



Article Risk Factors Associated with Opportunistic Infections among People Living with HIV/AIDS and Receiving an Antiretroviral Therapy in Gabon, Central Africa

Augustin Mouinga-Ondeme ^{1,*}, Neil Michel Longo-Pendy ², Ivan Cyr Moussadji Kinga ³, Barthélémy Ngoubangoye ³, Pamela Moussavou-Boundzanga ¹, Larson Boundenga ^{2,4}, Abdoulaye Diane ¹, Jeanne Sica ⁵, Ivan Sosthene Mfouo-Tynga ^{1,†} and Edgard Brice Ngoungou ^{6,†}

- ¹ Unité des Infections Rétrovirales et Pathologies Associées, Centre Interdisciplinaire de Recherches Médicales de Franceville (CIRMF), Franceville BP 769, Gabon; pamelamoussavoub@gmail.com (P.M.-B.); dianeyabdoulaye@gmail.com (A.D.); tivansdavids2012@gmail.com (I.S.M.-T.)
- ² Unité de Recherches en Ecologie de la Santé, Centre Interdisciplinaire de Recherches Médicales de Franceville (CIRMF), Franceville BP 769, Gabon; longo2michel@gmail.com (N.M.L.-P.); boundenga@gmail.com (L.B.)
- ³ Centre de Primatologie, Centre Interdisciplinaire de Recherches Médicales de Franceville (CIRMF), Franceville BP 769, Gabon; cyrth01@yahoo.fr (I.C.M.K.); genistha@hotmail.com (B.N.)
- ⁴ Département d'Anthropologie, Université de Durham, South Road, Durham DH1 3LE, UK
- ⁵ Centre de Traitement Ambulatoire, Franceville BP 277, Gabon; jeannesic@yahoo.fr
- ⁶ Département d'Epidémiologie, Biostatistiques et Informatique Médicale (DEBIM)/Unité de Recherche en Epidémiologie des Maladies Chroniques et Santé Environnement (UREMCSE), Faculté de Médecine, Université des Sciences de la Santé, Libreville-Owendo BP 18231, Gabon; ngoungou2001@yahoo.fr
- Correspondence: ondeme@yahoo.fr
- These authors contributed equally to this work.

Abstract: The Human Immunodeficiency Virus/Acquired Immunodeficiency Syndrome (HIV/AIDS) is still one of the main causes of death in sub-Saharan Africa. Antiretroviral therapies (ARTs) have significantly improved the health conditions of people living with HIV/AIDS (PLWHA). Consequently, a significant drop in morbidity and mortality, along with a reduced incidence of opportunistic infections (OIs), has been observed. However, certain atypical and biological profiles emerge in ART patients post-examination. The objective of this study was to identify the risk factors that contributed to the onset of OIs in HIV patients undergoing ART in Gabon. Epidemiological and biological data were obtained from medical records (2017 to 2019) found at the outpatient treatment centre (CTA) of Franceville in Gabon. Samples for blood count, CD4, and viral load analysis at CIRMF were collected from PLWHA suffering from other pathogen-induced conditions. A survey was carried out and data were analysed using Rstudio 4.0.2 and Excel 2007 software. Biological and socio-demographic characteristics were examined concerning OIs through both a univariate analysis via Fisher's exact tests or chi² (χ^2), and a multivariate analysis via logistic regression. Out of the 300 participants initially selected, 223 were included in the study, including 154 (69.05%) women and 69 (30.95%) men. The mean age was 40 (38.6; 41.85), with individuals ranging from 2 to 77 years old. The study cohort was classified into five age groups (2 to 12, 20 to 29, 30 to 39, 40 to 49, and 50 to 77 years old), among which the groups aged 30 to 39 and 40 to 49 emerged as the largest, comprising 68 (30.5%) and 75 (33.6%) participants, respectively. It was noted that 57.9% of PLWHA had developed OIs and three subgroups were distinguished, with parasitic, viral, and bacterial infections present in 18%, 39.7%, and 55.4% of cases, respectively. There was a correlation between being male and having a low CD4 T-cell count and the onset of OIs. The study revealed a high overall prevalence of OIs, and extending the study to other regions of Gabon would yield a better understanding of the risk factors associated with the onset of these infections.

Keywords: HIV-1; PLWHA; opportunistic infections; Gabon



Citation: Mouinga-Ondeme, A.; Longo-Pendy, N.M.; Moussadji Kinga, I.C.; Ngoubangoye, B.; Moussavou-Boundzanga, P.; Boundenga, L.; Diane, A.; Sica, J.; Mfouo-Tynga, I.S.; Ngoungou, E.B. Risk Factors Associated with Opportunistic Infections among People Living with HIV/AIDS and Receiving an Antiretroviral Therapy in Gabon, Central Africa. *Viruses* 2024, 16, 85. https://doi.org/ 10.3390/v16010085

Academic Editor: Carlo Contini

Received: 19 October 2023 Revised: 11 November 2023 Accepted: 11 November 2023 Published: 4 January 2024



Copyright: © 2024 by the authors. Licensee MDPI, Basel, Switzerland. This article is an open access article distributed under the terms and conditions of the Creative Commons Attribution (CC BY) license (https:// creativecommons.org/licenses/by/ 4.0/).

1. Introduction

The vulgarization of antiretroviral therapy (ART) has improved the living conditions of people living with HIV/AIDS (PLWHA) and has led to a significant decrease in morbidity and mortality [1–3]. Factors related to ART—late initiation, discontinuation, poor adherence, low CD4 T lymphocyte count, inadequate virological monitoring, isoniazid preventive therapy, gender, age, place of residence, and functional or disclosure or nutritional status—have been associated with the emergence of opportunistic infections (OIs) among adults (PLWHA) post-ART [4–7]. Frequently, OIs occur at all stages of AIDS, leading to increased gravity of the condition and worse health status of PLWHA [8]. This constitutes a major cause of hospitalization, and subsequent death in patients, highlighting the issue of management [9]. The prevalence of OIs among ART-treated adults varies and depends on the specificity of each country; for example, it was estimated at 22.4% in the south-east of Nigeria, from 33.6 to 88.44% in Ethiopia, and at 33.51% in seven provinces of Indonesia [10,11]. Some of the commonly described OIs in PLWHA are salmonella infection, chronic diarrhoea, candidiasis, toxoplasmosis, tuberculosis, dermatitis, and pneumonia [1].

