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Accessibility
HIV-Care Outcome in Saudi Arabia; a Longitudinal Cohort

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Abstract

Background—Clinical characteristics of HIV-1 infection in people inhabiting Western, Sub-Saharan African, and South-East Asian countries are well recognized. However, very little information is available with regard to HIV-1 infection and treatment outcome in MENA countries including the Gulf Cooperation Council (GCC) states.

Methods—Clinical, demographic and epidemiologic characteristics of 602 HIV-1 infected patients followed in the adult Infectious Diseases Clinic of King Faisal Specialist Hospital and Research Centre, in Riyadh, Kingdom of Saudi Arabia a tertiary referral center were longitudinally collected from 1989 to 2010.

Results—Of the 602 HIV-1 infected patients in this observation period, 70% were male. The major mode of HIV-1 transmission was heterosexual contact (55%). At diagnosis, opportunistic infections were found in 49% of patients, most commonly being pneumocystis. AIDS associated neoplasia was also noted in 6% of patients. A hundred and forty-seven patients (24%) died from the cohort by the end of the observation period. The mortality rate peaked in 1992 at 90 deaths per 1000 person-year, whereas the mortality rate gradually decreased to <1% from 1993-2010. In 2010, 71% of the patients were receiving highly active retroviral therapy.

Conclusions—These data describe the clinical characteristic of HIV-1-infected patients at a major tertiary referral hospital in KSA over a 20-year period. Initiation of antiretroviral therapy resulted in a significant reduction in both morbidity and mortality. Future studies are needed in the
design and implementation of targeted treatment and prevention strategies for HIV-1 infection in KSA.

Keywords
HIV; Human immunodeficiency Virus-1; CD4+T cells; Antiviral therapy; Viral load; Opportunistic infections; AIDS; Acquired immunodeficiency syndrome

Introduction

Human immunodeficiency virus (HIV) infection represents one of the preeminent global health problems and a major health delivery challenge. According to the World Health Organization (WHO) and the joint United Nations program on HIV/AIDS (UNAIDS), 33 million people worldwide are currently infected with HIV (http://www.unaids.org). The use of antiretroviral therapy has dramatically improved the outcome and survival of HIV-infected persons.

A number of analyses of treatment outcomes in longitudinal cohorts from both well-resourced and resource-limited settings have been published [1-9]. However, data on HIV treatment outcomes in the MENA region are limited. In the decade ending in 2009, approximately 24,000 out of the estimated 460,000 people with HIV in the MENA region died of AIDS. Although estimates of the prevalence of the disease remain low (approximately 0.1%-0.2% in Egypt, Iran, Jordan, Libya, Morocco, Syria, Tunisia, Gulf Cooperation Council (GCC) countries, and Yemen), the number of new infections in the MENA region is increasing [10].

HIV is increasingly recognized as a major health problem in Kingdom of Saudi Arabia (KSA) and in the rest of member states in GCC [11]. The first official case of HIV infection in KSA was reported in 1984 in a blood transfusion recipient [12]. By 1986, 13 HIV-infected patients in KSA had been documented, all linked to contaminated blood products [13]. Between 1984 and 2001, 6,046 HIV infections were reported in KSA, 1,285 (21%) of whom were Saudi citizens [14, 15]. In response to the epidemic, a national AIDS program was established within the Saudi Ministry of Health (MOH) to oversee and coordinate country-wide efforts to prevent, diagnose and treat HIV. One of the first care provision facilities was the King Faisal Specialist Hospital and Research Centre (KFSHRC), where the first case had been reported. HIV-infected patients from all provinces of KSA are referred to this institution for counselling, diagnostic evaluation, and treatment. Currently, one out of four HIV-infected patients in KSA is receiving care in this facility. Clinical and demographic data from these patients have been entered longitudinally into a central database. This database is primary data source for HIV treatment outcomes in Saudi Arabia. The purpose of this study is to describe HIV treatment outcomes in a cohort of 602 prospectively enrolled, longitudinally-followed HIV patients, whom all are Saudi National from different social- economic background.
Methods and Materials

Patient Cohort

HIV-infected patients were collected in this observational cohort from 1989 to 2010 at the KFSHRC. Over the 20-year period, 602 patients all of whom are Saudi Nationals were enrolled representing 1,200 annual patient-visits. Broadly speaking, patients in the cohort belonged to three groups: 1) newly diagnosed HIV patients within the hospital’s catchment area (around 5 new cases per year, mostly contacts of recently diagnosed patients-spouses, parents or children); 2) referral from the Ministry of Health (MOH) or other care providers; and 3) self-referrals. The data were entered into a customized database that consisted of 150 variables of patient information; including demographic data such as geographic origin, age, and gender. In addition, patient’s medical information was collected by chart review to include age at presentation, mode of transmission, symptoms, laboratory analyses, antiretroviral therapy, and opportunistic infections.

