



**Regional differences in AIDS and non-AIDS related mortality in HIV-positive individuals across Europe and Argentina
the EuroSIDA study**

Reekie, Joanne; Kowalska, Justyna Dominika; Karpov, Igor; Rockstroh, Jurgen; Karlsson, Anders; Rakhmanova, Aza; Horban, Andrzej; Kirk, Ole; Lundgren, Jens; Mocroft, Amanda

Published in:
P L o S One

DOI:
[10.1371/journal.pone.0041673](https://doi.org/10.1371/journal.pone.0041673)

Publication date:
2012

Document version
Publisher's PDF, also known as Version of record

Citation for published version (APA):
Reekie, J., Kowalska, J. D., Karpov, I., Rockstroh, J., Karlsson, A., Rakhmanova, A., ... Mocroft, A. (2012). Regional differences in AIDS and non-AIDS related mortality in HIV-positive individuals across Europe and Argentina: the EuroSIDA study. *P L o S One*, 7(7), e41673. <https://doi.org/10.1371/journal.pone.0041673>

Regional Differences in AIDS and Non-AIDS Related Mortality in HIV-Positive Individuals across Europe and Argentina: The EuroSIDA Study

Joanne Reekie^{1*}, Justyna D. Kowalska², Igor Karpov³, Jurgen Rockstroh⁴, Anders Karlsson⁵, Aza Rakhmanova⁶, Andrzej Horban⁷, Ole Kirk^{2,8}, Jens D. Lundgren^{2,8}, Amanda Mocroft¹, for EuroSIDA in EuroCoord

1 Medical School, University College London, London, United Kingdom, **2** Copenhagen HIV Programme, University of Copenhagen, Copenhagen, Denmark, **3** Belarus State Medical University, Minsk, Belarus, **4** University of Bonn, Bonn, Germany, **5** Venhaelsan-Sodersjukhuset, Stockholm, Sweden, **6** Medical Academy Botkin Hospital, St Petersburg, Russia, **7** Warsaw Medical University, Hospital of Infectious Diseases, Warsaw, Poland, **8** Department of Infectious Diseases, Rigshospitalet, Copenhagen, Denmark

Abstract

Background: Differences in access to care and treatment have been reported in Eastern Europe, a region with one of the fastest growing HIV epidemics, compared to the rest of Europe. This analysis aimed to establish whether there are regional differences in the mortality rate of HIV-positive individuals across Europe, and Argentina.

Methods: 13,310 individuals under follow-up were included in the analysis. Poisson regression investigated factors associated with the risk of death.

Findings: During 82,212 person years of follow-up (PYFU) 1,147 individuals died (mortality rate 14.0 per 1,000 PYFU (95% confidence interval [CI] 13.1–14.8). Significant differences between regions were seen in the rate of all-cause, AIDS and non-AIDS related mortality (global $p < 0.0001$ for all three endpoints). Compared to South Europe, after adjusting for baseline demographics, laboratory measurements and treatment, a higher rate of AIDS related mortality was observed in East Europe (IRR 2.90, 95%CI 1.97–4.28, $p < .0001$), and a higher rate of non-AIDS related mortality in North Europe (IRR 1.51, 95%CI 1.24–1.82, $p < .0001$). The differences observed in North Europe decreased over calendar-time, in 2009–2011, the higher rate of non-AIDS related mortality was no longer significantly different to South Europe (IRR 1.07, 95%CI 0.66–1.75, $p = 0.77$). However, in 2009–2011, there remained a higher rate of AIDS-related mortality (IRR 2.41, 95%CI 1.11–5.25, $p = 0.02$) in East Europe compared to South Europe in adjusted analysis.

Interpretations: There are significant differences in the rate of all-cause mortality among HIV-positive individuals across different regions of Europe and Argentina. Individuals in Eastern Europe had an increased risk of mortality from AIDS related causes and individuals in North Europe had the highest rate of non-AIDS related mortality. These findings are important for understanding and reviewing HIV treatment strategies and policies across the European region.

Citation: Reekie J, Kowalska JD, Karpov I, Rockstroh J, Karlsson A, et al. (2012) Regional Differences in AIDS and Non-AIDS Related Mortality in HIV-Positive Individuals across Europe and Argentina: The EuroSIDA Study. PLoS ONE 7(7): e41673. doi:10.1371/journal.pone.0041673

Editor: Patrick S. Sullivan, Rollins School of Public Health, Emory University, United States of America

Received: February 6, 2012; **Accepted:** June 25, 2012; **Published:** July 23, 2012

Copyright: © 2012 Reekie et al. This is an open-access article distributed under the terms of the Creative Commons Attribution License, which permits unrestricted use, distribution, and reproduction in any medium, provided the original author and source are credited.

Funding: This work was supported by the European Commission BIOMED 1 (CT94-1637), BIOMED 2 (CT97-2713), the 5th Framework (QLK2-2000-00773) and the 6th Framework (LSHP-CT-2006-018632), and the 7th Framework (FP7/2007-2013, EuroCoord n° 260694) programmes. The participation of centres from Switzerland was supported by The Swiss National Science Foundation (Grant 108787). The funders had no role in study design, data collection and analysis, decision to publish, or preparation of the manuscript.

Competing Interests: The authors have declared that no competing interests exist.

* E-mail: j.reekie@ucl.ac.uk

Introduction

Mortality in HIV-positive individuals continues to decrease due to improvements in combination antiretroviral therapy (cART) and its availability, although it remains higher than in the general population [1]. A high proportion of the mortality observed in HIV-positive individuals is now due to non-AIDS related causes [2,3]. However, the incidence of non-AIDS related mortality, from causes such as cardiovascular disease (CVD), liver disease, and cancer, has also declined [3]. Drug specific toxicity, lifestyle factors (smoking, drug and alcohol misuse), co-infection with other viruses

(hepatitis B and C, human papillomavirus) and residual immunodeficiency are all thought to contribute to the higher relative risk of non-AIDS related mortality observed in HIV-positive individuals compared to the general population [4].

Research investigating regional differences across Europe, particularly allowing a comparison with East Europe, is limited. In Western and Central Europe, widespread access to antiretroviral therapy has led to considerable drops in AIDS related mortality [5]. However, in Eastern Europe, a region with one of the fastest growing HIV epidemics worldwide [6], the uptake of cART remains low and previous studies have reported substantial

differences in access to care and treatment compared to the rest of Europe [7,8]. Shortly after the introduction of cART, the EuroSIDA study reported that the all-cause mortality rate in Western Central Europe was significantly lower than in North and South Europe [9]. Since then, the study has grown substantially, particularly in Eastern Europe where few cohort studies are established. Additionally, to allow for comparisons outside of Europe the study includes data from Argentina, where the uptake of cART is high, with more than 95% of those in need of cART (CD4 count <350 cells/mm³) receiving it [10]. The study provides a unique opportunity to establish whether, in the current treatment era, regional differences in the mortality rate of HIV-positive individuals remain.

Methods

Ethics Statement

Informed consent and ethics committee approval were obtained in each participating centre according to national guidelines. Full study details are available at www.cphiv.dk.