At the end of 2022, the World Health Organization (WHO) estimated that 39 million people were HIV-infected and 650,000 died from AIDS-related illnesses worldwide; the fatalities were mainly due to OIs [12]. During the last Demographic and Health Survey (DHS), HIV-1 infection was reported in 4.1% of the general population aged 15 to 49 years old in Gabon [13]. The distribution of HIV-1 across the country showed that the Haut-Ogooué province, in the south-eastern part of the country, was one of the most affected regions, with 4.2% of HIV-1 infected people, and around 54% of PLWHA receiving ART in Gabon [13]. Treatment administration in this country follows WHO recommendations for resource-limited settings/countries and includes generalized first-line ART regimens: Recommended first-line regimens until 2019 included two nucleoside reverse transcriptase inhibitors plus one non-nucleoside reverse transcriptase inhibitor (2NRTI + 1 NNRTI) [14]. Due to mutations associated with drug resistance, WHO recommended a transition to integrase strand transfer inhibitor (INSTI)-based first-line regimens [15]. Studies have been conducted on HIV in co-infection with other pathogens in Gabon, but there is only one study, conducted in Libreville (capital city), highlighting OIs among PLWHA [16]. No study on OIs was found concerning PLWHA in rural or semi-rural areas, including the Haut-Ogooué province of Gabon. Recently, we reported a high prevalence of HTLV-1/HIV-1 co-infection among PLWHA followed-up at the outpatient or ambulatory treatment centre (CTA) of Franceville, the major city of Haut-Ogooué province, Gabon [17]. The need for preventive strategies to effectively manage the scenarios of co-infections in PLWHA was clearly identified and recommended.

Our intent was to assess the incidence and associated risk factors of OIs, so we considered the HIV-1-infected patients receiving ART monitoring at the CTA of Franceville. In this sanitary facility, specific care has been provided for several HIV/AIDS patients of all ages, both male and female, including pregnant women, from the entire Haut-Ogooué province.

2. Materials and Methods

2.1. Study Design and Sampling

From January to June 2020, we conducted a retrospective and cross-sectional study of PLWHA who were monitored at the CTA in Franceville, a city of roughly 7000 inhabitants in the south-eastern region of Gabon. Patients were recruited between June 2018 and September 2019 to perform a one-time CD4 count and HIV-1 RNA viral load measurement at diagnosis (as described in Section 2.3) free of cost to patients at the Centre Interdisciplinaire de Recherches Médicales de Franceville (CIRMF). According to medical records (2017–2019) filed at the CTA of Franceville, the recruitment criteria were as follows: (i) being confirmed HIV-1-positive, (ii) being on antiretroviral therapy (ART), and (iii) providing informed consent. The additional information collected included gender, age, ART regimen, and CD4 count at ART initiation. Patients who did not show a CD4 count and/or viral

load during diagnosis of the opportunistic diseases, and those who had interrupted their follow-up at the CTA, were excluded from the study. Figure 1 presents the timeline of the study.

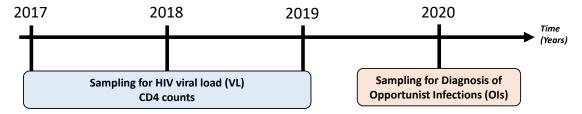


Figure 1. Schematic representation of the timeline of the study. Progress in carrying out examinations on the people living with HIV/AIDS (PLWHA) throughout the study period.

2.2. Ethical Consideration

The study was approved by the Gabon National Ethics Committee for Research (approval registered as PROT N°0011/2013/SG/CNE) and all participants gave verbal consent before participation in the study.

2.3. Sampling for HIV Viral Load and CD4 Counts

For determining HIV viral load (VL) and CD4 counts, blood specimens were collected from each patient using 5 mL EDTA tubes and transferred to a reference national laboratory, located at the Unit of Retroviral Infection and Associated Pathology, CIRMF. On arrival at the facility, samples were centrifuged, and then plasma and buffy coat fractions were collected and stored at -80 °C until testing, as previously described [17]. Specifically, CD4 counts at enrolment were determined using flow cytometry (Fascount, Becton Dickinson, San Jose, CA, USA). Plasma HIV-1 VL was determined using the Generic HIV Viral Load test (Biocentric, Bandol, France) performed according to the manufacturer's instructions, using 200 μ L of plasma and the protocol with a detection limit of 300 copies/mL [17,18].

2.4. Diagnosis of Other Pathogenic (Viral, Parasitic, and Bacteriological) Infections

The search for opportunistic infections (OIs) was carried out in 2020 in patients initially recruited between 2017 and 2019 (as shown in Figure 1). Plasma samples from patients were screened for hepatitis B virus infection (HBV surface antigen [19]) by simultaneously using a rapid diagnosis test (RDT) (AgHBs 2 DETERMINETM, Abbott Rapid Diagnostics Ltd, Stockport, UK) and an automated quantitative test (ù). Similarly, the screening for hepatitis C virus (HCV) infection was based on the antibodies anti-HCV by simultaneously using an RDT (Accu-Tell[®] HCV Rapid Test Cassette/Strip (Serum/Plasma), Accubiotech, Beijing, China) and an automated quantitative test (VIDAS[®] Anti-HCV (HCV), bioMérieux, Marcy-L'Etoile, France). For parasitic infection, toxoplasmosis serological analysis was performed with an automated quantitative test (VIDAS® Toxo IgG II and VIDAS® ToxoIgM kits, bioMérieux, Marcy-L'Etoile, France). The presence of IgG meant that the organism showed an acquired immunity with respect to Toxoplasma, while the presence of IgM indicated a sign of acute Toxoplasma infection. For bacteriological diagnosis, Chlamydia infection was screened via serological diagnosis, which evaluated the presence of IgG and IgM specific to Chlamydia, by using CHLAMYCHECK IgG and IgA. On the other hand, the "treponemal test" and the automated quantitative Elecsys[®] Syphilis for cobas E411 (Roche Diagnostics, Mannheim, Germany) were used for the diagnosis of syphilis. In case of positivity, a second test called the "non-treponemal test" (VDRL assay with the SIMPLI RPR CARBO kit from MEDIFF°, Aubagne, France) was performed to determine the status of infection (active or not). The direct diagnosis of gonorrhoea infection was performed via urethral cytological examination for male and endo-cervical or vaginal for female samples. An attempt to determine the presence of Gram-negative Cocci in diplococci "coffee bean", was performed using a bacteriological culture of genital samples on VCA3 medium [20]. Finally, tuberculosis (TB) diagnosis was performed by using chest X-rays, skin tests, and sputum examinations in a specialized care centre.

