO [4]. Clinical failure was defined as new or recurrent WHO stage 4 condition, after at least 6 months of ART. Immunological failure was defined as fall of CD4 counts to pre-therapy baseline or below; 50% fall from the on-treatment peak value; and persistent CD4 levels below 100 cells/mm³.

Virological failure was defined as plasma viral load >5,000 copies /ml, after at least 6 months of Art.

**Statistical analysis**

The data was analysed by using MS excel. The data were presented by mean ± standard deviation for continuous variables. Chi-square test and P values were used to assess the statistical significance of the findings.
Table 1: Baseline characteristics of patients

<table>
<thead>
<tr>
<th>Variable</th>
<th>Value</th>
</tr>
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<tbody>
<tr>
<td>Males</td>
<td>27,797</td>
</tr>
<tr>
<td>Females</td>
<td>29,877</td>
</tr>
<tr>
<td>Mean baseline CD4 count ± SD (months)</td>
<td>191.63±181.86</td>
</tr>
<tr>
<td>Mean duration of ART±SD (months)</td>
<td>53±23.44</td>
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</tbody>
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RESULTS

Out of total 57,674 individuals, 244 had failure, with prevalence of failure of first-line ART 0.47%. There was no significant association of duration of ART with first-line ART failure with \( \chi^2 = 0.0069; P > 0.05 \) (Table 2).

Males had higher incidence of first-line ART failure than females with statistically significant values, \( \chi^2 = 84.03 \) and \( p < 0.05 \) (Table 3).

First-line ART failure was found in 180 subjects with less than mean CD4 count and in 64 subjects with more than mean CD4 count. There was significant association between low mean CD4 count and first-line ART failure with \( \chi^2 = 15.48 \) and \( p < 0.05 \) (Table 4).

DISCUSSION

To our knowledge ours is the ever largest study on the prevalence of failure of first-line ART, conducted in combined Andhra Pradesh (before bifurcation). Major problem of long-run programs is treatment failure. Prevalence of treatment failure need not be uniform all over country. Prevalence of treatment failures may be heterogeneous and may vary from region to region or states. So it is desired to estimate prevalence of treatment failure at frequent intervals, so that second-line drugs can be introduced only in regions with high prevalence of first-line ART failure.

India has the second greatest HIV burden in the world after South Africa. Monitoring the efficacy of treatment in countries with limited resources is difficult. Unnecessary switching to second-line ART drugs may be a burden in countries like India, with financial constraints. So continuation of first-line ART is cost effective in resource limited settings, especially in
places where prevalence of first-line ART failure is low. Accordingly specific group of patients who meet the NACO definitions of failure criteria, may be switched over to second-line ART.

Some of the studies published their findings about first-line ART failure. In a study from South India in 2004, 40 of 1,443 patients (14%) experienced treatment failure at a mean of 406 days [5]. Dragsted et al. have reported that the incidence of treatment failure at 12 months was 11.6 per 100 person-years of follow-up, and that reduced over time [6]. Rajasekaran et al. reported in their study that cumulative incidence of treatment failure among 1370 patients was 3.9% [3]. Treat Asia HIV observational database (TAHOD) showed that the rate of clinical failure was 7.3 per 100 people-years [7]. Our study was the largest study, covering a period of a decade, including 57,674 subjects. Our study documented that prevalence of failure of first-line ART was only 0.47%, indicating the efficacy of first-line ART.

There were reports about predictors of failure of first-line and ART. Rajasekaran et al. reported that negative change in absolute lymphocyte count, hemoglobin concentrations and body weight during follow-up as inexpensive predictors of failure of first-line ART [3]. Male patients had a 3.5 times significantly greater hazard ratio for treatment failure compared with female patients [3]. Patients from urban areas had 1.9 times greater hazard ratio compared to rural areas [3, 5], which may reflect possible involvement in high-risk behavior.

Anup Singh et al. reported that low baseline CD4 count, lower peak CD4 count achieved, early and lower level of plateau of CD4 count were as important predictors of first-line ART failure [2]. They also documented that co-infection with tuberculosis, older age, and male sex were important predictors of first-line ART failure. Laurence Ahoua et al, found that older age group of patients of >35 years of age were more likely to fail with first-line ART [8]. Deeks et al. have shown that base line CD4 count was associated with treatment failure [9]. These predictors can be used for close follow-up to identify the treatment failure early and assist policy makers in planning second-line treatment regimens where required. Some of the recent cohort studies and RCTs suggested that baseline and time-updated CD4 cell counts are better predictors of HIV-1 disease progression than are Plasma Viral Loads (PVLs) [10, 11]. A lower risk of clinical progression was reported among female patients with intermediate baseline viral load than in males [12].

Adherence is important in achieving suppression with an ART regimen. Adherence begins to wane after the first month of therapy. So, closer assessment of adherence particularly after first month of therapy is important [13]. Suboptimal adherence to these regimens has been postulated to be one of the main factors associated with decreased HIV suppression as well as for the emergence of resistant virus [14, 15].

Prevalence of primary HIV drug resistance mutations (DRMs) in northern India is 2.9% and <5% in Kakinada of southern India, which are within the threshold limit of <5% [16, 17]. This finding reinforces the National ART program’s effort in maintaining low level of primary ART drug resistance. Until the baseline HIV genotyping becomes more affordable, there is a need for Periodical studies to assess primary ART drug resistance from different regions of India.

Kumaraswamy et al. reported that 20% of patients modified their first-line ART regimen and the most common reason for modifying therapy was development of adverse reactions and treatment failure was less common cause [5, 18].

The association between HIV infection and tuberculosis is complex and bi-directional [19]. The effect of tuberculosis on HIV replication is mediated by cytokines IL-1, IL-6, and TNFα, which in turn results in enhanced viral replication [20].

TAHOD study has shown that treatment with the simple fixed-dose combination Stavudine, Lamivudine, and Nevirapine was safe and effective, with good adherence and tolerability and also adverse events are the common reason for treatment change [7, 21].

CONCLUSION

- Prevalence of failure of first-line ART was 0.47%.
- No significant association between mean duration and failure of first-line ART.
- Males had higher incidence of first-line ART failure, compared to females.
- Low mean CD4 counts were associated with higher failure rates.

Low prevalence of failure in our study indicates that first-line ART is still effective and can be continued, especially in resource limited settings like India. But as duration of therapy is longer of many years, there is a chance for development of resistance. As a result, prevalence of first-line ART failure need to be assessed periodically in different parts of the country, so that second-line ART may be needed only in regions with high failure rates. Routine virological monitoring and genotyping resistance is not practicable at present in our country. So it is ideal to monitor ART with watchfulness and using non-expensive criteria like thorough clinical history, adherence to treatment, hemoglobin level, total lymphocyte count, body weight measurement, peak CD4 count, and treatment of opportunistic infections [22]. Good adherence also reduces incidence of drug resistance. This is the ideal procedure until routine virological monitoring and
testing for primary drug resistance becomes affordable in our country.

REFERENCES

9. Deeks SG, Barbour JD, Grant RM, Martin JN; Duration and predictors of CD4 T cell gains in patients who continue combination therapy despite detectable plasma viremia. AIDS, 2002; 16(2): 201-207.