HIV diagnostics

Until 1996, HIV diagnosis was established by enzyme immunoassay with confirmation by immunoblot assay. After 1997, HIV polymerase chain reaction (PCR) was used to confirm the diagnosis in patients with reactive serologic tests.

Initial serologic testing for HIV 1 and 2 was performed using a micro-particle enzyme immunoassay (MEIA; AxSYM HIV 1/2 gO, Abbott Laboratories, Abbott Park, USA). If serology was reactive, specimens were confirmed using the Chiron RIBA HIV-1/HIV-2 strip immunoblot assay.

For the purpose of this report, we used the CDC definition of AIDS [16]. Opportunistic infections are defined by microbiological confirmation (by stain for Pneumocystis jirovecii, culture for M. tuberculosis, atypical mycobacteria, and fungi).

Molecular testing included HIV PCR, which uses three sets of four primers from envelope, polymerase, reverse transcriptase, and core protein genes. Viral load was measured with the branched DNA method (VER SANT HIV-1 RNA 3.0 Assay, [bDNA], Bayer Diagnostics, Berkeley, CA, USA). Results were reported in a range from 50 to >500,000 copies/ml. CD4+ count and percentage was obtained by standard flow cytometric methods.

Statistical analysis

The software package SAS version 9.2 was used to perform the statistical analysis (Statistical Analysis System, SAS Institute Inc., Cary, NC, USA). Descriptive statistics for the continuous variables were reported as mean ± standard deviation. Categorical variables were reported as frequencies and percentages and the Chi-square test was used to test for statistical significance. The statistical level of significance was set at p < 0.05.
Ethics

This study was conducted in accordance with the Helsinki Declaration and approved by the Research Ethics Committee and Research Advisory Council (ORA # 2031 031). Consent was waived by the KFSHRC IRB as the study was a retrospective chart review study.

Results

Patient Characteristics

Of the 602 patients in the cohort, 420 (70%) were male. The mean age at diagnosis was 30 years (standard deviation (SD) =14, range=1–91). KFSHRC is located within the metropolitan area of Riyadh (the country’s capital) and serves as a referral hospital for the central region of KSA. Almost one-third of the patients in the cohort were originally from this part of the country (33%, n=196). The origin of the remaining patients was broadly diversified: southern region (27%, n=161); western region (22%, n=131); eastern region (16%, n=94); and northern region (3%, n=19). During the observation period, 147 (24%) patients died and 83 (14%) were lost to follow up (Table 1) with an average of 3 patients lost to follow up per year with a peak loss of 12 patients in 2002. However, after 2010 most of the patients (62%, n=372) were still being seen for follow-up at the infectious disease clinic at KFSHRC.

A number of patients diagnosed with HIV also had HIV positive family members including spouses (14%, n=84), parents (6%, n=37), or children (5%, n=28). The major route of transmission was heterosexual contact (55%, n=329) followed by blood transfusion (24%, n=146). The paediatric age group (<14 years of age) consisted of 93 (15%) patients. Perinatal transmission was identified as the route of transmission in 59 (63%) children.

The annual frequency of new HIV-1 infections initially peaked between 1986 and 1990 (Figure 1). Most HIV-1 infections observed during this time were causally linked to transfusions with blood and blood products (Table 2). One patient was infected during bone marrow transplantation. Six patients acquired HIV-1 infection after kidney transplantation; two of these kidney transplants were performed as cadaveric kidneys in the United States in 1985 and four patients received the kidneys from commercial non-related donors in India and Egypt. The second peak in new HIV-1 infection occurred from 1999 to 2003 and was predominantly referrals from other hospitals within KSA. Most of these infections were linked to heterosexual transmission (65%, n=141). Since 2004, there has been a slow but steady decline in new HIV-1 patients referred from other institutions as new HIV/AIDS treatment centres have been opened around the country.