2.5. Occurrence of Opportunistic Infections

In order to emphasize the co-occurrence between HIV infection and OIs, a Venn diagram was generated using the "Venn Diagram" package. Furthermore, a binomial generalized linear model (GLM) was performed using the "stats" package to investigate the effect of seven factors as explanatory variables in the GLM, namely immunological (CD4 T lymphocyte count), virological (viral load), gender (sex), age, marital status, level of ART participation/consistency/lost to follow-up, and geographical origin, on the likelihood of OIs in PLWHA, which serves as the dependent variable in the model [21]. The GLM holds that any observed response is a linear sum of multiple individuals' underlying responses. The GLM was used to determine how the predetermined factors (7) might have combined and/or contributed to the occurrence of OIs.

2.6. Statistical Analysis

The information collected was recorded in an Excel file and analysis was conducted with R software version 4.0.2 (http://cran.r-project.org/bin/windows/base/old/4.0.2/ NEWS.R-4.0.2.html, accessed on 22 June 2020). The interaction level among variables (factors) was estimated with the Akaike information criterion (AIC) for model selection, and the AIC changes were analysed by utilizing the dredge function within the MuMIn package [21]. As a consequence of the AIC change analysis, only the variables (two factors) that had an impact on the incidence of OIs were chosen and included in the model, out of the original seven explanatory variables. The model with the lowest AIC was considered the best, while Δ AIC values below 2 were also retained [22]. Subsequently, we conducted the chi² significance test (*p* < 0.05) to compare the proportional variance among the various forms of OIs.

3. Results

3.1. Characteristics of Patients

Of the 300 patients selected, 223 were included in the study; the participants had a median age of 41 (IQR 27.5–54.5) and their ages ranged from 2 to 77 years old, comprising 69 (30.9%) males and 154 (69.1%) females. The study cohort was categorised into five age groups (2 to 12, 20 to 29, 30 to 39, 40 to 49, and 50 to 77 years old) with the two most populated age groups (30 to 39 and 40 to 49) emerging as prominent, comprising 68 (30.5%) and 75 (33.6%) participants, respectively. Participants had their CD4 T-cell counts available upon inclusion, and the median count was determined to be 419 (93–745). The median duration of ART was 20 months, and all participants were taking ART as a first line of treatment. Two ART regimens were the most prevalent, consisting of zidovudine (AZT) plus lamivudine (3TC) along with efavirenz (EFV)/nevirapine (NVP) (25.8%), and tenofovir (TDF) plus 3TC/emtricitabine (FTC) plus EFV (15%) (Table 1).

Table 1. Patient characteristics associated with opportunistic infections.

Variables		OIs Prevalence	Virological OIs	Bacteriological OIs	Parasitic OIs
Gender	n(%)	n/N (%)	n (% [95% CI])	n (% [95% CI])	n (% [95% CI])
Men Women	69 (30.9) 154 (69.1)	47/69 (68.1) 82/154 (53.2)	11 (16 [8.6–27.2]) 23 (15 [9.9-21.8])	35 (50.7 [38.5–62.9]) 58 (37.7 [30.1–45.9])	11 (16 [8.6–27.2]) 21 (13.6 [8.8–20.3])
Total	223 (100)	129/223 (57.9)	34 (15 [11–20.8])	93 (41.7 [35.2–48.5])	32 (14.3 [10.2–19.8])
Age (Years)					
2–12	9(4.1)	6/9 (66.7)	1 (11 [0.6–49.3])	4 (44.4 [15.3–77.3])	3 (33.3 [9–69.1])
20–29	27(12.1)	19/27 (70.4)	7 (26 [11.9–46.6])	13 (48.1 [29.2–67.6])	4 (14.8 [4.9–34.6])
30–39	68(30.5)	35/68 (51.5)	14 (20.6 [12.1–32.5	19 (28 [18.1–40.3])	6 (8.8 [3.6–18.9])
40-49	75(33.6)	42/75 (56)	8 (10.7 [5–20.5])	37 (49.3 [37.7–61])	11 (14.7 [7.9–25.2])
50-77	44(19.7)	27/44 (61.4)	4 (9.1 [3–22.6])	20 (45.5 [30.7–61])	8 (18.2 [8.7–33.2])
Median (IQR) 41 (27.5–54.5)	. /	,			

Variables		OIs Prevalence	Virological OIs	Bacteriological OIs	Parasitic OIs
CD4 cell count, cells/mm ³ <200 [200–499] ≥500 Median (IQR) 419 (93–745)	35 (15.7) 127 (57) 61 (27.3)	24/35 (68.6) 88/127 (69.3) 17/61 (27.9)	8 (22.9 [11–40.6]) 22 (17.3 [11.4–25.3]) 4 (6.6 [2.1–16.7])	20 (57.1 [39.5–73.2) 62 (48.8 [40–57.8]) 11 (18 [9.8–30.4])	8 (22.9 [11–40.6]) 22 (17.3 [11.4–25.3]) 2 (3.3 [0.6–12.4])
ART regimen AZT – 3TC – EFV TDF – 3TC – EFV AZT + 3TC + NVP TDF + FTC + EFV Others	94 (42.2) 13 (5.8) 21 (9.4) 54 (24.2) 41 (18.4)	54/94 (57.4) 8/13 (61.5) 10/21 (47.6) 31/54 (57.4) 26/41 (63.4)	15 (16 [9.5–25.3]) 2 (15.4 [2.7–46.3]) 4 (19 [6.3–42.6]) 7 (13 [5.8–25.5]) 6 (14.6 [6.1–29.9])	40 (42.6 [32.5–53.2]) 8 (61.5 [32.3–84.9]) 5 (23.8 [9.1–47.5]) 22 (40.7 [27.9–54.9]) 18 (44 [28.8–60.1])	16 (17 [10.3–26.5]) 1 (7.7 [0.4–38]) 2 (9.5 [1.7–31.8]) 7 (13 [5.8–25.5]) 6 (14.6 [6.1–30])

Table 1. Cont.

3TC: lamivudine, EFV: efavirenz, TDF: tenofovir, AZT: zidovudine, FTC: emtricitabine, NVP: nevirapin.

3.2. Prevalence, Distribution, and Spectrum of Opportunistic Infections

Approximately 57.9% of PLWHA were diagnosed with OIs, and pulmonary TB (55.4%) was the most prevalent OI after ART initiation. This condition is considered as the primary bacterial infection. The viral (HBV and HCV) infections (39.7%) and parasitic infections (mainly oral candidiasis; 18%), ranked second and third, respectively (Table 1).

3.3. Different Combinations of Opportunistic Infections among People Living with HIV

The Venn diagram (Figure 2) and proportions comparison indicate that the populace consists of HIV-infected patients linked with other OIs (57.9%) against HIV-mono-infected patients ((42.1%) (Chi² = 10.368, *p*-value = 0.001)). In OIs in HIV bi-infections, bacterial conditions illnesses were more frequent, followed by virus-induced ones (Chi² = 78.624, *p*-value < 0.0001). Interestingly, the comorbidity made up of tri-infections of HIV, bacteria, and parasites was the most prevalent, with only two individuals in the studied population being tetra-infected with HIV, bacteria, virus, and parasites (Figure 2).