Inclusion Criteria

The EuroSIDA study is a prospective study of individuals with HIV-1 in more than 100 centres across Europe, Israel and Argentina. To be eligible for inclusion into EuroSIDA, individuals must be aged 16 or older and have a pre-booked outpatient appointment at the centre. Starting in May 1994, there have been eight different periods of enrolment in EuroSIDA (cohorts I–VIII). Currently, 16,597 individuals are enrolled in EuroSIDA. The study has expanded into Eastern Europe with half of the individuals originating from Eastern European countries in Cohort VIII. Data is collected and updated at 6-monthly intervals on standardised recruitment and follow-up forms. Full study details are available at www.cphiv.dk. The incidence of LTFU in EuroSIDA has previously been reported to be fairly consistent over time at $<5\%$ per 100 PYFU [11]. An extensive quality assurance program has been established that includes data checking at the coordinating office, as well as regular monitoring visits with source verification of all new major events, plus a random selection of individuals followed at the clinics.

Statistical Methods

Individuals were included from baseline, defined as either 1st January 2002 or enrolment into EuroSIDA, whichever occurred later, this was when EuroSIDA expanded its network in East Europe. Additionally, previous EuroSIDA studies have reported mortality rates in earlier time periods [3,12]. Follow-up was until death, or 6 months after their last recorded visit (to allow for delays in reporting death), whichever occurred first. The incidence of mortality was calculated per 1,000 person years of follow-up (PYFU) and stratified by region. Individuals were divided into 6 regions according to country of residence as follows:

- South Europe: Greece, Israel, Italy, Portugal, Spain.
- West Central Europe: Austria, Belgium, France, Germany, Luxembourg, Switzerland.
- North Europe: Denmark, Finland, Ireland, the Netherlands, Norway, Sweden, the United Kingdom
- East Central Europe: Bulgaria, Croatia, the Czech Republic, Hungary, Poland, Romania, Serbia, Slovakia
- Eastern Europe: Belarus, Estonia, Latvia, Lithuania, the Russian Federation, Ukraine
- Argentina

In order to compare the regions of Europe and Argentina, South Europe was chosen as the reference group as this group contained the largest number of individuals.

For individuals who died, date and cause of death are reported by the site investigator and, since 2004, a Coding Causes of Death in HIV (CoDe) case report form is additionally completed for each fatal case and CoDe methods used to determine the underlying cause of death [13]. Deaths reported without a known cause were classified as either unknown AIDS or unknown non-AIDS mortality according to earlier developed methods [14].

Poisson regression analyses were used to investigate factors associated with the incidence of mortality. Variables investigated were age, gender, ethnic origin, smoking status, anaemia, hypertension, diabetes, hepatitis B and C status, HIV transmission group, CD4 count, any previous AIDS defining illnesses and year of follow-up. Age and year of follow-up were treated as continuous variables. CD4 count was categorised (<200 , 200–349, 350–500, ≥ 500 cell/mm³, missing) to allow for patients with missing values to remain in the analysis. Further treatment for HIV was split into 4 categories, not on cART, on cART with a viral load <500 copies/ml, on cART with a viral load ≥ 500 copies/ml, or on cART with no viral load measurement available. cART was defined as receiving ≥ 3 antiretrovirals. A cut-off of 500 copies/ml was chosen as this was lower limit of detection of the viral load assay for some individuals. Both current (time-updated) and baseline values were investigated for variables that could change over time. Factors that were significant in univariate analysis ($p < 0.1$) were included in multivariate analyses.

Individuals were classed as Hepatitis B positive if they had a positive HBV surface antigen test recorded and Hepatitis C positive if they had a positive HCV antibody test. Hypertension was defined as a diastolic blood pressure ≥ 90 or a systolic blood pressure ≥ 140 (mmHg) or receiving anti-hypertensive medication. If a diagnosis of insulin dependent diabetes was reported by the investigating centre or if an individual was receiving anti-diabetic medication they were classed as diabetic. Anaemia was defined as a haemoglobin level ≤ 12 or ≤ 14 (mg/dl) for females and males respectively [15]. Smoking status was defined as either never, current, previous or unknown.

All analysis was performed using SAS 9.1 (SAS institute, Cary North Carolina, USA).

Results

13,310 HIV-positive individuals were included in the analysis. There were clear differences in demographics across the regions (table 1). In East Europe, individuals were more likely to be younger (median age 30), female (42%) and infected via intravenous drug use [IDU] (49%) or heterosexual sex (41%). In contrast the majority of the individuals in West Central (45%), and North Europe (59%) were infected through homosexual sex. Similar baseline CD4 counts were observed across the regions. In South, West Central, North, East Central Europe and Argentina around 75–80% were on cART at baseline compared to only 30% in East Europe. In these individuals on cART, 62% in East Europe had a viral load <500 copies/ml compared to 70–85% in the other regions.

From 2002 until a median last follow-up date of September 2011 (interquartile range [IQR] April 09–December 2011), individuals were followed for a total of 82,212 PYFU. During this time 1,147 patients died, the crude mortality rate was 14.0 per 1,000 PYFU (95% confidence interval [CI] 13.1–14.8). The majority (837, 73%) were attributed to non-AIDS related causes, the crude mortality rate was 10.2 non-AIDS related deaths per

Table 1. Baseline characteristics (baseline was defined as either 1st January 2002 or enrolment into EuroSIDA which ever occurred later).

