

Delayed HIV diagnosis and initiation of antiretroviral therapy: inequalities by educational level, COHERE in EuroCoord

The Socio-economic Inequalities and HIV Writing Group for Collaboration of Observational HIV Epidemiological Research in Europe (COHERE) in EuroCoord*

Objectives: In Europe and elsewhere, health inequalities among HIV-positive individuals are of concern. We investigated late HIV diagnosis and late initiation of combination antiretroviral therapy (cART) by educational level, a proxy of socioeconomic position.

Design and methods: We used data from nine HIV cohorts within COHERE in Austria, France, Greece, Italy, Spain and Switzerland, collecting data on level of education in categories of the UNESCO/International Standard Classification of Education standard classification: non-completed basic, basic, secondary and tertiary education. We included individuals diagnosed with HIV between 1996 and 2011, aged at least 16 years, with known educational level and at least one CD4⁺ cell count within 6 months of HIV diagnosis. We examined trends by education level in presentation with advanced HIV disease (AHD) (CD4⁺ <200 cells/μl or AIDS within 6 months) using logistic regression, and distribution of CD4⁺ cell count at cART initiation overall and among presenters without AHD using median regression.

Results: Among 15 414 individuals, 52, 45, 37, and 31% with uncompleted basic, basic, secondary and tertiary education, respectively, presented with AHD (*P* trend <0.001). Compared to patients with tertiary education, adjusted odds ratios of AHD were 1.72 (95% confidence interval 1.48–2.00) for uncompleted basic, 1.39 (1.24–1.56) for basic and 1.20 (1.08–1.34) for secondary education (*P* < 0.001). In unadjusted and adjusted analyses, median CD4⁺ cell count at cART initiation was lower with poorer educational level.

Conclusions: Socioeconomic inequalities in delayed HIV diagnosis and initiation of cART are present in European countries with universal healthcare systems and individuals with lower educational level do not equally benefit from timely cART initiation.

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Introduction

Socioeconomic status is inversely associated with access to and use of health services by the general population and, in Europe and elsewhere, health inequalities in the general and in subpopulations such as HIV-positive people are a

growing concern [1–3]. Members of disadvantaged socioeconomic groups face barriers to health services [2,4], even in European countries with universal and public health insurance [5]. In spite of the existence of universal access to confidential HIV testing, HIV care and combination antiretroviral therapy (cART) in most

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European countries [6], a patient's socioeconomic status may influence the ease or difficulty of access. Since the HIV epidemic is entrenched among the socially vulnerable, including homosexual men, injecting drug users (IDUs) and migrants [7,8], with access to healthcare increasingly limited for undocumented migrants [9,10], questions about the effect of socio-economic status on the diagnosis and treatment of HIV-positive persons are pressing.

Timely diagnosis of HIV infection is the first step in the HIV treatment cascade, but the COHERE collaboration of HIV cohorts of routine clinical practice in Europe recently reported that there is a very high rate of late presentation [11]. Those who do not know they have an HIV infection are less likely to take steps to prevent its transmission. Late diagnoses delay cART initiation and raise the rates of hospitalization, morbidity and mortality [12,13]. Universal access does not alone ensure that health services will be utilized. Measuring socioeconomic status is difficult. In adults, educational level is widely used as a proxy given that it is fairly stable beyond early adulthood and is therefore less likely to be affected by reverse causation [14–17], that is, if the association between socioeconomic status exists, because better health status allows to achieve a better socioeconomic status. Previous studies on the association between educational level, the timing of HIV diagnosis and initiation of cART lack a pan-European perspective to present a clear picture of the problem [18–20]. These studies have been conducted in single countries, each applying a different classification of educational level. Without a clear understanding of the role that socioeconomic inequality plays in access to HIV diagnosis and treatment, we cannot formulate effective policy and programs, or efficiently allocate resources, particularly at times of economic crisis [21].

We investigated the association between socioeconomic status and delayed HIV diagnosis and initiation of cART in COHERE, a large European collaboration of HIV cohort studies. We chose educational level as a proxy for socioeconomic position and harmonized data on education across cohorts. Our working hypothesis was that lower educational level is linked to higher risk of delayed diagnosis and cART initiation. Because of the widespread practice of antenatal testing in the study setting and the different testing strategies for the different HIV transmission categories, our secondary aims were to examine whether the association varied by sex and transmission category.