Opportunistic infections and AIDS associated malignancies

In the 1990’s, we recorded 184 episodes of opportunistic infections and 23 AIDS associated malignancies, resulting in an OI rate of 10.6 cases per 100 patient years. Pneumocystis jirovecii pneumonia (PCP) was the most frequently diagnosed opportunistic disease (27%, n=50), followed by Candida esophagitis (25%, n=45). Mycobacterium tuberculosis (MTB) and cytomegalovirus (CMV) infections was recorded in 29 patients each and represented 16% of all opportunistic infections. Most cases of CMV infection involved retinitis (72%,
n=21), pneumonitis (14%, n=4), colitis (7%, n=2), hepatitis (7%, n=2). Of the MTB infections (n=29), 59% represented pulmonary TB, 24% with disseminated infections, 10% involved the central nervous system (CNS) and 7% other sites. Atypical mycobacterial infections were diagnosed in 17 patients (9% of opportunistic infection). M. avium-intracellulare (71%), M. kansasii (12%), and M. chelonae (6%) were the most frequently isolated species. Other reported opportunistic diseases included cryptosporidiosis (n=7, 4% of opportunistic infections, four cases with intestinal disease and three cases with gastric disease), and cryptococcal infection (n=6, 3% of opportunistic infections, four cases with disseminated disease and two cases of meningitis).

Reports of AIDS associated malignancies including Kaposi sarcomas were less frequent reported and accounted for 3% in our cohort of patients (1.5% skin, 0.5% pulmonary, 0.7% gastro-intestinal, and 0.3% disseminated). Lymphoma was reported in approximately 1% of patients, 0.2% Hodgkin’s disease and 0.7% Non-Hodgkin’s Lymphoma). The mortality rate for associated malignancies is listed in Table 3.

Treatment characteristics

Antiretroviral drugs have been available in KSA since 1987. They are used according to international standards and guidelines, for the treatment of adult and paediatric patients. Overall, the number of patients receiving antiretroviral therapy rose continuously for this cohort over the observation period (Figure 2). Zidovudine (AZT) mono-therapy was started in 1989 and administered to 20 patients. Subsequently, combination treatments with two nucleoside reverse transcriptase inhibitors (NRTI) were provided to a total of 56 patients between 1990 and 1996. Highly active antiretroviral therapy (HAART) began in 1996 as a combination of three or four antiretroviral drugs. Clinically, the initiation of HAART had dramatic impact in the mortality rate of HIV infected individuals in KSA. In our cohort, the mean CD4 count at the time of diagnosis is 350 cells/mm$^3$ (performed on 244 patients at diagnosis) with an average viral load of 183, 879 copies per ml (available on 134 patients at diagnosis). Since 1996, the number of patients presenting with CD4 <200 has fallen (data not shown), and more importantly, mortality declined to 1% (Figure 3).

Mortality

The mortality in this patient cohort was highest in the period between 1989 and 1993 peaking at 90 deaths per 1000 person-years in 1992 (Figure 3). Mortality in this cohort decreased from 90 to 20 deaths per 1000 person-years between 1993 and 2000, respectively. From 2001 to 2010 mortality declined remaining between 0 to 20 deaths per 1000 person-years, but remains significantly higher when compared to the United States rate of 0.025 deaths per 1000 person-years (http://www.cdc.gov/nchs). In addition, despite increased survival after initiation of HAART, the mortality rate of our HIV cohort at 1% remains substantially higher than the average general KSA population mortality rate 0.3% as reported by the MOH (http://data.worldbank.org).
Discussion

According to the latest MOH report there has been a 0.5-2.5% increase in the incidence of HIV infection between 1984 and 2009 [17]. During this period, 15,213 HIV-infected persons were diagnosed with a breakdown of 4,019 Saudi nationals and 11,194 non-Saudi citizens (Annual AIDS Report-National AIDS Program www.moh.gov.sa). Given that KFSHRC being a main referral centres for HIV infection in KSA, the patient population reflected in this cohort does represent from many of the diverse geographic regions in KSA. Despite the diversity of patients, we recognize the limitation of a single centre cohort. In addition, longitudinal data was not complete for all patients, particularly those patients who may have transferred care to another facility. Even with these limitations, the longitudinal data presented for this cohort are robust and reliable given the standardization of laboratory testing, therapy regimens, and close monitoring and follow up. To our knowledge, this report is the first observational study to describe the outcome of HIV care and related mortality among a cohort of Saudi national HIV-infected patients receiving their care in KSA [14].

