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An evaluation of health-related quality of life and its determinants among people living with HIV/AIDS attending tertiary care hospital of Kurnool, India

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ABSTRACT

Background: HIV/AIDS is known to affect an individual not only physically but also mentally, socially, and financially. It is a syndrome that builds a vacuum in a person affecting his/her life as a whole. Combined with ART, Quality of life (QoL) is an important component in the evaluation of the wellbeing of people living with HIV/AIDS (PLHIV). The objective of this study is to assess health related quality of life of HIV/AIDS patients attending ART clinic Tertiary care hospital, GGH, Kurnool and to determine the association of socio-demographic and disease related variables with health related quality of life.

Methods: A cross-sectional study was conducted from April 2019 to June 2019 involving 400 purposively selected HIV-positive patients of age >18 years, who were taking highly active anti-retroviral therapy for the past 6months from the ART center, GGH, Kurnool. After obtaining IEC clearance and informed consent, WHOOOL-BREF instrument was used for data collection. Data analysis was performed using IBM SPSS version 26.0.

Results: Out of the 400 participants, 60% were males. The mean age of the participants was 38.5±10.54 years. Overall quality of life had a mean score of 69.71. Physical (82.57) and level of independence (78.78) domains showed higher mean score when compared to psychological (63.82), environmental (61.49) and Social (60.26) domains.

Conclusions: Among study subjects 15.5% had excellent QoL (≥80), nearly 69.75% had good QoL (60-79) and 14.75% had poor QoL (<60). Low QoL scores were seen in the social domain, suggesting that more social interventions are required in this population.

Keywords: Health related quality of life, Patients living with HIV/AIDS, WHOQOL-BREF

INTRODUCTION

Acquired immune deficiency syndrome (AIDS) is a syndrome caused by human immunodeficiency virus (HIV). HIV or AIDS is one of the serious public health problems with severe impact on various facets of life. Globally, 36.9 million people were living with HIV at the end of 2017. An estimated 0.8% of adults aged 15-49 years worldwide are living with HIV.1 Around 59% of people living with HIV were receiving antiretroviral

treatment by the end of 2017. In India, an estimated 21.40 lakhs people living with HIV (2017) with the prevalence of 0.22% of adult population. Andhra Pradesh state stands among top five highest prevalent states with a prevalence of 0.63%.² According to recent National AIDS Control Organization data, India is estimated to have around 87.58 (36.45-172.90) thousand new HIV infections in 2017, showing new HIV infection decline by 85% since the peak of 1995 and by 27% between 2010 to 2017. As per UNAIDS report in 2017 there were a total 69000 number of HIV/AIDS related deaths in India. Since the peak, the number of annual AIDS-related deaths has declined by almost 71%.²

Antiretroviral drugs have revolutionized the treatment for HIV by increasing the average life span of HIV positive individuals. India has launched national ART (antiretroviral therapy) program in 2004 with the aim to provide free ART drugs to the patients suffering from HIV or AIDS. At present there are 448 ART centers providing access to patients to get free ART drugs. With the introduction of antiretroviral treatment the course of disease has drastically changed from a fatal to a chronic and potentially manageable disease. However people with HIV or AIDS continue to have substantially lower quality of life than general population, even where the majority of those living with HIV have virological control and immunologically stable. Combined with ART, improving quality of life is central to the care and support of people living with HIV (PLHIV).³ Improved access to ART and other biomedical interventions must be adequately matched with the requisite psychosocial treatment to help improve the effectiveness of such interventions and for overall better outcome of the disease.4

Optimizing care for PLHIV requires an understanding of the factors that contribute to physical health, psychological wellbeing, social relationships, and quality of life. In view of achievable longer life span among people using ART, quality of life (QOL) has emerged as a significant medical outcome measure and its enhancement has an important goal. The World Health Organization (WHO) defined QOL as individuals perception of their position in life in the context of the culture and value systems in which they live and relation to their goals, standards, expectations and concerns.⁵ It is a broad ranging concept affected in a complex way by the person's physical health, psychological state, personal beliefs, social relationships and their relationship to salient features of their environment. 5The second objective of the National AIDS control program phase IV of India aims to provide comprehensive care to people living with HIV or AIDS (PLHIV). Improving QOL is in line with comprehensive care. Studies in context of assessing QOL among PLHIV are few in India and very few in Andhra Pradesh state which stand among one of the highest prevalence states in the country. Hence this study was aimed to assess health related quality of life of HIV or AIDS patients attending ART clinic, Tertiary care hospital, GGH, Kurnool and to determine the association of socio-demographic and disease related variables with health related quality of life.