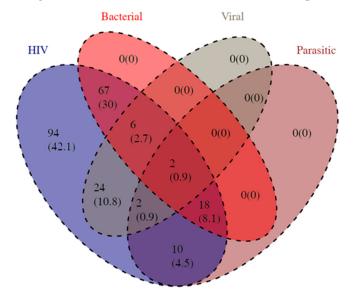


Figure 2. Different combinations of opportunistic infections among people living with HIV. The Venn diagram illustrates the proportions of individuals infected with HIV only (blue), those afflicted with bacterial infections only (red), those with viral infections only (grey), and those diagnosed with parasitic infections only (indigo/purple). The mixed colours indicated the proportions of PLWHA co-infected with HIV and one OI group, two OI groups, three OI groups, or different combinations between OI groups. The resulting and merged intermediate colours reflect the proportions and combinations observed.

The incidence of OIs was higher among men (68.1%) than women (53.2%). Moreover, the age group consisting of 20-29-year-old individuals had the highest OI prevalence rate, estimated at 70.4%. Participants with a CD4 T-cell count value of [200-499] and <200 cells/mm³ exhibited a greater likelihood of developing OIs than those with higher CD4 T-cell count values (value > 500 cells/mm³). Similarly, a relatively high incidence of OIs was observed in patients who were administered TDF-3TC-EFV as their ART regimen (61.5%), closely followed by TDF + FTC + EFV and AZT-3TC-EFV, with an estimated rate of 57.4% for both.

The best model ($\Delta AIC = 0$) resulting from our generalized linear model (GLM) indicates that two out of the seven sociobiological factors identified among PLWHA receiving ART were linked to the development of OIs. These factors were identified as gender and the initial value of the CD4 T-cell count (Table 2 and Figure 3). Consequently, male patients (Figure 3A) with suboptimal CD4 lymphocyte levels (Figure 3B) had a greater risk of developing OIs.

Table 2. Predictive model explaining the occurrence of opportunistic infections.

Model No. (Rank) –	Fixed Effects			Dí		Alcoilco Maiobt
	Intercept	CD4 Rate	Sex	– Df	ΔΑΙΟ	Akaike Weight
4 (1)	1.534	-0.0031	+	3	0	0.587
2 (2)	1.726	-0.0032		2	0.71	0.413

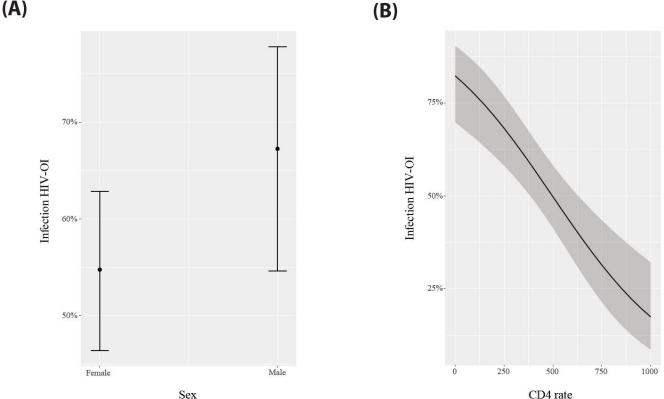


Figure 3. Probability of occurrence of opportunistic infections among people living with HIV. The combined effect of being male (A) with a low CD4 T-cell count (B) appears to be the most critical joint-risk factor associated with the development of opportunistic infections (OIs).

(B)

4. Discussion

In this study, we describe opportunistic infections (OIs) among people living with HIV/AIDS (PLWHA) in the south-east region of Gabon, with the majority of participants consisting of women, as reported in previous studies on this population. Most HIV-1 infected patients receiving ART in that CTA in Gabon are women, and there is no new finding in this regard [17,18,23]. Previous studies conducted in other countries on a similar type (female vs. male) of population have also described that women were the predominant gender [24,25]. We found that 57.9% of PLWHA had developed OIs, of which 18% were parasitic, 39.7% viral, and 55.4% bacterial infections. Being a male with a low CD4 T-cell count are the main joint-risk factors associated with the development of OIs.

The same population has been considered to study other pathogen-induced infections, including hepatitis viruses (HCV), and human T-lymphotropic virus type 1 (HTLV-1) [17]. The prevalence rate of 57.9% for OIs found among infected people confirms the vulnerability of PLWHA in Gabon with a high risk of exposure to viral, parasitic, and bacteriological agents. This reported rate is higher than those previously described in Gabon [16,26]. There is regularly a shortage of ARVs experienced in Gabon, and that might assuredly constitute a significant contributor and possible explanation for the high rates of OIs. Furthermore, starting ART at later stages of HIV infection, on the one hand, and the disparity observed in the level of engagement in HIV care facilities in poor countries or/and resource-limited areas, on the other hand, limit the effectiveness of the administered therapy [1,23]. Additionally, the emerging or increasing levels of HIV drug resistance to NNRTIs in resource-limited countries could also be considered as restraining factors [15]. At the time of the study, all participants were receiving the recommended first-line regimens, which included 2 NRTI + 1 NNRTI in Gabon. The obtained prevalence rate of OIs is comparable to the 55.3% that was reported among HIV/AIDS patients on ART in the southern zone Tigray, and the 60.97% rate in the western part of Ethiopia [24,27]. However, it is higher than the prevalence reported in similar studies conducted in other regions of Ethiopia, and in Nigeria and Taiwan [10,28–32]. In addition, our findings are lower than the 83.81% and 88.44% rates reported among HIV/AIDS patients in other distinctive parts of Ethiopia [5].

Analysis of bacteriological, viral, and parasitic OIs found that pulmonary TB, HBV, HCV, and oral candidiasis were the commonly occurring OIs in the studied population of Franceville. Bacteriological infections represented only by pulmonary TB (55.4%) were the most prevalent OI among PLWHA. Viral infections with HBV and HCV (39.7%) and parasitic infections with oral candidiasis (18%) ranked second and third, respectively. This finding about pulmonary TB in our study was congruent with another study conducted in Libreville, the Capital city of Gabon, where this OI was reported as the major cause of morbidity [16]. Other previous studies conducted in Gabon showed that the prevalence rate of TB/HIV co-infection was estimated at 26% to 42% [33–36]. Pulmonary TB is considered a leading cause of death in PLWHA, and having the highest prevalence reported in the current study highlights the risk related to this vulnerable population progressing rapidly towards the AIDS stage and developing further complications [37]. These findings emphasize the need for more surveillance and delicate care of PLWHA.