The modes of HIV transmission in this KSA cohort are distinct from reports in other MENA countries. In addition, we note that transmission mode followed a biphasic distribution in earlier years as compared to the recent periods. Overall, blood transfusions and blood products account for the highest fraction of HIV transmission, approximately 54%, followed by heterosexual activity at 30%. From 1984-1996, the major mode of HIV transmission was through transfusion of blood and blood products and likely reflects the lack of appropriate blood supply testing during this period. In 1996, the KSA and other GCC states established the Gulf Committee for blood banks that now follows international screening guidelines for blood banking with a subsequent drop in HIV transmission from blood products. From 1994-2010, heterosexual activity remains the dominant risk factor for HIV transmission in our cohort. Compared to other MENA countries, this result is a striking difference where intravenous drug use (IVDU) and men who have sex with men (MSM) are major modes of transmission [18,19]. However, while acquisition of HIV through homosexual activity represents a minor fraction of HIV (1.7%) transmission in our KSA cohort, this finding may also reflect reporting biases from local perceptions and stigma related to homosexuality [20].

Our data also indicate that there has been a steady increase in numbers of HIV infections acquired through IVDU with a peak in 2003 (3%) that subsequently fell. Interestingly, the rise in 2003 IVDU-related HIV acquisition increase may reflect the initiation of a screening program for HIV in the IVDU population presenting for substance abuse treatment in KSA (MOH report for UNAIDS on HIV/AIDS, 2010). With that said, there are limitations in HIV testing that could influences our cohort including the timing of HIV diagnosis, which may reflect differences in both HIV incidence and testing patterns.

In the early 1990s, the rate of opportunistic infections in our cohort is 49.7 episodes per 100 patient-years. These rates are higher than reported from developed countries at the same time period where the average opportunistic rate is 10.6 per 100 patient years [21-29]. In fact, our KSA cohort opportunistic infection rate more closely resembles those from developing countries [30-34] likely reflecting a large proportion of patients presenting with AIDS at the time of initial evaluation. On the other hand, AIDS associated malignancies are
at a rate of 6 per 100 patient years, which is significantly lower compared to other studies [35] but may be due to patient attrition to medical facilities in their home regions.

The initiation of antiretroviral therapy since 1996 has been encouraging, with a general reduction of CD4+ T lymphocytes < 200 and decrease in mortality rate in this cohort. This low rate is a clear programmatic improvement in HIV treatment and intervention. In concert with the findings of CD4+ T cell preservation, for the first time in the MENA region - we identified mortality at <1% in the latter part of the cohort between 1993 and 2010. We believe the free access to healthcare, education and the availability of antiviral therapy accounts for the maintenance of immune reconstituted HIV-infected individuals. In addition, a higher proportion of screening and treatment of infected partners contributes to this finding. However, relative to general population, mortality rates remain three times higher among HIV-infected persons (1% in HIV versus 0.3% general population). Given the growing evidence of increased cardiovascular risk, and malignancy in well-controlled HIV-infected patients, there is need for a more in depth analysis of mortality and morbidity in immune reconstituted, virally suppressed individuals in the KSA and MENA region.

Conclusion

In conclusion, this report from a single-centre in KSA describes the dominant risk for HIV transmission from the period 1984-2010. Results confirm previous observations of delayed identification and treatment of HIV-infection, both resulting in high rates of opportunistic infections and mortality. Following initiation of antiretroviral therapy, patients experienced immune reconstitution with marked reduction in mortality. While our findings do not directly support a strategy for early case-detection of HIV, programs including a nation-wide registry and awareness campaign such as the “Get Tested” concept introduced by the CDC that recommends all individuals between the ages of 13 and 64 be tested for HIV at least once as part of routine health care - and with higher frequency if high risk factors are identified - are prudent next-steps toward improving the health status of HIV infected individuals. In addition, awareness programs and national cohort studies would help initiate early HIV treatment interventions. Based on this cohort study, future investigations into the comorbidities of well-controlled HIV infected individuals as well as those with co-infections (HBV, HCV and MTB) in the KSA and MENA region are warranted.