METHODS

A cross-sectional study was conducted among people living with HIV positive/AIDS aged above 18 yrs and on ART drugs for more than 6months duration attending to

ART centre of tertiary care hospital, Kurnool from April 1st to June 30th 2019.

Ethical issues

Ethical clearance was obtained from the Ethical Committee of Kurnool Medical College before starting the study proper. The purpose of the study was clearly explained to all study subjects in their local language and interviewed after taking informed consent.

Sample size

Sample size was calculated by using, a z value of 1.96 as the degree of accuracy at 95% confidence interval, proportion of PLWHA with QOL level of better than average was assumed to be 50% and 10% allowable error. Based on this the sample size was calculated to be 392. in the present study, 400 study subjects were enrolled.

Inclusion criteria

People living with HIV or AIDS (PLHIV) older than 18 years and on ART drugs for more than 6 months duration attending to ART centre, GGH, Kurnool.

Exclusion criteria

People living with HIV/AIDS (PLHIV) less than 18 years, patients on ART drugs less than 6 months duration, those not registered at the center and patients who were severely ill were excluded from the study.

Study tool

Study subjects were interviewed by using World Health HIV (WHOQoL-HIV) Organization OoL questionnaire. There was an orientation session for the filling up of Study tool for the project team members. The questionnaire consisted of 31 items in six domains (physical health, psychological health, level of independence, social relationships, environment, and religious/personal beliefs).8 Each item contained a 5-point Likert type scale that is best represented on their opinion. On the scale one (1) indicates low and negative perceptions, scale five (5) indicates high and positive perceptions, which denoted better QOL. Negatively worded items were scored in reverse order, and all the scores were checked for appropriate range between 1and 5. Quality of life scores were categorized into three sections with scores ≥80 denoting excellent QoL, 60 to 79 denoting a good QoL, and <60 representing poor QoL.

Other data that were collected include socio-demographic information such as age, gender, education level, employment, per capita monthly income, ART duration, WHO clinical stage, opportunistic infections (both present and past), recent CD4 count.

Statistical analysis

Analysis was performed using Statistical Package for Social Sciences (SPSS) version 26.0. Data were represented using frequency, percentages and mean scores with standard deviation. Tests of significance were performed at each level and a p value less than 0.05 is considered statistically significant; homogeneity of variance was tested and considered appropriately. Percentage scores were calculated as the sum of individual scores in a domain divided by total attainable score in that domain multiplied by 100. Thus the percentage scores ranges from minimum of 25 to a maximum of $100.^{9.10}$ Internal consistency reliability scale was examined using Cronbach's α . Cronbach's α of 0.6 or above was considered acceptable.

RESULTS

Out of the 400 participants, 240 (60) were males and 160 (40) were females. Age ranged from 18 to 70 years. The mean age of the participants was 38.5±10.54 years. Most of them were in age group 31-45 years (48.7) (Figure 1). The majority of the patients were Hindus 345 (86); 360

(90) were married, 229 (57.2) were illiterates. Unemployed and unskilled workers together account for 207 (52). Most of them 161 (40.2) belonged to lower-middle socio-economic class and nearly 287 (72) were in Stage 1 (Table 1).

Internal consistency reliability scale was examined using Cronbach's α. Cronbach's α of 0.6 or above was considered acceptable. The internal consistency of all the domains of the instrument (WHOQOL-HIV bref) was found to be good. Average of internal consistency of all domains of the instrument was found to be between 0.65 and 0.83. The mean±SD of overall quality of life was 69.71±10.15. Physical (82.57) and level of independence (78.78) domains showed higher mean score when compared to psychological (63.82), environmental (61.49) and social (60.26) domains which had lowest mean scores (Table 2).

Among the study population, 62 (15.5) were graded as presenting with an excellent (\geq 80) overall QoL, 279 (69.75) presented with good overall QoL (60-79), and 59 (14.75) had poor QoL (<60) (Table 3).

Table 1: Socio-demographic and disease related characteristics of the study subjects.