		South Europe		West central Europe		North Europe		East Central Europe		East Europe		Argentina	
Total (N,% of total)		3246	24.4%	2920	21.9%	2866	21.5%	1700	12.8%	2082	15.6%	496	3.7%
Gender (N.%)	Male	2391	73.7%	2246	76.9%	2332	81.4%	1221	71.8%	1171	57.7%	308	62.1%
Race (N.%)	White	2971	91.5%	2102	72.0%	2426	84.6%	1674	98.5%	2074	102.3%	489	98.6%
Exposure Group (N.%)	Homosexual	1101	33.9%	1307	44.8%	1693	59.1%	627	36.9%	154	7.6%	126	25.4%
	Injection Drug User	949	29.2%	406	13.9%	317	11.1%	449	26.4%	996	49.1%	62	12.5%
	Heterosexual	988	30.4%	849	29.1%	684	23.9%	490	28.8%	830	40.9%	292	58.9%
Prior AIDS (N.%)		853	26.3%	951	32.6%	915	31.9%	424	24.9%	402	19.8%	161	32.5%
Hepatitis B (N.%)	Negative	2186	67.3%	2308	79.0%	2339	81.6%	1408	82.8%	1653	81.5%	357	72.0%
	Positive	162	5.0%	202	6.9%	206	7.2%	81	4.8%	112	5.5%	26	5.2%
	Unknown	898	27.7%	410	14.0%	321	11.2%	211	12.4%	317	15.6%	113	22.8%
Hepatitis C (N.%)	Negative	1595	49.1%	1982	67.9%	1839	64.2%	995	58.5%	699	34.5%	295	59.5%
	Positive	755	23.3%	460	15.8%	329	11.5%	480	28.2%	1014	50.0%	91	18.3%
	Unknown	896	27.6%	478	16.4%	698	24.4%	225	13.2%	369	18.2%	110	22.2%
Hypertension	No	1629	50.2%	1136	38.9%	862	30.1%	1045	61.5%	1295	63.9%	292	58.9%
	Yes	488	15.0%	538	18.4%	505	17.6%	327	19.2%	119	5.9%	42	8.5%
	Unknown	1129	34.8%	1246	42.7%	1499	52.3%	328	19.3%	668	32.9%	162	32.7%
Diabetes	No	2904	89.5%	2012	68.9%	2614	91.2%	1609	94.6%	2061	101.6%	489	98.6%
	Yes	180	5.5%	148	5.1%	93	3.2%	44	2.6%	5	0.2%	5	1.0%
	Unknown	162	5.0%	760	26.0%	159	5.5%	47	2.8%	16	0.8%	2	0.4%
Anaemia	No	2194	67.6%	1904	65.2%	1642	57.3%	776	45.6%	197	9.7%	187	37.7%
	Yes	782	24.1%	747	25.6%	911	31.8%	295	17.4%	110	5.4%	205	41.3%
	Unknown	270	8.3%	269	9.2%	313	10.9%	629	37.0%	1775	87.5%	104	21.0%
Smoking	Never	706	21.7%	765	26.2%	576	20.1%	416	24.5%	387	19.1%	146	29.4%
	Current	1495	46.1%	1179	40.4%	1156	40.3%	872	51.3%	1142	56.3%	169	34.1%
	Previous	623	19.2%	669	22.9%	643	22.4%	306	18.0%	369	18.2%	146	29.4%
	Unknown	422	13.0%	307	10.5%	491	17.1%	106	6.2%	184	9.1%	35	7.1%
On cART (N.%)		2558	78.8%	2170	74.3%	2286	79.8%	1269	74.6%	603	29.7%	367	74.0%
On cART with viral load<500copies/ml (N, % on cART)		1795	70.2%	1732	79.3%	1927	84.3%	978	77.0%	371	61.5%	253	68.9%
Missing viral load on cART (N, % on cART)		16	0.6%	5	0.2%	11	0.5%	20	1.6%	40	6.6%	9	2.5%
Age (median, IQR)		40	35–46	42	37–49	43	37–50	35	30–42	30	25–36	37	31–43
CD4 count (median, IQR)		450	291–650	432	286–621	420	279–598	373	242–543	407	259–575	342	201–510
Missing CD4 count (N, %)		20	0.6%	10	0.3%	15	0.5%	6	0.4%	96	4.6%	4	0.8%
Baseline date (median, IQR)		1/02	1/02–12/03	1/02	1/02–12/05	1/02	1/02–12/03	1/04	1/02–12/08	2/06	4/04–6/08	2/06	11/03–6/06

doi:10.1371/journal.pone.0041673.t001

1,000 PYFU (95% CI 9.5–10.9) compared to 3.8 AIDS related deaths per 1,000 PYFU (95% CI 3.4–4.2). Figure 1 shows the crude mortality rate for all cause, AIDS and non-AIDS related causes by region. Significant differences in mortality were seen across the 6 different regions (global $p < 0.0001$ for all three endpoints). Higher all-cause mortality rates were observed in North and East Europe compared to the other regions. In East Europe a higher rate of AIDS related mortality was observed, the crude mortality rate due to AIDS related causes in East Europe was over three times higher than in South Europe (incidence rate ratio [IRR] 3.41, 95%CI 2.48–4.68, $p < 0.0001$). In contrast, a higher rate of non-AIDS related mortality was observed in North

Europe, compared to South Europe, the crude rate of non-AIDS related mortality in North Europe was 1.4 times higher (IRR 1.41, 95% CI 1.18–1.69, $p < 0.0001$).

Table 2 shows the characteristics of the individuals who died at the time of their death in each of the 6 regions. Those in East Europe were younger at the time of their death and had lower CD4 counts. A higher proportion on individuals in East Europe (66%) and Argentina (68%) had experienced an AIDS event prior to their death than in the other regions. In contrast, in those dying in South, West Central, North, East Central Europe and Argentina around 30–40% had been diagnosed with a non-AIDS

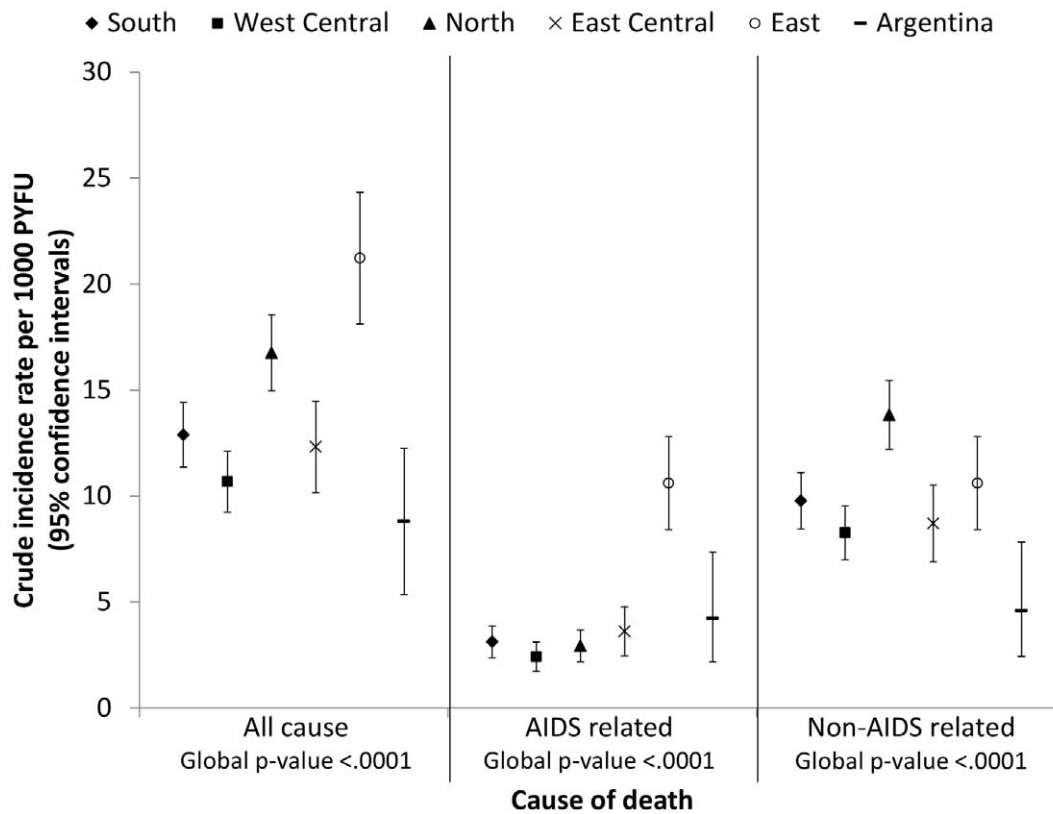


Figure 1. Crude mortality rate across the six different regions. Cause of death are reported by the site investigator and, since 2004, a Coding of Death in HIV (CoDe) case report form is additionally completed for each fatal case and CoDe methods used to determine the underlying cause of death.

doi:10.1371/journal.pone.0041673.g001

event prior to their death, whereas only 17% had been diagnosed in East Europe.