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Abbreviations

| Abbreviation | Definition |
|--------------|------------|
| CDC          | Centers for Disease Control and Prevention |
| HAART        | Highly Active Antiviral Therapy |
IVDU  Intravenous Drug Users

MENA  Middle East and North Africa (refers to the following countries: Bahrain, Egypt, Iran, Iraq, Palestine, Jordan, Kuwait, Lebanon, Libya, Morocco, Oman, Qatar, KSA, Syria, Tunisia, UAE, and Yemen)

KSA  Kingdom of Saudi Arabia

PCR  Polymerase Chain Reaction

References

1. May MT, Ingle SM, Costagliola D, Justice AC, de Wolf F, et al. Cohort profile: Antiretroviral Therapy Cohort Collaboration (ART-CC). Int J Epidemiol. 2014; 43:691–702. [PubMed: 23599235]

2. Crystal S, Sambamoorthi U, Merzel C. The diffusion of innovation in AIDS treatment: zidovudine use in two New Jersey cohorts. Health Serv Res. 1995; 30:593–614. [PubMed: 7591783]

3. Egger M, May M, Chêne G, Phillips AN, Ledergerber B, et al. Prognosis of HIV-1-infected patients starting highly active antiretroviral therapy: a collaborative analysis of prospective studies. Lancet. 2002; 360:119–129. [PubMed: 12126821]

4. Coetze D, Hildebrand K, Boulle A, Maartens G, Louis F, et al. Outcomes after two years of providing antiretroviral treatment in Khayelitsha, South Africa. AIDS. 2004; 18:887–895. [PubMed: 15060436]

5. Farmer P, Léandre F, Mukherjee J, Gupta R, Tarter L, et al. Community-based treatment of advanced HIV disease: introducing DOT-HAART (directly observed therapy with highly active antiretroviral therapy). Bull World Health Organ. 2001; 79:1145–1151. [PubMed: 11799447]

6. Kumarasamy N, Solomon S, Chaguturu SK, Mahajan AP, Flanigan TP, et al. The safety, tolerability and effectiveness of generic antiretroviral drug regimens for HIV-infected patients in south India. AIDS. 2003; 17:2267–2269. [PubMed: 14523288]

7. Bourgeois A, Laurent C, Mougnotou R, Nkoué N, Lactuock B, et al. Field assessment of generic antiretroviral drugs: a prospective cohort study in Cameroon. Antivir Ther. 2005; 10:335–341. [PubMed: 15865228]

8. Chene G. Prognostic importance of initial response in HIV-1 infected patients starting potent antiretroviral therapy: analysis of prospective studies. Lancet. 2003; 362:679–686. [PubMed: 12957089]

9. Zhou J, Kumarasamy N, Ditangco R, Kamarulzaman A, Lee CK, et al. The TREAT Asia HIV Observational Database: baseline and retrospective data. J Acquir Immune Defic Syndr. 2005; 38:174–179. [PubMed: 15671802]

10. Shawky S, Soliman C, Kassak KM, Oraby D, El-Khoury D, et al. HIV surveillance and epidemic profile in the Middle East and North Africa. J Acquir Immune Defic Syndr. 2009; 51(Suppl 3):S83–95. [PubMed: 19553783]

11. Obermeyer CM. HIV in the Middle East. BMJ. 2006; 333:851–854. [PubMed: 17053246]

12. Kingston ME, Harder EJ, Al-Jaberi MM, Bailey TM, Roberts GT, et al. Acquired immune deficiency syndrome in the Middle East from imported blood. Transfusion. 1985; 25:317–318. [PubMed: 4024227]

13. Harfi HA, Fakhry BM. Acquired immunodeficiency syndrome in Saudi Arabia. The American-Saudi connection. JAMA. 1986; 255:383–384. [PubMed: 3941519]

14. Madani TA, Al-Mazrou YY, Al-Jeffri MH, Al Huzaim NS. Epidemiology of the human immunodeficiency virus in Saudi Arabia: 18-year surveillance results and prevention from an Islamic perspective. BMC Infect Dis. 2004; 4:25. [PubMed: 15298719]