	Male	Female	Total	
Characteristics	N (%)	N (%)	N (%)	
Age group (years)				
15-30	47 (11.7)	63 (16.7)	110 (27.5)	
31-45	121 (30.2)	74 (18.5)	195 (48.7)	
46-60	65 (16.2)	21 (5.2)	86 (21.4)	
>60	7 (1.7)	2 (0.5)	9 (2.2)	
Marital status	·	•		
Un-married	18 (4.5)	6 (1.5)	24 (6)	
Married	219 (54.7)	141 (35.2)	360 (90)	
Divorce	1 (0.25)	2 (0.5)	3 (0.75)	
Widowed	2 (0.25)	11 (2.7)	13 (3.2)	
Religion				
Hindu	208 (52)	137 (34)	345 (86)	
Muslim	17 (4.2)	11 (2.7)	28 (7)	
Christian	15 (3.7)	12 (3)	27 (6.7)	
Education status				
Illiterate	118 (29.5)	111 (27.7)	229 (57.2)	
Primary	13 (3.2)	5 (1.2)	18 (4.5)	
middle	20 (5)	10 (2.5)	30 (7.5)	
High school	65 (16.2)	26 (6.5)	91 (22.7)	
Intermediate	16 (4)	5 (1.2)	21 (5.2)	
Graduate and above	8 (2)	3 (0.7)	11 (2.7)	
Employment status				
Professional	0	0		
Semi professional	20 (5)	3 (0.7)	23 (5.7)	
Skilled worker	64 (16)	11 (2.7)	75 (18)	
Semi-skilled worker	69 (17.2)	26 (6.5)	95(23.7)	
Un skilled worker	77 (19.2)	63 (15.7)	140 (35)	
Un-employed	10 (2.5)	57 (14.2)	67 (16.7)	

Continued.

Characteristics	Male	Female	Total	
SES				
Upper	7 (1.7)	1 (0.25)	8 (2)	
Upper-middle	80 (20)	42 (10.5)	122 (30.5)	
Lower-middle	93 (23.2)	68 (17)	161 (40.2)	
Upper-lower	27 (6.7)	20 (5)	47 (11.7)	
Lower	33 (8.2)	29 (7.2)	62 (15.5)	
Stage of HIV				
Stage 1	159 (39.7)	128 (32)	287 (71.7)	
Stage 2	57 (14.2)	25 (6.2)	82 (20.5)	
Stage 3	23 (5.7)	7 (1.7)	30 (7.5)	
Stage 4	1 (0.25)	0	1 (0.25)	

Table 2: Mean (SD) of domains of the quality of life with scales description.

QoL-domain	Mean (SD)	Cronbach's α	Correlation with total score
Physical domain	82.57±15.61	0.683	0.679
Psychological domain	63.82±12.47	0.676	0.774
Level of independence	78.78±13.81	0.689	0.665
Social relation	60.26±15.32	0.829	0.287
Environmental domain	61.49 ±13.07	0.717	0.558
Spiritual domain	72.47±17.48	0.695	0.679
Overall HQOL	82.57±15.61	0.653	1.000

Table 3: Components of the quality of life of people living with HIV or AIDS in study subjects.

Parameter	Excellent	Good	Poor
rarameter	N (%)	N (%)	N (%)
Physical domain	254 (63.5)	112 (28)	34 (8.5)
Psychological domain	43 (10.7)	250 (6.2)	97 (24.2)
Level of independence	241 (60.2)	127 (31.7)	32 (8)
Social relation	70 (17.5)	145 (36.2)	183 (45.7)
Environmental	33 (8.2)	224 (56)	143 (35.7)
Spiritual	165 (41.2)	152 (38)	83 (20.7)
Overall	62 (15.5)	279 (69.75)	59 (14.75)

Table 4: Comparison of the mean±SD of different domains based on demographic variables, disease related characteristics.