Viral OIs including HBV and HCV were the second most prevalent and induced diseases in the current study with a rate of 39.7%. Both hepatitis viruses are prevalent in Gabon and have already been described in co-infection with HIV [19,38,39]. The HIV/HCV and HIV/HBV co-infections could result from unprotected sexual intercourse; these diseases have been shown to be transmitted through unprotected sex with infected partners [40]. As in mono-infection, viral hepatitis B- and C-related diseases were estimated to be responsible for a significantly high number of deaths [41]. HBV and HCV infections can cause acute and chronic hepatitis, leading to liver fibrosis, cirrhosis, hepatocellular carcinoma (HCC), and end-stage liver disease over time [42]. In the era of ART, these diseases are considered to account for an increased probability of deaths among PLWHA [43–45]. Therefore, continuous surveillance must be conducted among PLWHA to clarify the current status of hepatitis viral diseases as well as parasitic infections. Parasitic infection, or oral candidiasis (18%), is one of the main OIs, together with the above-mentioned ones. Generally, oral candidiasis is prevalent among PLWHA, as previously described in Gabon (14.19%), as well as reported in India and Ethiopia [16,25,46]. The prevalence rate reported in the present study was higher than that previously reported in Gabon (14.19% vs. 18%). All these findings could be explained by the difference between a relatively rural area (current study) and more urbanized settlements. This coincided with previous studies from Ethiopia and the USA, where it was demonstrated that rural/countryside patients with lower levels of education, low income, and limited access to healthcare facilities more readily develop OIs than others [25,47–49].

Our study showed that being a male with a low CD4 T-cell count constitutes a higher risk that is associated with the development of OIs. We have already indicated that most HIV-1 infected patients and monitored ART recipients at most care centres (CTA) in Gabon are women [17,18,23]. Despite receiving ART, OIs were reported in the current study. The fact that men are more likely to develop OIs could be explained by the decision to adhere to and receive ART at the latter stages or/and the poor adherence and subsequent side effects [1]. It is recognised that, unlike women who often attend healthcare facilities as quickly as possible when they get ill/diagnosed, their male counterparts are prompt to seek medical advice when facing health-related complications or severe cases [50]. Information collected from participants showed that most men unfortunately do not accept the reality of being infected with HIV/AIDS, and so delay taking and adhering to recommended actions. Not only do they refuse or delay ARTs or/and take them discontinuously, but sometimes they continue to engage in inappropriate conduct such as unprotected sex. The obvious consequence is the emergence of HIV-associated complications, and sometimes mutations with drug resistance reported in Africa, as it is mainly observed in Gabon [14,18]. In Gabon, ART guidelines at the time of this study were based on WHO recommendations for resource-limited settings/countries and included generalized first-line ART regimens: Recommended first-line regimens included 2 NRTI + 1 NNRTI [14]. Due to mutations associated with drug resistance, WHO recommended the transition to integrase strand transfer inhibitor (INSTI)-based first-line regimens [15]. The high prevalence of OIs among patients receiving TDF-3TC/FTC+EFV was probably also due to possible mutations associated with resistance. And the low CD4 T-cell count could be interrelated with treatment failure. CD4 T-cell counts of [200–499] and <200 cells/mm³ were reported in our study, and similar counts are most likely reported in cases of development of OIs. This result is reasonable as CD4 T-cells play critical roles in the induction of both humoral and cellular immunity to combat OIs [1]. Comparatively, other authors found that PLWHA with a CD4 T-cell count of 200 cells/mm³ were at higher risk of developing advanced forms of OIs such as pulmonary TB [51]. Also, previous studies described that there are patients who develop OIs despite the high CD4 T-cell count [52,53]. This finding was in concordance with our results, showing that the risk of developing OIs is not only due to low CD4 T-cell count. In fact, a high CD4 T-cell count could be an abnormal increase in case of co-infection between HIV-1 and HTLV-1 conferring false immunity to the patients, as recently reported in a similar population type [17].

One limitation of this study was the relatively small number of participants, in comparison to the entire active patient database of the CTA, with a significant number of "lost to follow-up" individuals, and thus not consistently monitored. Furthermore, the study was solely conducted in a semi-rural location of a single CTA, making it difficult to evaluate how the obtained results might translate to urban areas or other regions of the country with PLWHA. Hence, as this study was retrospective, certain characteristics of the patients that were not documented, including their level of knowledge, marital status, and professional category or geographical location could influence and skew the analysis of the results. Therefore, the necessity of improving the management of PLWHA will enhance treatment outcomes. In the future, it will be critically important to consider all aspects of OIs, drug resistance, patient behaviour regarding treatment compliance, and attendance at CTAs when devising strategies for the care of PLWHA in Gabon. Poor compliance or medication abandonment may also heighten the likelihood of OIs. When developing such approaches, the involvement of personnel such as health staff, researchers, social workers, politicians, and traditional practitioners should be considered as all of these people may influence and have a significant impact on the healthcare of those specific groups of PLWHA.

5. Conclusions

Our study reports a significant occurrence of opportunistic infections (OIs) amongst HIV-1-infected patients receiving first-line ART and being monitored at CTA of Franceville in the south-east part of Gabon. We examine the range and risk factors related to the development of these OIs. The major OIs observed in the study were pulmonary TB, hepatitis B and C, and oral candidiasis. Additionally, the development of OIs was linked to male gender and low CD4 T-cell count. It is imperative to devise strategies to tackle the burden of OIs and improve outcomes, specifically for those with severely weakened immune systems. Our research emphasises the necessity of offering enhanced monitoring and care for HIV-positive individuals. Furthermore, carrying out extensive investigations in various provinces across the country will aid in identifying and comprehending all OIs that are prevalent, ultimately enabling the development of national strategies to mitigate their impacts on the health of PLWHA.

Author Contributions: Conceptualization, A.M.-O., N.M.L.-P., I.S.M.-T. and E.B.N.; data curation, I.C.M.K. and A.D.; formal analysis, N.M.L.-P., I.C.M.K. and A.D.; investigation, I.C.M.K.; methodology, A.M.-O.; project administration, A.M.-O., I.S.M.-T. and E.B.N.; resources, J.S.; supervision, A.M.-O., B.N., I.S.M.-T. and E.B.N.; validation, A.M.-O., N.M.L.-P., I.S.M.-T. and E.B.N.; writing—original draft, A.M.-O., N.M.L.-P., B.N., P.M.-B., L.B., I.S.M.-T. and E.B.N.; writing—review and editing, A.M.-O., B.N., P.M.-B., L.B., I.S.M.-T. and E.B.N.; writing—to the published version of the manuscript.