Methods

Patients

COHERE in EuroCoord is a collaboration of 35 observational cohorts covering 32 European countries, within the framework of the EuroCoord network of excellence (<http://www.eurocoord.net>). Each cohort

submits data in a standardized format (the HIV Collaboration Data Exchange Protocol, <http://www.hicdep.org>) to co-ordinating centres at the Copenhagen HIV Programme, Denmark, or the Institut de Sante Publique d'Epidémiologie et de Développement (Bordeaux School of Public Health), Bordeaux, France. The Regional Coordinating Centres ensure adherence to strict quality assurance guidelines and perform data checks, including the removal of duplicate records. Data include information on patients' characteristics (age, sex, geographical origin, and transmission category), use of cART (type of regimes and dates of start and discontinuation), CD4⁺ cell counts and plasma HIV-RNA over time and their dates, AIDS-defining conditions and deaths. Further information is available at www.cohere.org.

In 2012, EuroCoord defined socioeconomic variables with the intent of standardizing and harmonizing collection of socioeconomic data across cohorts in European countries. Educational level was the only available socioeconomic variable whose harmonization across European countries could be performed retrospectively. The definition of the variable 'maximum attained level of education' was based on the UNESCO/International Standard Classification of Education (ISCED) standard classification, and was classified as 'uncompleted basic' (ISCED 0), 'basic' (ISCED 1 and 2), 'secondary' (ISCED 3 and 4) and 'tertiary' (ISCED 5 and 6), based on data on education systems and reforms available from the European Encyclopaedia on National Education Systems (http://eacea.ec.europa.eu/education/eurydice/eurydice_en.php).

These analyses included data from nine cohorts in six European countries (Austria, France, Greece, Italy, Spain and Switzerland) that collected data on maximum attained educational level. Patients were included if they were diagnosed with HIV between 1 January 1996 and 31 December 2011, were aged at least 16 years, were enrolled within 6 months of HIV diagnosis, and had at least one measured CD4⁺ cell count within 6 months of their HIV diagnosis while cART-naïve and the level of education was known [11]. Patients from COHERE's three seroconverter cohorts were excluded; by definition, they could not qualify as late presenters. Ethics approval was granted by the Ethic Committees of each of the participating cohorts according to country regulations. Signed informed consent was obtained from all patients.

Data were pooled in September 2011 within COHERE in EuroCoord (www.cohere.org and www.EuroCoord.net) and additional data on educational level data were received from the cohorts in 2012.

Statistical analyses

Delayed HIV presentation

We defined delayed HIV presentation based on definitions proposed by the European Late Presenter

Consensus Working group: presentation with advanced HIV disease (AHD) as CD4⁺ cell count below 200 cells/ μ l or an AIDS-defining event in the 6 months following presentation; and presentation with late HIV disease (LHD) as CD4⁺ cell count below 350 cells/ μ l or an AIDS-defining event in the 6 months following diagnosis [22].

We used logistic regression models to explore the association between educational level with AHD and LHD, adjusting for the following potential confounders, chosen *a priori*: calendar period of HIV diagnosis (<2001 versus \geq 2001, to reflect the introduction of optimal cART regimen, including boosted protease inhibitors and non-nucleoside reverse transcriptase inhibitors); transmission category (MSM, heterosexuals, IDUs, other/unknown); geographical origin (Europe, non-European, unknown); age at HIV diagnosis (<35 versus \geq 35 years, median age of the dataset); sex; and cohort. Likelihood ratio tests were used to determine if sex and transmission category were effect modifiers for the association between educational level and delayed HIV diagnosis. We also refit the models treating age at HIV diagnosis as a continuous variable (linear and fractional polynomial [23]), and, since the younger patients might not have completed education yet, restricting to individuals with age above 25 years. Finally, we described differences by broad cohort geographical areas defined *a priori* as Austria, France and Switzerland; and Greece, Italy and Spain. These analyses were descriptive and not adjusted for potential confounders. We present detailed results for the AHD analyses and a summary of LHD analyses.