Table 3 gives the results of the Poisson regression analysis and presents the incidence rate ratios for all-cause, AIDS and non-AIDS related mortality for the five regions compared to South Europe. Adjusting for gender, age, HIV exposure group, AIDS diagnosis prior to baseline, non-AIDS event prior to baseline, hepatitis B and C status, hypertension, anaemia, baseline CD4 count, baseline date and baseline on cART/viral load status accounted for some of the higher rate of AIDS-related mortality observed in East Europe, though the rate of AIDS-related mortality was still more than double the rate in South Europe (IRR 2.27, 95%CI 1.42–3.46, $p = 0.0006$). Further, modelling age, hepatitis B and C status, hypertension, anaemia, CD4 count, year of follow-up and on cART viral load as time updated covariates the rate of AIDS related mortality was more than three times higher in East Europe compared to South.

A higher incidence of non-AIDS related mortality was observed, after adjusting for baseline variables (listed in the footnote of table 3), in North Europe (IRR 1.63, 95%CI 1.34–1.99, $p < 0.0001$), East Central Europe (IRR 1.31, 95%CI 1.01–1.70, $p = 0.04$) and East Europe (IRR 1.81, 95%CI 1.27–2.57, $p = 0.001$) compared to South Europe (Table 3). After including time-updated covariates in the model the increased rate was no longer significant in East Central Europe (IRR 1.22, 95% CI 0.95–1.58, $p = 0.11$) or East Europe (IRR 1.29, 95% CI 0.98–1.71, $p = 0.07$) but there remained a significantly increased rate of mortality in North Europe (IRR 1.51, 95% CI 1.24–1.82, $p < 0.0001$).

In the adjusted analyses, increasing year of follow-up was associated with a significantly lower rate of both AIDS (IRR 0.91, 95%CI 0.87–0.95, $p < 0.0001$) and non-AIDS related mortality (IRR 0.90, 95%CI 0.88–0.92, $p < 0.0001$). The test for interaction between region and calendar year of follow-up was significant for all-cause ($p = 0.02$) and AIDS related ($p = 0.05$) mortality, indicating that the effect of calendar year of follow-up on mortality rate may be different in each region, although it was non-significant for non-AIDS related mortality ($p = 0.10$). Figure 2 shows the adjusted incidence rate ratio for mortality per calendar year of follow-up stratified by region, from the fully adjusted model. Significant decreases over time in all-cause mortality rates were observed in West Central (IRR 0.90 per calendar year later, 95%CI 0.85–0.95, $p = 0.0002$), North (IRR 0.85 per calendar year later, 95%CI 0.82–0.89, $p < 0.0001$), and East Europe (IRR 0.89 per calendar year later, 95%CI 0.84–0.95, $p = 0.0005$). No significant changes over time in all-cause mortality were observed in South ($p = 0.32$) or East Central Europe ($p = 0.37$). Argentina was excluded from this analysis due to the limited number of patients. Despite the observed decrease in mortality in East Europe over time, in the most recent period, 2009–2011, there remained a higher rate of all-cause (IRR 1.50, 95%CI 1.50–2.38, $p = 0.08$) and AIDS-related mortality (IRR 2.41, 95%CI 1.11–5.25, $p = 0.02$) compared to South Europe in adjusted analysis. However, in 2009–2011, the significantly higher rate in North Europe was no longer observed for all-cause (IRR 0.91, 95%CI 0.58–1.42, $p = 0.18$) or non-AIDS related mortality (IRR 1.07, 95%CI 0.66–1.75, $p = 0.77$), suggesting that the differences in North Europe, relative to South, are decreasing over time.

Table 2. Characteristics at time of death of 1147 patients who died between 2002–2011 across Europe and Argentina.

	Region						P-value*
	South	West Central	North	East Central	East	Argentina	
N died (% of total in each region)	273 (8.4%)	208 (7.1%)	337 (11.8%)	126 (7.4%)	178 (8.6%)	25 (5.0%)	<.0001
Age (median, IQR)	47 (42–57)	48 (43–59)	50 (43–59)	42 (36–54)	35 (30–41)	41 (34–48)	<.0001
CD4 count, cells/mm ³ (median, IQR)	208 (78–392)	249 (85–450)	260 (113–452)	236 (90–416)	156 (45–296)	58 (24–136)	0.0002
Missing CD4 count (N, %)	56 (20.5%)	39 (18.8%)	78 (23.2%)	23 (18.3%)	82 (46.1%)	11 (44.0%)	
Ever started cART (N, %)	261 (95.6%)	195 (93.8%)	325 (96.4%)	112 (89.9%)	89 (50.0%)	24 (96.0%)	<.0001
HIV viral load <500 copies/ml (N,% with viral load measured)	128 (63.1%)	101 (61.6%)	187 (71.4%)	48 (58.5%)	17 (37.0%)	6 (60.0%)	0.0005
Missing viral load (N,%)	70 (25.6%)	44(21.2%)	75 (22.3%)	44 (34.9%)	132 (74.2%)	15 (60.0%)	
AIDS reported prior to death (N, %)	159 (58.2%)	123 (59.1%)	168 (49.9%)	75 (59.5%)	118 (66.3%)	17 (68.0%)	<.0001
Most recent AIDS diagnosis (N, % of AIDS diagnosis)							
Oesophageal candidiasis	12 (7.6%)	23 (18.7%)	32 (19.1%)	7 (9.3%)	26 (22.0%)	2 (11.8%)	
PCP	14 (8.8%)	9 (7.3%)	25 (14.9%)	7 (9.3%)	8 (6.8%)	2 (11.8%)	
HIV wasting syndrome	8 (5.0%)	12 (9.8%)	10 (6.0%)	6 (8.0%)	18 (15.3%)	1 (5.9%)	
Pulmonary TB	18 (11.3%)	5 (4.1%)	6 (3.6%)	7 (9.3%)	21 (17.8%)	1 (5.9%)	
NHL	24 (15.1%)	10 (8.1%)	20 (11.9%)	8 (10.7%)	2 (1.7%)	1 (5.9%)	
Non-AIDS event prior to death (N,%)	109 (39.9%)	63 (30.3%)	107 (31.8%)	45 (35.7%)	31 (17.4%)	8 (32.0%)	<.0001
Non-AIDS events reported (N,%)							
NADM	49 (17.9%)	24 (11.5%)	45 (13.3%)	18 (14.3%)	4 (2.2%)	4 (16%)	
Cardiovascular disease	32 (11.7%)	30 (14.4%)	52 (15.4%)	19 (15.1%)	3 (1.7%)	1 (4%)	
Pancreatitis	10 (3.7%)	7 (3.4%)	7 (2.1%)	4 93.2%)	10 (5.6%)	0 (0%)	
Liver related event	25 (9.2%)	9 (4.3%)	11 (3.3%)	10 (7.9%)	17 (9.6%)	4 (16%)	
End-stage renal disease	10 (3.7%)	7 (3.4%)	5 (1.5%)	0 (0%)	1 (0.6%)	0(0%)	

*Chi-squared test for categorical variables and Kruskal-Wallis for continuous variables. IQR: Inter-quartile range, PCP: Pneumocystis jiroveci pneumonia, TB: mycobacterium tuberculosis, NHL: non-Hodgkin's lymphoma, NADM: non-AIDS defining malignancy.
doi:10.1371/journal.pone.0041673.t002

Discussion

EuroSIDA is unique amongst studies of HIV-positive individuals in terms of its coverage across the entire continent of Europe. The results of this study with over 67,000 PYFU found significant differences in mortality rates across regions of Europe and Argentina. In particular, individuals in East Europe were found to have a higher mortality rate from AIDS related causes and individuals in North Europe a higher rate of non-AIDS related mortality. Similar rates of both AIDS and non-AIDS related mortality were seen across the other regions of Europe and Argentina. The mortality rate decreased over time particularly in West Central, North and East Europe.