Funding: This research received no funding. Our research is supported by the Centre Interdisciplinaire de Recherches Médicales de Franceville (CIRMF), Gabon. This Centre is supported by Total Gabon and the Gabonese Government.

Institutional Review Board Statement: Not applicable.

Informed Consent Statement: Not applicable.

Data Availability Statement: The data that support the findings of this study are available on request from the corresponding author.

Acknowledgments: We are grateful to all the members of our team, mainly OLOUMBOU Eldridge Fedricksen and ENGONE-ONDO Dimitri, for their support. We apologize for the numerous contributions that may not have been cited in this review.

Conflicts of Interest: The authors declare no conflict of interest.

References

- Woldegeorgis, B.Z.; Kerbo, A.A.; Obsa, M.S.; Mokonnon, T.M. A systematic review and meta-analysis of antimicrobial resistance knowledge, attitudes, and practices: Current evidence to build a strong national antimicrobial drug resistance narrative in Ethiopia. *PLoS ONE* 2023, *18*, e0287042. [CrossRef] [PubMed]
- Palella, F.J., Jr.; Baker, R.K.; Moorman, A.C.; Chmiel, J.S.; Wood, K.C.; Brooks, J.T.; Holmberg, S.D. Mortality in the highly active antiretroviral therapy era: Changing causes of death and disease in the HIV outpatient study. *J. Acquir. Immune Defic. Syndr.* 2006, 43, 27–34. [CrossRef]
- Mirani, G.; Williams, P.L.; Chernoff, M.; Abzug, M.J.; Levin, M.J.; Seage, G.R., 3rd; Oleske, J.M.; Purswani, M.U.; Hazra, R.; Traite, S.; et al. Changing Trends in Complications and Mortality Rates Among US Youth and Young Adults with HIV Infection in the Era of Combination Antiretroviral Therapy. *Clin. Infect. Dis.* 2015, *61*, 1850–1861. [CrossRef] [PubMed]
- Arefaine, Z.G.; Abebe, S.; Bekele, E.; Adem, A.; Adama, Y.; Brockmeyer, N.H.; Coenenberg, J.; Potthoff, A.; Gebremeskel, T.G. Incidence and predictors of HIV related opportunistic infections after initiation of highly active antiretroviral therapy at Ayder Referral Hospital, Mekelle, Ethiopia: A retrospective single centered cohort study. *PLoS ONE* 2020, *15*, e0229757. [CrossRef] [PubMed]

- Solomon, F.B.; Angore, B.N.; Koyra, H.C.; Tufa, E.G.; Berheto, T.M.; Admasu, M. Spectrum of opportunistic infections and associated factors among people living with HIV/AIDS in the era of highly active anti-retroviral treatment in Dawro Zone hospital: A retrospective study. *BMC Res. Notes* 2018, 11, 604. [CrossRef] [PubMed]
- Dagnaw Tegegne, K.; Cherie, N.; Tadesse, F.; Tilahun, L.; Kassaw, M.W.; Biset, G. Incidence and Predictors of Opportunistic Infections among Adult HIV Infected Patients on Anti-Retroviral Therapy at Dessie Comprehensive Specialized Hospital, Ethiopia: A Retrospective Follow-Up Study. *HIV AIDS* 2022, *14*, 195–206. [CrossRef] [PubMed]
- 7. Zhou, J.; Paton, N.I.; Ditangco, R. AIDS-defining illness diagnosed within 90 days after starting highly active antiretroviral therapy among patients from the TREAT Asia HIV Observational Database. *Int. J. STD AIDS* **2007**, *18*, 446–452. [CrossRef]
- 8. Cuvelier, M.L.; Léonard, P.; Rikir, E.; Belachew, S. When HIV gets into the brain. Rev. Med. Liege 2008, 63, 338–341.
- 9. Niriella, M.A.; Kasturiratne, A.; Pathmeswaran, A.; De Silva, S.T.; Perera, K.R.; Subasinghe, S.; Kodisinghe, S.K.; Piyaratna, T.; Vithiya, K.; Dassanayaka, A.S.; et al. Lean non-alcoholic fatty liver disease (lean NAFLD): Characteristics, metabolic outcomes and risk factors from a 7-year prospective, community cohort study from Sri Lanka. *Hepatol. Int.* **2019**, *13*, 314–322. [CrossRef]
- Iroezindu, M.O.; Agbaji, O.O.; Daniyam, C.A.; Isiguzo, G.C.; Isichei, C.; Akanbi, M.O. Liver function test abnormalities in Nigerian patients with human immunodeficiency virus and hepatitis B virus co-infection. *Int. J. STD AIDS* 2013, 24, 461–467. [CrossRef]
- 11. Putranti, A.; Asmarawati, T.P.; Rachman, B.E.; Hadi, U. Oral candidiasis as clinical manifestation of HIV/AIDS infection in Airlangga University hospital patients. *IOP Conf. Ser. Earth Environ. Sci.* **2018**, 125, 012063. [CrossRef]