Variable	Physical domain	Psychological domain	Level of independence	Social relation	Environ- mental	Spiritual	Overall score
Gender							
Male (n=240)	82.88±15.94	63.83±14.04	78.12±15.08	62.47±15.69	61.23±13.65	73.77±17.97	70.13±11.20
Female (n=160)	82.09±15.13	63.80±9.70	79.69±11.62	56.96±14.16	61.89±12.20	70.52±16.58	69.08±8.29
P value	0.62	0.98	0.24	0.000*	0.62	0.068	0.28
Age (years)							
≤30	84.6±15.9	65.06±13.1	80.85±12.6	61.45±14.6	63.14±14.2	68.47±18.5	70.35±9.9
>30	81.7±15.4	63.34±12.2	77.9±14.1	59.81±15.5	60.87±12.5	73.99±16.8	69.47±10.2
P value	0.10	0.21	0.06	0.34	0.12	0.005*	0.43
Marital status	3						
Married	82.20±15.61	64.01±12.18	78.67±13.75	60.81±15.09	61.84±13.06	72.72±16.95	69.84±9.90
Un-married	89.78±14.57	67.22±13.38	83.26±10.40	61.13±17.26	61.89±11.99	76.52±20.91	73.26±11.04
Divorce	78.33±25.68	46.67±23.09	66.67±32.14	33.33±11.54	53.00±18.24	50.00±8.66	54.66±19.73
Widowed	80.77±13.51	56.54±12.01	75.77±14.41	49.77±9.61	53.15±12.41	63.62±21.92	63.23±8.19
P value	0.138	0.007**	0.147	0.001**	0.077	0.022**	0.002**

Continued.

Variable	Physical domain	Psychological domain	Level of independence	Social relation	Environ- mental	Spiritual	Overall score		
Education star	Education status								
Illiterate	81.80±15.58	63.06±11.20	77.85±13.19	58.85±14.93	60.09±12.78	71.35±17.20	68.82±9.76		
Primary	79.72±14.60	61.67±15.03	76.50±14.93	58.00±15.48	54.47±10.70	72.50±16.56	66.94±11.70		
Middle	84.67±14.67	61.47±15.26	79.20±15.20	58.43±16.40	61.76±10.88	70.50±20.98	68.70±10.98		
High school	84.39±15.41	67.02±13.12	80.61±14.21	64.23±15.19	64.56±13.54	74.13±16.75	72.00±10.02		
Intermediate or diploma	80.24±19.00	60.19±12.55	77.86±17.36	59.90±17.53	63.73±14.00	74.81±19.97	69.14±10.89		
Graduate and above	86.91±15.49	70.27±15.07	86.36±6.74	66.64±12.01	72.18±12.89	83.18±13.46	77.90±7.58		
P value	0.518	0.021**	0.257	0.056	0.001**	0.248	0.009**		
Socioeconomic	c status								
Upper	86.88±16.02	61.50±12.81	83.75±10.93	62.75±16.43	65.25±19.31	75.00±17.11	73.12±10.03		
Upper-middle	82.69±14.52	65.48±11.85	79.56±13.34	60.13±14.31	65.33±10.49	74.16±14.11	70.89±8.84		
Lower-middle	83.29±16.10	63.99±12.89	77.98±14.88	59.96±16.03	61.37±13.24	72.37±18.11	69.44±10.74		
Upper-lower	80.96±14.50	63.98±12.24	78.72±12.66	58.00±13.11	59.31±13.01	69.79±19.99	68.53±9.59		
Lower	81.53±17.09	60.37±12.44	78.52±13.25	62.69±16.90	55.28±13.94	70.85±19.78	68.55±11.38		
P value	0.786	0.126	0.740	0.583	0.000**	0.583	0.396		
Stage of diseas	se								
Stage 1 (n=287)	85.07 ±14.88	65.71±11.02	81.18±12.08	60.97±15.39	63.14±12.37	74.55±16.50	71.50±9.20		
Stage 2 (n=82)	78.18 ±15.18	61.07±14.32	74.61±15.16	58.54±14.75	58.53±13.59	67.79±18.41	66.14±10.44		
Stage 3 (n=30)	71.67 ±15.72	54.27±13.55	67.40±16.99	57.37±15.51	53.45±14.27	66.43±19.86	62.80±12.48		
Stage 4 (n=1)	-	-	-	-	-	-	-		
P value	<0.001**	<0.001**	<0.001**	0.149	<0.001**	<0.001**	<0.001**		
Duration of A	RT (years)								
0.5-4	82.58±15.41	63.16±13.28	79.01±13.15	60.23±15.75	61.16±13.13	71.26±17.61	69.11±10.13		
4.1-8	84.24±14.56	64.95±11.16	80.06±12.80	59.04±14.23	61.37±12.30	73.67±17.14	70.68±9.50		
8.1-12	80.63±16.67	64.03±12.48	76.54±16.90	61.69±15.93	62.57±14.32	73.22±17.11	69.51±11.19		
>12	78.95±19.40	62.11±12.67	76.11±13.20	63.05±15.78	61.52 ±13.35	73.68 ±20.26	70.16±10.62		
P value	0.316	0.593	0.296	0.565	0.893	0.638	0.609		
CD4 count									
<350/μl	79.97±17.07	61.32±12.94	74.90±16.73	58.67±14.38	58.18±13.15	71.91±17.64	66.93±10.35		
>350/µl	83.61±14.88	64.82±12.16	80.30±12.13	60.91±15.65	62.83±12.83	72.69±17.44	70.83 ± 9.86		
	0.046*	0.011*	0.002*	0.187	0.001*	0.686	0.000*		