Differences in the incidence of AIDS related mortality could partly be explained by variations in patient demographics including CD4 count and use of effective cART regimens. The epidemic in Eastern Europe is more recent, from 1995 there has been a rapid increase in HIV infections [16]. Since then a number of countries in the region have expanded access to antiretroviral coverage although treatment coverage remains low. UNAIDS estimated that in 2009 19% of those eligible for treatment in East Europe were accessing it, an estimated increase of 34% since the previous year [16]. In our study, after adjustment, the rate of AIDS related mortality appeared to be decreasing over time in East Europe, which may reflect improvements in antiretroviral coverage. However, after adjustment for time-updated CD4 count, use of cART, viral load and other factors, regional differences in

AIDS related mortality were still significant, indicating that the differences cannot be fully explained by those variables collected and included in our models.

A high proportion of patients in East Europe are infected via IDU [17]. This was accounted for in the adjusted analysis but, IDU populations may differ across the regions. Several factors have been identified as contributing to the excess risk of mortality observed in IDUs in the cART era including decreased access and adherence to cART [18,19], and more comorbidities such as co-infection with hepatitis C [20,21]. In West Central Europe coverage of HIV services in IDU populations has been reported to be high, particularly access to cART [22]. However in East Europe services are low, there are low levels of needle exchange programmes [23] and very few IDUs have been found to be receiving antiretroviral treatment [22]. Opiate substitution programmes are available in most countries but may be very limited in East Europe. Detailed information on opiate substitution programmes or whether individuals were actively injecting drug users was not available in this analysis. However, a sensitivity analysis, excluding those where the reported transmission route was IDU, found a similar increased incidence rate for East compared to South Europe, indicating that the higher prevalence of IDU does not account for the excess risk.

State-of-the-art care of HIV patients requires the utilisation of multiple health care interventions, such as the procurement of antiretroviral drugs, laboratory equipment, proper monitoring of antiretroviral treatment effectiveness and safety, and health care

Table 3. Results from univariate and multivariate Poisson regression analysis investigating mortality rates by region (South is the reference region).

All-cause mortality									
Region	Univariate			Multivariate^a			Multivariate^b		
	IRR	95% CI	p-value	IRR	95% CI	p-value	IRR	95% CI	p-value
South Europe	1.00	–	–	1.00	–	–	1.00	–	–
West Central Europe	0.83	0.69–0.99	0.04	0.99	0.82–1.20	0.90	0.85	0.70–1.03	0.09
North Europe	1.30	1.11–1.53	0.001	1.50	1.26–1.78	<.0001	1.37	1.15–1.62	0.0003
East Central Europe	0.96	0.77–1.18	0.67	1.31	1.05–1.64	0.01	1.28	1.03–1.59	0.02
East Europe	1.65	1.36–1.99	<.0001	2.07	1.57–2.73	<.0001	1.77	1.42–2.20	<.0001
Argentina	0.68	0.45–1.03	0.06	0.99	0.65–1.52	0.97	0.77	0.50–1.16	0.21
AIDS related mortality									
Region	Univariate			Multivariate^c			Multivariate^d		
	IRR	95% CI	p-value	IRR	95% CI	p-value	IRR	95% CI	p-value
South Europe	1.00	–	–	1.00	–	–	1.00	–	–
West Central Europe	0.77	0.53–1.13	0.18	0.84	0.57–1.23	0.36	0.72	0.49–1.06	0.09
North Europe	0.94	0.66–1.34	0.73	1.09	0.75–1.58	0.66	1.01	0.70–1.46	0.95
East Central Europe	1.16	0.78–1.74	0.46	1.29	0.84–1.97	0.24	1.43	0.93–2.19	0.10
East Europe	3.41	2.48–4.68	<.0001	2.27	1.42–3.64	0.0006	2.90	1.97–4.28	<.0001
Argentina	1.36	0.73–2.51	0.33	1.03	0.54–1.95	0.93	1.01	0.53–1.90	0.98
Non-AIDS related mortality									
Region	Univariate			Multivariate^a			Multivariate^b		
	IRR	95% CI	p-value	IRR	95% CI	p-value	IRR	95% CI	p-value
South Europe	1.00	–	–	1.00	–	–	1.00	–	–
West Central Europe	0.85	0.69–1.04	0.11	1.01	0.81–1.26	0.92	0.88	0.70–1.10	0.27
North Europe	1.41	1.18–1.69	0.0002	1.63	1.34–1.99	<.0001	1.51	1.24–1.82	<.0001
East Central Europe	0.89	0.69–1.14	0.35	1.31	1.01–1.70	0.04	1.22	0.95–1.58	0.11
East Europe	1.09	0.85–1.39	0.51	1.81	1.27–2.57	0.001	1.29	0.98–1.71	0.07
Argentina	0.47	0.27–0.82	0.008	0.87	0.49–1.54	0.62	0.61	0.35–1.09	0.09

^aadjusted for baseline variables gender, age, HIV exposure group, hepatitis B and C status, prior AIDS diagnosis, prior non-AIDS diagnosis, hypertension, diabetes, anaemia, smoking status, CD4 count, baseline date and on cART viral load.

^badjusted as in a but age, hepatitis B and C status, hypertension, diabetes, anaemia, smoking status, CD4 count, year of follow-up and on cART viral load were included as time-updated covariates.

^cadjusted for baseline variables gender, age, HIV exposure group, hepatitis B and C status, prior AIDS diagnosis, prior non-AIDS diagnosis, hypertension, diabetes, anaemia, CD4 count, baseline date and on cART viral load.

^dadjusted as in a but age, hepatitis B and C status, hypertension, diabetes, anaemia, CD4 count, year of follow-up and on cART viral load were included as time-updated covariates.

doi:10.1371/journal.pone.0041673.t003

infrastructure in general [16]. Not all differences in patient care will have been captured by the use of effective cART, and overall lower levels of patient care in East Europe, may play a significant role in the higher rate of AIDS related mortality. Socio-economic and infrastructure aspects, for which we do not have data [24], may also play a role.