- 12. UNAIDS. Dernières Statistiques sur L'état de L'épidémie de Sida. 13 July 2023. Available online: https://www.unaids.org/fr/resources/documents/2023/UNAIDS_FactSheet (accessed on 9 October 2023).
- 13. EDSG. Prévalence de L'infection à VIH au Gabon. 2012. Available online: https://gabon.unfpa.org/sites/default/files/pub-pdf/ EDS%20II.pdf (accessed on 9 October 2023).
- Engone-Ondo, J.D.; Mouinga-Ondémé, A.; Lékana-Douki, S.E.; Diané, A.; Mamimandjiami, A.I.; Banga, O.; Ndong-Atome, G.R.; Aghokeng, A.F. High rate of virological failure and HIV drug resistance in semi-rural Gabon and implications for dolutegravirbased regimen efficacy. J. Antimicrob. Chemother. 2021, 76, 1051–1056. [CrossRef]
- 15. WHO. Update of Recommendations on First- and Second-Line Antiretroviral Regimens; WHO: Geneva, Switzerland, 2019.
- Okome-Nkoumou, M.; Guiyedi, V.; Ondounda, M.; Efire, N.; Clevenbergh, P.; Dibo, M.; Dzeing-Ella, A. Opportunistic diseases in HIV-infected patients in Gabon following the administration of highly active antiretroviral therapy: A retrospective study. *Am. J. Trop. Med. Hyg.* 2014, 90, 211–215. [CrossRef] [PubMed]
- 17. Mouinga-Ondémé, A.; Boundenga, L.; Koumba Koumba, I.P.; Idam Mamimandjiami, A.; Diané, A.; Engone-Ondo, J.D.; Djuicy, D.D.; Sica, J.; Mombo, L.E.; Gessain, A.; et al. Human T-Lymphotropic virus type 1 and human immunodeficiency virus co-infection in rural Gabon. *PLoS ONE* **2022**, *17*, e0271320. [CrossRef] [PubMed]
- Liégeois, F.; Vella, C.; Eymard-Duvernay, S.; Sica, J.; Makosso, L.; Mouinga-Ondémé, A.; Mongo, A.D.; Boué, V.; Butel, C.; Peeters, M.; et al. Virological failure rates and HIV-1 drug resistance patterns in patients on first-line antiretroviral treatment in semirural and rural Gabon. J. Int. AIDS Soc. 2012, 15, 17985. [CrossRef]
- Bivigou-Mboumba, B.; Amougou-Atsama, M.; Zoa-Assoumou, S.; M'Boyis Kamdem, H.; Nzengui-Nzengui, G.F.; Ndojyi-Mbiguino, A.; Njouom, R.; François-Souquière, S. Hepatitis B infection among HIV infected individuals in Gabon: Occult hepatitis B enhances HBV DNA prevalence. *PLoS ONE* 2018, *13*, e0190592. [CrossRef]
- Van Der Pol, B.; Ferrero, D.V.; Buck-Barrington, L.; Hook, E.W.; Lenderman, C.; Quinn, T.C.; Gaydos, C.A.; Lovchik, J.C.; Schachter, J.; Moncada, J.; et al. Multicenter Evaluation of the BDProbeTec ET System for Detection of Chlamydia trachomatis and Neisseria gonorrhoeae in Urine Specimens, Female Endocervical Swabs, and Male Urethral Swabs. J. Clin. Microbiol. 2001, 39, 1008–1016. [CrossRef]
- Barton, K. MuMIn: Multi-Model Inference. 2009. Available online: http://r-forge.r-project.org/projects/mumin/ (accessed on 30 January 2009).
- 22. Burnham, K.P.; Anderson, D.R. Multimodel Inference: Understanding AIC and BIC in Model Selection. *Sociol. Methods Res.* 2004, 33, 261–304. [CrossRef]
- Mongo-Delis, A.; Mombo, L.E.; Mickala, P.; Bouassa, W.; Bouedy, W.S.; M'Batchi, B.; Bisseye, C. Factors associated with adherence to ARV treatment in people living with HIV/AIDS in a rural area (Koula-Moutou) in East Gabon. *Afr. J. AIDS Res.* 2019, 18, 51–57. [CrossRef]
- 24. Weldearegawi, T.Z.; Gerensea, H.; Berihu, H.; Gidey, G.; Welearegay, M.Z. The magnitude of opportunistic infections and associated factors in HIV-infected adults on antiretroviral therapy in southern zone Tigray, Ethiopia: A cross-sectional study. *Pan Afr. Med. J.* 2020, *35*, 126. [CrossRef]
- Girma, D.; Dejene, H.; Adugna Geleta, L.; Tesema, M.; Bati, F. Time to occurrence, predictors, and patterns of opportunistic infections incidence among HIV-positive patients attending Antiretroviral Therapy Clinic of Salale University Comprehensive Specialized Hospital: A retrospective cohort study. *Medicine* 2022, 101, e29905. [CrossRef] [PubMed]
- Charleine, M.B.; Bridy, M.D.; Coella, M.J.; Marion, N.O.M.; Magalie, E.M.; Michelle, N.O.G.; Philomene, K.N.; Marielle, B.A. Adverse effects related to HAART in Gabonese PLHIV: Frequency, severity and relationship with observance. *Health Sci. Dis.* 2023, 24, 16–23.
- 27. Fite, M.B.; Aga, D.J. Spectrum of Opportunistic Disease and Associated Factors among Patients Attending ART Clinic, Nekemte Specialized Hospital, Western Ethiopia. *bioRxiv* 2020. [CrossRef]