15. Al-Mazrou YY, Al-Jeffri MH, Fidail AI, Al-Huzaim N, El-Gizouli SE. HIV/AIDS epidemic features and trends in Saudi Arabia. Ann Saudi Med. 2005; 25:100–104. [PubMed: 15977685]
16. 1993 revised classification system for HIV infection and expanded surveillance case definition for AIDS among adolescents and adults. MMWR Recomm Rep. 1992; 41:1–19.
17. Mazroa MA, Kabbash IA, Felemban SM, Stephens GM, Al-Hakeem RF, et al. HIV case notification rates in the Kingdom of Saudi Arabia over the past decade (2000-2009). PLoS One. 2012; 7:e45919. [PubMed: 23049892]
18. Elmir E, Nadia S, Ouafae B, Rajae M, Amina S, et al. HIV epidemiology in Morocco: a nine-year survey (1991-1999). Int J STD AIDS. 2002; 13:839–842. [PubMed: 12537738]
19. DeJong J, Jawad R, Mortagy I, Shepard B. The sexual and reproductive health of young people in the Arab countries and Iran. Reprod Health Matters. 2005; 13:49–59. [PubMed: 16035597]
20. Gray PB. HIV and Islam: is HIV prevalence lower among Muslims? Soc Sci Med. 2004; 58:1751–1756. [PubMed: 14990375]
21. Schwarcz L, Chen MJ, Vittinghoff E, Hsu L, Schwarz S. Declining incidence of AIDS-defining opportunistic illnesses: results from 16 years of population-based AIDS surveillance. AIDS. 2013; 27:597–605. [PubMed: 10988097]
22. Mocroft A, Katlama C, Johnson AM, Pradier C, Antunes F, et al. AIDS across Europe, 1994-98: the EuroSIDA study. Lancet. 2000; 356:291–296. [PubMed: 11071184]
23. Baril L, Jouan M, Agher R, Cambau E, Caumes E, et al. Impact of highly active antiretroviral therapy on onset of Mycobacterium avium complex infection and cytomegalovirus disease in patients with AIDS. AIDS. 2000; 14:2593–2596. [PubMed: 11101072]
24. Bacellar H, Muñoz A, Miller EN, Cohen BA, Besley D, et al. Temporal trends in the incidence of HIV-1-related neurologic diseases: Multicenter AIDS Cohort Study, 1985-1992. Neurology. 1994; 44:1892–1900. [PubMed: 7936243]
25. Alarcón JO, Freimanis-Hance L, Krauss M, Reyes MF, Cardoso CA, et al. Opportunistic and other infections in HIV-infected children in Latin America compared to a similar cohort in the United States. AIDS Res Hum Retroviruses. 2012; 28:282–288. [PubMed: 21902581]
26. Naba MR, Kanafani ZA, Awar GN, Kanj SS. Profile of opportunistic infections in HIV-infected patients at a tertiary care center in Lebanon. J Infect Public Health. 2010; 3:130–133. [PubMed: 20869674]
27. Al-Mozaini et al. Page 9
28. Gortmaker SL, Hughes M, Cervia J, Brady M, Johnson GM, et al. Effect of combination therapy including protease inhibitors on mortality among children and adolescents infected with HIV-1. N Engl J Med. 2001; 345:1522–1528. [PubMed: 11794218]
29. Candiani TM, Pinto J, Cardoso CA, Carvalho IR, Dias AC, et al. Impact of highly active antiretroviral therapy (HAART) on the incidence of opportunistic infections, hospitalizations and mortality among children and adolescents living with HIV/AIDS in Belo Horizonte, Minas Gerais State, Brazil. Cad Saude Publica. 2007; 23:414–423.
34. Duan S, Guo HY, Pang L, Yuan JH, Jia MH, et al. Analysis of the epidemiologic patterns of HIV transmission in Dehong prefecture, Yunnan province. Zhonghua Yu Fang Yi Xue Za Zhi. 2008; 42:866–869. [PubMed: 19141217]