An increased rate of non-AIDS mortality was found in patients in North Europe compared to South Europe, and these differences could not be explained by adjusting for differences in patient demographics, treatment or other risk factors that were measured such as smoking status, co-infection and CVD risk factors, either at baseline or time-updated. However, the rate of non-AIDS related mortality does appear to be decreasing over time in North Europe with no difference observed in the most recent time period. The differences in non-AIDS mortality are probably not driven by poor patient care but by underlying differences in population morbidity

and mortality. Smoking and excessive alcohol are found at an increased level in the HIV-positive population compared to the general population [25,26] and could not be fully adjusted for in these analyses. Further, Mediterranean diet has been found to be associated with a lower risk of all-cause mortality, in particular due to CVD and cancer [27]. A recent study found that middle-aged men in North Europe had a 5% and woman 28% higher risk of all-cause mortality compared to South [28]. Although we have accounted for CVD factors such as diabetes and hypertension, these data were not available for all patients. Additionally, data on alcohol use was not available for this analysis although it has recently been added to the EuroSIDA follow-up forms. Furthermore, if HIV infection is accelerating the aging process [29], certain factors in North Europe such as a high intake of animal fat, smoking, and alcohol use, may be acting with this and could partly explain the differences observed, although we found no evidence

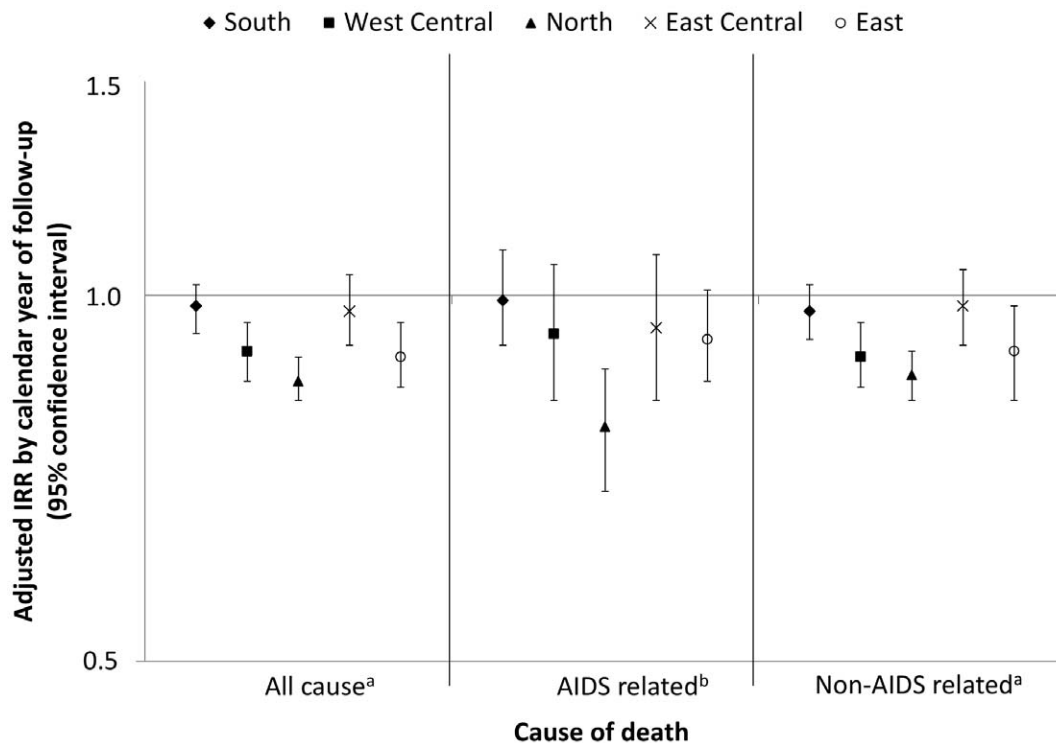


Figure 2. Incidence rate ratio (IRR) for all cause, AIDS and non-AIDS related mortality per calendar year of follow-up stratified by region. (Argentina was excluded from this analysis due to limited power). ^aadjusted for gender, age*, HIV exposure group, hepatitis B* and C* status, prior AIDS diagnosis, prior non-AIDS diagnosis, hypertension*, diabetes*, anaemia*, smoking status*, CD4 count*, baseline date* and on cART viral load*. ^badjusted for gender, age*, HIV exposure group, hepatitis B* and C* status, prior AIDS diagnosis, prior non-AIDS diagnosis, hypertension*, diabetes*, anaemia*, CD4 count*, baseline date* and on cART viral load*. *time-updated. doi:10.1371/journal.pone.0041673.g002

that the relationship between age and mortality differed between regions (data not shown).

Another possibility is that the higher observed rate in North Europe is due to under ascertainment or incomplete reporting of mortality in other regions. Centres in some countries may be linked to national death registers allowing more complete reporting of mortality or have better methods of finding patients lost to follow-up (LTFU). LTFU in EuroSIDA is low, with individuals in North and West Central Europe having the lowest and those in East Europe the highest rates of LTFU in the study [11]. To investigate biases caused by under ascertainment of death we varied the right censoring interval from 6 months after last follow-up visit to 3 months and 1 year. The results remained consistent with the main analysis, suggesting that these differences aren't solely due to underreporting in other regions. An analysis comparing the general HIV-negative population in these regions may also help clarify the reasons for these differences.

Demographically, individuals enrolled in the EuroSIDA study are fairly representative of each region based on comparisons with data from UNAIDS [30]. However, centres included in EuroSIDA are often university-associated clinics in larger cities, thus individuals in these clinics may have better access to care and may not be representative of all clinics in Europe, with potentially better clinical outcomes. This raises most concern in Eastern Europe, where the least favourable outcomes were seen, suggesting that improvements in this region are urgently needed. Further, within each of the regions some countries maybe quite heterogeneous in terms of their political and economic situation, however, the groupings provide a valuable regional overview of the differences in mortality rate across Europe.

In conclusion, differences were observed in the rate of all-cause mortality among HIV-positive individuals across different regions of Europe. Individuals in Eastern Europe had an increased risk of mortality from AIDS related causes in part due to differences in use of effective cART. Individuals in North Europe had the highest rate of non-AIDS related mortality which could not easily be explained but did appear to decrease over time. These findings are important for understanding and reviewing HIV treatment strategies and policies across the European region.

Acknowledgments

The multi-centre study group on EuroSIDA (national coordinators in parenthesis).

Argentina: (M Losso), C Elias, Hospital JM Ramos Mejia, Buenos Aires. **Austria:** (N Vetter), Pulmologisches Zentrum der Stadt Wien, Vienna; R Zangerle, Medical University Innsbruck, Innsbruck. **Belarus:** (I Karpov), A Vassilenko, Belarus State Medical University, Minsk, VM Mitsura, Gomel State Medical University, Gomel; O Suetnov, Regional AIDS Centre, Svetlogorsk. **Belgium:** (N Clumeck), S De Wit, M Delforge, Saint-Pierre Hospital, Brussels; R Colebunders, Institute of Tropical Medicine, Antwerp; L Vandekerckhove, University Ziekenhuis Gent, Gent. **Bosnia-Herzegovina:** (V Hadziosmanovic), Klinicki Centar Univerziteta Sarajevo, Sarajevo. **Bulgaria:** (K Kostov), Infectious Diseases Hospital, Sofia. **Croatia:** (J Begovac), University Hospital of Infectious Diseases, Zagreb. **Czech Republic:** (L Machala), D Jilich, Faculty Hospital Bulovka, Prague; D Sedlacek, Charles University Hospital, Plzen. **Denmark:** (J Nielsen), G Kronborg, T Benfield, M Larsen, Hvidovre Hospital, Copenhagen; J Gerstoft, T Katzenstein, A-B E Hansen, P Skinhoj, Rigshospitalet, Copenhagen; C Pedersen, Odense University Hospital, Odense; L Ostergaard, Skejby Hospital, Aarhus. **Estonia:** (K Zilmer), West-Tallinn Central Hospital, Tallinn; Jelena Smidt, Nakkusosakond Siseklinik, Kohtla-Järve. **Finland:** (M Ristola),