- 28. Mitiku, H.; Weldegebreal, F.; Teklemariam, Z. Magnitude of opportunistic infections and associated factors in HIV-infected adults on antiretroviral therapy in eastern Ethiopia. *HIV AIDS* **2015**, *7*, 137–144. [CrossRef] [PubMed]
- 29. Moges, N.A.; Kassa, G.M.; Boneya, D.J. Rate of HIV transmission and associated factors among HIV-exposed infants in selected health facilities of East and West Gojjam Zones, Northwest Ethiopia; retrospective cohort study. *BMC Infect. Dis.* 2017, 17, 475. [CrossRef] [PubMed]
- Damtie, D.; Yismaw, G.; Woldeyohannes, D.; Anagaw, B. Common opportunistic infections and their CD4 cell correlates among HIV-infected patients attending at antiretroviral therapy clinic of Gondar University Hospital, Northwest Ethiopia. BMC Res. Notes 2013, 6, 534. [CrossRef]
- Dereje, N.; Moges, K.; Nigatu, Y.; Holland, R. Prevalence and Predictors of Opportunistic Infections among HIV Positive Adults on Antiretroviral Therapy (On-ART) versus Pre-ART in Addis Ababa, Ethiopia: A Comparative Cross-Sectional Study. *HIV AIDS* 2019, 11, 229–237. [CrossRef]
- Sun, H.Y.; Chen, M.Y.; Hsieh, S.M.; Sheng, W.H.; Chang, S.Y.; Hsiao, C.F.; Hung, C.C.; Chang, S.C. Changes in the clinical spectrum of opportunistic illnesses in persons with HIV infection in Taiwan in the era of highly active antiretroviral therapy. *Jpn. J. Infect. Dis.* 2006, *59*, 311–316.
- Bélard, S.; Remppis, J.; Bootsma, S.; Janssen, S.; Kombila, D.U.; Beyeme, J.O.; Rossatanga, E.G.; Kokou, C.; Osbak, K.K.; Obiang Mba, R.M.; et al. Tuberculosis Treatment Outcome and Drug Resistance in Lambaréné, Gabon: A Prospective Cohort Study. Am. J. Trop. Med. Hyg. 2016, 95, 472–480. [CrossRef]
- 34. Nkoghe, D.; Toung Mve, M.; Nnegue, S.; Okome Nkoume, M.; Iba, B.J.; Hypolite, J.; Leonard, P.; Kendjo, E. HIV seroprevalence among tuberculosis patients in Nkembo Hospital, Libreville, Gabon. Short note. *Bull. Soc. Pathol. Exot.* **2005**, *98*, 121–122.
- 35. Kombila, D.U.; Moussavou-Kombila, J.B.; Grobusch, M.P.; Lell, B. Clinical and laboratory features of tuberculosis within a hospital population in Libreville, Gabon. *Infection* **2013**, *41*, 737–739. [CrossRef] [PubMed]
- Stolp, S.M.; Huson, M.A.; Janssen, S.; Beyeme, J.O.; Grobusch, M.P. Tuberculosis patients hospitalized in the Albert Schweitzer Hospital, Lambaréné, Gabon-a retrospective observational study. *Clin. Microbiol. Infect.* 2013, 19, E499–E501. [CrossRef] [PubMed]
- 37. Corbett, E.L.; Watt, C.J.; Walker, N.; Maher, D.; Williams, B.G.; Raviglione, M.C.; Dye, C. The growing burden of tuberculosis: Global trends and interactions with the HIV epidemic. *Arch. Intern. Med.* **2003**, *163*, 1009–1021. [CrossRef] [PubMed]
- Abe, H.; Ushijima, Y.; Bikangui, R.; Ondo, G.N.; Pemba, C.M.; Zadeh, V.R.; Mpingabo, P.I.; Ueda, H.; Agnandji, S.T.; Lell, B.; et al. Genetic Diversity of Hepatitis B and C Viruses Revealed by Continuous Surveillance from 2015 to 2021 in Gabon, Central Africa. *Microorganisms* 2023, 11, 2046. [CrossRef] [PubMed]
- 39. Groc, S.; Abbate, J.L.; Le Gal, F.; Gerber, A.; Tuaillon, E.; Albert, J.L.; Nkoghé, D.; Leroy, E.M.; Roche, B.; Becquart, P. High prevalence and diversity of hepatitis B and hepatitis delta virus in Gabon. *J. Viral Hepat.* **2019**, *26*, 170–182. [CrossRef]
- 40. Poudel, K.C.; Poudel-Tandukar, K.; Nakahara, S.; Yasuoka, J.; Jimba, M. Knowing the consequences of unprotected sex with seroconcordant partner is associated with increased safer sex intentions among HIV-positive men in Kathmandu, Nepal. *J. Health Popul. Nutr.* **2011**, *29*, 191–199. [CrossRef]
- Devarbhavi, H.; Asrani, S.K.; Arab, J.P.; Nartey, Y.A.; Pose, E.; Kamath, P.S. Global burden of liver disease: 2023 update. J. Hepatol. 2023, 79, 516–537. [CrossRef]
- Hu, J.; Liu, K.; Luo, J. HIV-HBV and HIV-HCV Coinfection and Liver Cancer Development. *Cancer Treat. Res.* 2019, 177, 231–250. [CrossRef]
- 43. Rosenthal, E.; Roussillon, C.; Salmon-Céron, D.; Georget, A.; Hénard, S.; Huleux, T.; Gueit, I.; Mortier, E.; Costagliola, D.; Morlat, P.; et al. Liver-related deaths in HIV-infected patients between 1995 and 2010 in France: The Mortavic 2010 study in collaboration with the Agence Nationale de Recherche sur le SIDA (ANRS) EN 20 Mortalité 2010 survey. *HIV Med.* 2015, *16*, 230–239. [CrossRef]
- Goehringer, F.; Bonnet, F.; Salmon, D.; Cacoub, P.; Paye, A.; Chêne, G.; Morlat, P.; May, T. Causes of Death in HIV-Infected Individuals with Immunovirologic Success in a National Prospective Survey. *AIDS Res. Hum. Retroviruses* 2017, 33, 187–193. [CrossRef]
- 45. Ioannou, G.N.; Bryson, C.L.; Weiss, N.S.; Miller, R.; Scott, J.D.; Boyko, E.J. The prevalence of cirrhosis and hepatocellular carcinoma in patients with human immunodeficiency virus infection. *Hepatology* **2013**, *57*, 249–257. [CrossRef] [PubMed]
- 46. Srirangaraj, S.; Venkatesha, D. Opportunistic infections in relation to antiretroviral status among AIDS patients from south India. *Indian J. Med. Microbiol.* **2011**, *29*, 395–400. [CrossRef] [PubMed]
- Dembelu, M.; Woseneleh, T. Prevalence of and Factors Associated with Reoccurrence of Opportunistic Infections among Adult HIV/AIDS Patients Attending the ART Clinic at Public Health Facilities in Arba Minch Town, Southern Ethiopia. *HIV AIDS* 2021, 13, 867–876. [CrossRef] [PubMed]
- 48. Pellowski, J.A.; Kalichman, S.C.; White, D.; Amaral, C.M.; Hoyt, G.; Kalichman, M.O. Real-time medication adherence monitoring intervention: Test of concept in people living with HIV infection. *J. Assoc. Nurses AIDS Care* **2014**, *25*, 646–651. [CrossRef]
- Trepka, M.J.; Fennie, K.P.; Sheehan, D.M.; Lutfi, K.; Maddox, L.; Lieb, S. Late HIV diagnosis: Differences by rural/urban residence, Florida, 2007–2011. AIDS Patient Care STDS 2014, 28, 188–197. [CrossRef] [PubMed]
- 50. Baker, P. Men's health: An overlooked inequality. Br. J. Nurs. 2016, 25, 1054–1057. [CrossRef]
- 51. Saldanha, D.; Gupta, N.; Shenoy, S.; Saralaya, V. Prevalence of opportunistic infections in AIDS patients in Mangalore, Karnataka. *Trop. Dr.* **2008**, *38*, 172–173. [CrossRef]

- 52. Benson, C.A.; Kaplan, J.E.; Masur, H.; Pau, A.; Holmes, K.K. Treating opportunistic infections among HIV-infected adults and adolescents: Recommendations from CDC, the National Institutes of Health, and the HIV Medicine Association/Infectious Diseases Society of America. *MMWR Recomm. Rep.* **2004**, *53*, 1–112. [CrossRef]
- 53. Dang, L.V.P.; Nguyen, Q.H.; Ishizaki, A.; Larsson, M.; Vu, N.T.P.; Do Duy, C.; Olson, L.; Dinh, T.T. Prevalence of Opportunistic Infections and Associated Factors in HIV-Infected Men Who Have Sex with Men on Antiretroviral Therapy in Bach Mai Hospital, Hanoi, Vietnam: A Case-Control Study. *Am. J. Men's Health* **2020**, *14*, 1557988320926743. [CrossRef]

Disclaimer/Publisher's Note: The statements, opinions and data contained in all publications are solely those of the individual author(s) and contributor(s) and not of MDPI and/or the editor(s). MDPI and/or the editor(s) disclaim responsibility for any injury to people or property resulting from any ideas, methods, instructions or products referred to in the content.