^{*} P value (Student's t-test), **P value (ANOVA).

Physical domain which measured pain, discomfort, energy and sleep was affected by CD4 count (p value 0.046, Table 4) and stage of the disease (p<0.001) with the advancement of stage of disease mean score value is decreased significantly. Psychological domain which measured ability to concentrate, self-satisfaction, enjoyment and negative feelings was significantly affected by educational status, marital status (divorced individuals had lowest mean score 46.67 followed by widowed who had mean score of 56.54), stage of the disease and CD4 count (Table 4). In the Level of independence domain which measured dependence on medications, mobility, ability to perform daily activities and capacity to work, significantly lower mean scores

were observed with advancement of stage of disease and lower CD4 count (p<0.001).

Social relationship domain which measured personal relationships, sex life and social support has showed significant variation with gender (female had lower mean score 56.96 when compared to males 62.47, p<0.001, see Table 4), marital status (divorced individuals had lowest mean score of 33.33 followed by widowed who had a mean score of 49.77, p=0.001, Table 4). Educational status, socioeconomic status, stage of the disease and CD4 count does not appear to significantly alter the social relationship domain. Environmental domain which measured access to health care, transport, conditions of

living place, money for daily needs was significantly affected by educational status (p <0.001), socioeconomic status (p<0.001), stage of the disease (p<0.001) and CD4 count (p<0.001). Spiritual domain which measured blame, fear about future and death was affected by age (individuals aged <30 years had a mean of68.47 and individuals aged >30 years had a mean of 73.99, p<0.001), marital status and stage of the disease. Overall quality of life showed significant variation with marital status, educational status, stage of the disease and CD4 count. The mean score of all the domains significantly

decreased with advancement of stage of the disease. (Table 4).

Most commonly associated opportunistic infection was pulmonary TB (1 in 4 individuals had pulmonary TB) followed by oral Candidiasis (2) (Table 5).

CD4 count varied between 12-1290 cells/ mm³. Mean CD 4 counts in all study subjects was 492.66±220.51 cells/ mm³. Overall quality of life significantly improved with increased CD4 counts in PLHA (p value 0.019) (Table 6).

Table 5: Comparison of groups with different CD4 counts with overall quality of life score by one-way ANOVA.

CD4 cell count	No. of patients (%)	Overall HQOl Mean±SD	P value
<50	3 (0.75)	59.33±3.511	
51-100	7 (1.75)	63.14±9.13	
101-200	19 (4.75)	67.89±10.56	
201-300	48 (12)	68.85±8.30	0.019**
301-400	69 (17.25)	67.43±11.73	
401-500	66 (16.5)	71.91±10.72	
Above 500	188 (47)	70.59±9.52	

Table 6: Opportunistic infections (present and past) among patients.

Opportunistic infection (present/past)	Male (%)	Female (%)	Total (%)
Pulmonary TB	67 (16.75)	33 (7.25)	100 (25)
Oral candidiasis	6 (1.5)	2 (0.5)	8 (2)
Lymphadenopathy	6 (1.5)	0	6 (1.5)
Pneumonia	3 (0.75)	1 (0.25)	4 (1)
Herpes zoster	2 (0.5)	1 (0.25)	3 (0.75)
TB bone	1 (0.25)	0	1 (0.25)
Cryptococcal meningitis	1 (0.25)	0	1 (0.25)

DISCUSSION

In our study, the most affected domain was social relationship domain followed by environmental and psychological domains. Various other studies also showed low social and psychological domain scores. ¹²⁻¹⁵ This can be explained by societal discrimination, stigmatization, isolation, dissatisfaction with sex life and negative feelings. Poor environmental domain score in our study is in contradiction with other studies ^{11,15} as majority of people are from rural area because of which they have poor access to health care, poor environmental conditions and poor transport facilities.