Helsinki University Central Hospital, Helsinki. **France:** (C Katlama), Hôpital de la Pitié-Salpêtrière, Paris; J-P Viard, Hôpital Necker-Enfants Malades, Paris; P-M Girard, Hospital Saint-Antoine, Paris; JM Livrozet, Hôpital Edouard Herriot, Lyon; P Vanhems, University Claude Bernard, Lyon; C Pradier, Hôpital de l'Archet, Nice; F Dabis, D Neau, Unité INSERM, Bordeaux. **Germany:** (J Rockstroh), Universitäts Klinik Bonn; R Schmidt, Medizinische Hochschule Hannover; J van Lunzen, O Degen, University Medical Center Hamburg-Eppendorf, Infectious Diseases Unit, Hamburg; HJ Stellbrink, IPM Study Center, Hamburg; S Staszewski, JW Goethe University Hospital, Frankfurt; J Bogner, Medizinische Poliklinik, Munich; G. Fätkenheuer, Universität Köln, Cologne. **Greece:** (J Kosmidis), P Gargalianos, G Xylomenos, J Perdios, Athens General Hospital; G Panos, A Filandras, E Karabatsaki, 1st IKA Hospital; H Sambatakou, Ippokraton General Hospital, Athens. **Hungary:** (D Banhegyi), Szent László Hospital, Budapest. **Ireland:** (F Mulcahy), St. James's Hospital, Dublin. **Israel:** (I Yust), D Turner, M Burke, Ichilov Hospital, Tel Aviv; S Pollack, G Hassoun, Rambam Medical Center, Haifa; S Maayan, Hadassah University Hospital, Jerusalem. **Italy:** (S Vella), Istituto Superiore di Sanità, Rome; R Esposito, I Mazeu, C Mussini, Università Modena, Modena; C Arici, Ospedale Riuniti, Bergamo; R Pristera, Ospedale Generale Regionale, Bolzano; F Mazzotta, A Gabbuti, Ospedale S Maria Annunziata, Firenze; V Vullo, M Lichtner, University di Roma la Sapienza, Rome; A Chirrianni, E Montesarchio, M Gargiulo, Presidio Ospedaliero AD Cotugno, Monaldi Hospital, Napoli; G Antonucci, A Testa, P Narciso, C Vlassi, M Zaccarelli, Istituto Nazionale Malattie Infettive Lazzaro Spallanzani, Rome; A Lazzarin, A Castagna, N Gianotti, Ospedale San Raffaele, Milan; M Galli, A Ridolfo, Osp. L. Sacco, Milan; A d'Arminio Monforte, Istituto Di Clinica Malattie Infettive e Tropicale, Milan. **Latvia:** (B Rozentale), I Zeltina, Infectology Centre of Latvia, Riga. **Lithuania:** (S Chaplinskas), Lithuanian AIDS Centre, Vilnius. **Luxembourg:** (R Hemmer), T Staub, Centre Hospitalier, Luxembourg. **Netherlands:** (P Reiss), Academisch Medisch Centrum bij de Universiteit van Amsterdam, Amsterdam. **Norway:** (V Ormaasen), A Maeland, J Bruun, Ullevål Hospital, Oslo. **Poland:** (B Knysz) J Gasiorowski, Medical University, Wroclaw; A Horban, E Bakowska, Centrum Diagnostyki i Terapii AIDS, Warsaw; A Grzeszczuk, R Flisiak, Medical University, Bialystok; A Boron-Kaczmarek, M Pynka, M Parczewski, Medical University, Szczecin; M Beniowski, E Mularska, Osrodek Diagnostyki i Terapii AIDS, Chorzow; H Trocha, Medical University, Gdansk; E Jablonowska, E Malolepsza, K Wojcik, Wojewodzki Szpital Specjalistyczny, Lodz. **Portugal:** (F Antunes), M Doroana, L Caldeira, Hospital Santa Maria, Lisbon; K Mansinho, Hospital de Egas Moniz, Lisbon; F Maltez, Hospital Curry Cabral, Lisbon. **Romania:** (D Duiculescu), Spitalul de Boli Infectioase si Tropicale: Dr. Victor Babes, Bucarest. **Russia:** (A Rakhmanova), Medical Academy Botkin Hospital, St Petersburg; N Zakharova, St Petersburg AIDS Centre, St Peterburg; S

Buzunova, Novgorod Centre for AIDS, Novgorod. **Serbia:** (D Jevtovic), The Institute for Infectious and Tropical Diseases, Belgrade. **Slovakia:** (M Mokráš), D Staneková, Dérer Hospital, Bratislava. Slovenia: (J Tomazic), University Clinical Centre Ljubljana, Ljubljana. **Spain:** (J González-Lahoz), V Soriano, P Labarga, J Medrano, Hospital Carlos III, Madrid; S Moreno, J. M. Rodriguez, Hospital Ramon y Cajal, Madrid; B Clotet, A Jou, R Paredes, C Tural, J Puig, I Bravo, Hospital Germans Trias i Pujol, Badalona; JM Gatell, JM Miró, Hospital Clinic i Provincial, Barcelona; P Domingo, M Gutierrez, G Mateo, MA Sambeat, Hospital Sant Pau, Barcelona. **Sweden:** (A Karlsson), Venhaelsan-Sodersjukhuset, Stockholm; L Flamholc, Malmö University Hospital, Malmö. **Switzerland:** (B Ledergerber), R Weber, University Hospital, Zürich; P Francioli, M Cavassini, Centre Hospitalier Universitaire Vaudois, Lausanne; B Hirschel, E Boffi, Hospital Cantonal Universitaire de Geneve, Geneve; H Furrer, Inselspital Bern, Bern; M Battegay, L Elzi, University Hospital Basel. **Ukraine:** (E Kravchenko), N Chentsova, Kiev Centre for AIDS, Kiev; V Frolov, G Kutsyna, Luhansk State Medical University; Luhansk; S Servitskiy, Odessa Region AIDS Center, Odessa; M Krasnov, Kharkov State Medical University, Kharkov. **United Kingdom:** (S Barton), St. Stephen's Clinic, Chelsea and Westminster Hospital, London; AM Johnson, D Mercey, Royal Free and University College London Medical School, London (University College Campus); A Phillips, MA Johnson, A Mocroft, Royal Free and University College Medical School, London (Royal Free Campus); M Murphy, Medical College of Saint Bartholomew's Hospital, London; J Weber, G Scullard, Imperial College School of Medicine at St. Mary's, London; M Fisher, Royal Sussex County Hospital, Brighton; C Leen, Western General Hospital, Edinburgh.