35. Robbins HA, Shiels MS, Pfeiffer RM, Engels EA. Epidemiologic contributions to recent cancer trends among HIV-infected people in the United States. AIDS. 2014; 28:881–890. [PubMed: 24300545]
Figure 1. Incidence of new HIV-1 infection per year, stratified by risk group

Numbers of new HIV-1 infections reported between 1984 and 2010, and stratified by mode of transmission as collected through chart review.
Figure 2. Proportion and absolute frequency of HIV-1 patients treated with antiretroviral agents (ART)

The absolute number of patients are plotted by year with percentage of patients on ART treatment represented by line chart. Zidovudine (AZT) monotherapy began in 1989 (n=20). Between 1990 and 1996, two NRTI combination was used (n=56). Beginning in 1996, a combination of three or four different antiretroviral drugs (HAART) was administered.
Figure 3. Absolute and person-year death rate from time of diagnosis

Absolute number of HIV positive patient deaths per year (bar chart) compared to the death per 100 person-years per year (line chart).
Table 1
Socio-demographic characteristics of the HIV cohort (total of 602 patients).

| Gender          | Number (Percentile) |
|-----------------|---------------------|
| Female          | 182 (30.23 %)       |
| Male            | 420 (69.76 %)       |

| Age             | Number (Percentile) |
|-----------------|---------------------|
| Adults          | 507 (84.21 %)       |
| Pediatric       | 93 (15.44 %)        |

| Marital Status  | Number (Percentile) |
|-----------------|---------------------|
| Married         | 290 (48.25 %)       |
| Single          | 239 (39.77 %)       |
| Widowed         | 48 (7.99 %)         |
| Divorced        | 24 (3.99 %)         |

| Partner testing | Number (Percentile) |
|-----------------|---------------------|
| Positive        | 182 (30.23 %)       |
| Negative        | 121 (20.10 %)       |
| Unknown         | 299 (49.67 %)       |

| KSA Province    | Number (Percentile) |
|-----------------|---------------------|
| Central Province| 196 (32.56 %)       |
| Southern Province| 161 (26.74 %)   |
| Western Province | 131 (21.76 %)     |
| Eastern Province | 94 (15.61 %)      |
| Northern Province| 19 (3.16 %)       |
| Unknown         | 1 (0.17 %)          |

| Outcome         | Number (Percentile) |
|-----------------|---------------------|
| Alive           | 372 (62%)           |
| Death           | 147 (24%)           |
| Lost follow up  | 83 (14%)            |

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Table 2

Mode of HIV transmission

| Transmission                  | Number (Percentile) | Female: % & Male: % |
|-------------------------------|---------------------|---------------------|
| Hemophilia receiving blood products | 60 (9.97%)          | 0 (0%) & 60 (9.97%) |
| Blood Transfusion             | 86 (14.29%)         | 43 (7.14%) & 43 (7.14%) |
| Heterosexual                  | 329 (54.65%)        | 102 (16.94%) & 227 (37.71%) |
| MSM                           | 10 (1.66%)          |                    |
| Bisexual                      | 20 (3.32%)          | 0 (0%) & 20 (3.32%) |
| Perinatal                     | 59 (9.80%)          | 31 (5.15%) & 28 (4.65%) |
| Intravenous Drug Use (IVDU)   | 17 (2.82%)          | 0 (0%) & 17 (2.82%) |
| Organ Transplantation         | 7 (1.16%)           |                    |
| Unknown                       | 53 (10.1%)          | 5 (0.83%) & 48 (7.97%) |
Table 3
Mortality rate for opportunistic infections and HIV-associated malignancies

| Opportunistic Infections       | Total Number of Cases | Number of Death | Percentile (%) |
|--------------------------------|-----------------------|-----------------|----------------|
| Pneumocystis jirovecii         | 50                    | 4               | 8              |
| Candida esophagitis            | 45                    | Not recorded    | Not recorded   |
| Tuberculosis (MTB)             | 29                    | 9               | 31             |
| Cytomegalovirus (CMV)          | 29                    | 13              | 44.8           |
| Cryptosporidiosis              | 7                     | 4               | 57             |
| Cryptococcal                   | 6                     | 3               | 50             |
| Associated Malignancy          |                       |                 |                |
| Kaposi sarcomas                | 18                    | 3               | 17             |
| Lymphoma                       | 6                     | 0               | 0              |