In the present study, the average of internal consistency of all domains of the instrument was found to be between 0.65 and 0.83. The study done by Nojomi et al among people living with HIV/AIDS, Iran showed that the average of internal consistency of the four domains of the instrument was found between 0.64 and 0.83, which is similar to the present study. ¹⁶

Age and gender

The results of the present study showed that the majority of the patients were young men who were affected with disease during their active life. This result is agreed with other studies. ^{16,13} Age had a significant association with spiritual domain only. There is no significant variation in any domain of QoL by gender except in social relationships domain, this may be due to lack of social and family support, cultural beliefs in addition to stigma of the disease.

Marital status

Single individuals reported a better QoL score in each domain than married, Liping et al reported the same.¹⁷ This result was not consistent with the study of Shakirat et al, Dasgupta et al and Nojomi et al.^{11,15,16} Some other studies reported no significant association between marital status and QoL.¹⁰ Married persons had significantly scored better overall QoL than widowed and divorced which can be attributed to additional social

stigma and lack of physical and emotional support from their spouses as it was before.

Educational status

Overall quality of life significantly improved with higher educational status. This is because patients with higher education can have better understanding about their disease state, coping strategies and adherence to treatment. Primary education has no effect on quality of life as patients with only primary school education scored less than patients with no education. Higher education level had significant association with psychological and environmental domains. Study done by Liping et al showed that Those with higher education level had better scores in physical and environmental domains (p<0.05)

Socioeconomic status

Present study shows that overall quality of life does not have significant association with socio economic status of HIV/AIDS Patients. Socio economic status had a significant association with Environmental domain. This makes economically backward patients more prone to poor QOL in environmental domain. Similar findings observed in study done by Dasgupta et al.¹⁵

Stage of the disease and CD4 count

Both stage of the disease and CD4 count are indirectly related to each other. Significantly better scores were observed with higher CD4 count and with asymptomatic stage of disease except for the social domain. The significant impact of stage of illness reflects grievous and disabling nature of the disease on various aspects of quality of life. Stage of the illness does not appear to have a significant effect on personal relationships and sex life.

CONCLUSION

The present study shows the physical domain, level of independence and spirituality of the patients were relatively higher than psychological, environmental and social relationship domains. Age, sex, socioeconomic status was significantly associated with at least one domain of HQoL. Educational status was strongly associated with psychological, environmental domain. strongly status was associated Marital psychological, social and Spiritual domain. Moreover, WHO clinical stage was strongly associated with all HQoL domains except social domain.CD4 count was strongly associated with all HQoL domains except Spiritual and social domain. We observed that overall poor quality of life is related to educational status, marital status, CD4 cell count and stage of illness.

Recommendations

 The present findings highlight the need for enhanced socio psychological support and better environmental

- health. Strengthening social relations seems relatively more demanding than other domains.
- Awareness programs to address the issue of stigma and discrimination at level of family and community to reduce stigma to the barest minimum, which may help to create a sense of safe living conditions or help PLHIV to receive a necessary support from their family and community should be organized.
- Mental health care should be provided for all the PLHIV and suitable employment opportunities considering their functional capacities should be done thus focusing aspects of HIV beyond clinical care. Employment and financial security would lead to improve social health.
- Regular sensitization is needed for all health professionals and staff in ART centers about the condition of PLHA, so that at any time of the treatment they should maintain the focus, affection and empathy towards these patients.
- Clinical assessment of adverse reactions during the course of ART treatment and careful monitoring of patient after any ART switch could contribute to a better QOL, improve the patient—doctor relationship, and potentially maintain adherence with fewer undesired side effects.
- Sometime hospital-based care is not feasible or the patient is unable to come due to his personal reasons, home or community-based care is one appropriate option for better care and compliance.
- Policies should be made and rules should be enforced to protect the rights of PLHA in the work place as well as prevent any form of discrimination against them in the society.
- Therefore, the focus should be on various multidisciplinary care models (including primary medical care, social support, education and employment opportunities) with help of Government schemes and involving NGOs and care at home to promote HR-QOL in these patients.

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