Steering Committee: J Gatell, B Gazzard, A Horban, J Lundgren, I Karpov, B Ledergerber, M Lusso, A D'Arminio Monforte, C Pedersen, A Phillips, A Rakhmanova, M Ristola, P Reiss, J Rockstroh (Chair), S De Wit (Vice-Chair)

Coordinating Centre Staff: O Kirk, A Mocroft, A Cozzi-Lepri, D Grint, M Ellefson, D Podlekareva, J Kjør, L Peters, J Reekie, J Kowalska, J Nielsen, J Tverland, A H Fischer

EuroSIDA representatives to EuroCoord: O. Kirk, A. Mocroft, J. Grarup, S. deWitt, P. Reiss, A. Cozzi-Lepri, R. Thiebaut, J. Rockstroh, D. Burger, R. Paredes, J. Kjør

Author Contributions

Conceived and designed the experiments: JDL OK AM. Analyzed the data: JMR JDK. Wrote the paper: JMR. National data coordination: IK JR AK AR AH. Interpretation of results: JMR JDK JDL OK AM. Critically revised manuscript: JDK IK JR AK AR AH JDL OK AM. Approved the final version of the manuscript: JMR JDK IK JR AK AR AH JDL OK AM STUDY PI: JDL OK AM.

References

- Antiretroviral Therapy Cohort Collaboration (2008) Life expectancy of individuals on combination antiretroviral therapy in high-income countries: a collaborative analysis of 14 cohort studies. *Lancet* 372: 293–299.
- Lewden C, May T, Rosenthal E, Burty C, Bonnet F, et al. (2008) Changes in causes of death among adults infected by HIV between 2000 and 2005: The "Mortalite 2000 and 2005" surveys (ANRS EN19 and Mortavic). *J Acquir Immune Defic Syndr* 48: 590–598.
- Mocroft A, Brettle R, Kirk O, Blaxhult A, Parkin JM, et al. (2002) Changes in the cause of death among HIV positive subjects across Europe: results from the EuroSIDA study. *AIDS* 16: 1663–1671.
- Deeks SG, Phillips AN (2009) HIV infection, antiretroviral treatment, ageing, and non-AIDS related morbidity. *BMJ* 338: a3172.
- Joint United Nations Programme on HIV/AIDS (UNAIDS) (2009) Fact Sheet, North America and Western and Central Europe. Latest epidemiological trends.
- UNAIDS, World Health Organization (2009) 09 AIDS epidemic update. 1–100.
- Podlekareva D, Reekie J, Rakhmanova A, Horban A, Mocroft A, et al. (2008) Indicators of the use of healthcare interventions across Europe.
- Podlekareva D, Bannister W, Mocroft A, Abrosimova L, Karpov I, et al. (2008) The EuroSIDA study: Regional differences in the HIV-1 epidemic and treatment response to antiretroviral therapy among HIV-infected patients across Europe—a review of published results. *Cent Eur J Public Health* 16: 99–105.
- Chiesi A, Mocroft A, Dally LG, Miller V, Katlama C, et al. (1999) Regional survival differences across Europe in HIV-positive people: the EuroSIDA study. *AIDS* 13: 2281–2288.
- UNAIDS (2009) Argentina County Situation 2009.
- Mocroft A, Kirk O, Aldins P, Chiesi A, Blaxhult A, et al. (2008) Loss to follow-up in an international, multicentre observational study. *HIV Med* 9: 261–269.
- Mocroft A, Ledergerber B, Katlama C, Kirk O, Reiss P, et al. (2003) Decline in the AIDS and death rates in the EuroSIDA study: an observational study. *Lancet* 362: 22–29.
- Kowalska JD, Friis-Moller N, Kirk O, Bannister W, Mocroft A, et al. (2011) The Coding Causes of Death in HIV (CoDe) Project: initial results and evaluation of methodology. *Epidemiology* 22: 516–523.
- Kowalska JD, Mocroft A, Ledergerber B, Florence E, Ristola M, et al. (2011) A standardized algorithm for determining the underlying cause of death in HIV infection as AIDS or non-AIDS related: results from the EuroSIDA study. *HIV Clin Trials* 12: 109–117.
- Mocroft A, Kirk O, Barton SE, Dietrich M, Proenca R, et al. (1999) Anaemia is an independent predictive marker for clinical prognosis in HIV-infected patients from across Europe. EuroSIDA study group. *AIDS* 13: 943–950.
- UNAIDS (2010) Global Report:UNAIDS report on the global AIDS epidemic.
- UNAIDS (2009) Fact sheets. Eastern Europe and Central Asia. Latest epidemiological trends.
- van Asten LC, Boufassa F, Schiffer V, Brettle RP, Robertson JR, et al. (2003) Limited effect of highly active antiretroviral therapy among HIV-positive injecting drug users on the population level. *Eur J Public Health* 13: 347–349.
- Wood E, Montaner JS, Yip B, Tyndall MW, Schechter MT, et al. (2003) Adherence and plasma HIV RNA responses to highly active antiretroviral therapy among HIV-1 infected injection drug users. *CMAJ* 169: 656–661.
- Braitstein P, Justice A, Bangsberg DR, Yip B, Alfonso V, et al. (2006) Hepatitis C coinfection is independently associated with decreased adherence to

- antiretroviral therapy in a population-based HIV cohort. *AIDS* 20: 323–331. 10.1097/01.aids.0000198091.70325.f4 [doi];00002030-200602140-00003 [pii].
21. Murray M, Hogg R, Lima V, May M, Moore D, et al. (2011) The effect of injecting drug use history on disease progression and death among HIV-positive individuals initiating combination antiretroviral therapy: collaborative cohort analysis. *HIV Med.* 10.1111/j.1468-1293.2011.00940.x [doi].
 22. Mathers BM, Degenhardt L, Ali H, Wiessing L, Hickman M, et al. (2010) HIV prevention, treatment, and care services for people who inject drugs: a systematic review of global, regional, and national coverage. *Lancet* 375: 1014–1028. S0140-6736(10)60232-2 [pii];10.1016/S0140-6736(10)60232-2 [doi].
 23. Sarang A, Rhodes T, Platt L (2008) Access to syringes in three Russian cities: implications for syringe distribution and coverage. *Int J Drug Policy* 19 Suppl 1: S25–S36. S0955-3959(07)00244-7 [pii];10.1016/j.drugpo.2007.11.008 [doi].
 24. Hamers FF, Downs AM (2003) HIV in central and eastern Europe. *Lancet* 361: 1035–1044. S0140-6736(03)12831-0 [pii];10.1016/S0140-6736(03)12831-0 [doi].
 25. Friis-Moller N, Sabin CA, Weber R, d'Arminio MA, El-Sadr WM, et al. (2003) Combination antiretroviral therapy and the risk of myocardial infarction. *N Engl J Med* 349: 1993–2003.
 26. Lewden C, Salmon D, Morlat P, Bevilacqua S, Jouglu E, et al. (2005) Causes of death among human immunodeficiency virus (HIV)-infected adults in the era of potent antiretroviral therapy: emerging role of hepatitis and cancers, persistent role of AIDS. *Int J Epidemiol* 34: 121–130.
 27. Sofi F, Cesari F, Abbate R, Gensini GF, Casini A (2008) Adherence to Mediterranean diet and health status: meta-analysis. *BMJ* 337: a1344.
 28. Vollset SE (2010) Does the southern European cardiovascular mortality advantage extend to total mortality? 50-year trends in death risks between 40 and 70 years of age in Western European men and women. *Scand J Public Health* 38: 127–134.
 29. Deeks SG (2011) HIV infection, inflammation, immunosenescence, and aging. *Annu Rev Med* 62: 141–155.
 30. UNAIDS (2008) 08 Report on the global AIDS epidemic.