Because the analyses excluded patients with unknown educational level and complete case analyses may be biased if data on educational level are not missing completely at random (MCAR), we used the following sensitivity analyses. First, we used multiple imputation to impute educational level [24] assuming data are missing at random. After inspecting the characteristics of the individuals with available and with missing data on level of education, we decided to include in the imputation model CD4⁺ cell count at HIV diagnosis in addition to all variables used in the analysis models. Whereas age and CD4⁺ cell count were treated as continuous variables, all other variables were treated as categorical. Twenty imputed data sets were generated, analysed separately and then combined using Rubin's rule [25]. Second, we imputed missing data on educational level under the assumption of data missing not at random and we assigned all patients with unknown educational level to basic education, secondary education and tertiary education in separate analyses. These extreme case scenarios are unrealistic in practice, but provide an illustration of how sensitive the analyses may be to assumptions regarding missing data. Since we excluded patients without CD4⁺ cell count measured within 6 months of HIV, we also ran analyses in which we assumed these patients were all AHD and none AHD. Finally, we

reanalysed the data including patients who were enrolled and had more than one CD4⁺ cell count within 3 months and 12 months of HIV diagnosed rather than 6 months of HIV diagnosis as in the main analysis.

CD4⁺ cell count at initiation of antiretroviral therapy

We explored patterns of delayed cART initiation among patients included in the analyses of delayed HIV diagnosis. We further restricted the analysis to individuals who initiated cART while cART-naïve and who had their CD4⁺ cell count measured at least once, between 6 months before and 1 week after cART was initiated. Tests for trend and multiple median regression models estimated the association between level of education and CD4⁺ cell count at cART initiation overall and when restricted to individuals who did not present with AHD. All models were adjusted for sex, calendar year of cART initiation, transmission category, geographical origin, age at cART initiation, and cohort. Since there is no formal definition of late cART initiation, and national and international recommendations changed significantly across the study period, we decided that analyses of the distribution of CD4⁺ cell count at cART initiation with median regression would return more interpretable and robust results. As a sensitivity analysis, we defined late cART initiation as individuals with a CD4⁺ cell count below 350 cells/ μ l prior to initiation, and used logistic regression models adjusting for the same covariates above.

Analyses were performed using Stata version 11.0 (Stata Corp., College Station, Texas, USA).

Results

Of the 37 438 patients diagnosed with HIV during 1996–2011, 22 024 were excluded (12 656 enrolled >6 months of HIV diagnosis; 2459 had no CD4⁺ cell count measurements while cART-naïve within 6 months of HIV diagnosis; 6909 had unknown level of education). Of the remaining 15 414 patients, 9, 28, 47 and 15% had uncompleted basic, basic, secondary and tertiary education. Compared to the excluded individuals, these were more likely to be of European geographical origin, infected through sex between men and diagnosed with HIV after 2000. Median [interquartile range (IQR)] CD4⁺ cell count and age at HIV diagnosis were 304 (125–503) cells/ μ l and 35 [29–43] years; 76% were men, and 83% acquired HIV through heterosexual or MSM contact. Proportions with uncompleted basic, basic, secondary and tertiary education were 17, 39, 39 and 6% for women and 7, 25, 50 and 18% for men.

Delayed diagnosis

A total of 6129 (40%) patients presented with AHD, of which 5725 had CD4⁺ below 200 cells/ μ l, 404 had an AIDS-defining event, and 2522 had both. The proportion of AHD decreased with educational level: 52, 45,

37, and 31% for uncompleted basic, basic, secondary and tertiary education (P for trend <0.001). Although the proportion of patients presenting with AHD did not differ for men and women (39 and 41%, respectively), the proportion of AHD for uncompleted basic and tertiary education varied more for men (54–30%) than for women (49–41%) ($P < 0.001$ for interaction test). The gradient was more pronounced after 2001 ($P < 0.001$ test for interaction); the proportion of AHD with uncompleted basic and tertiary education was 55 and 43% before 2001 and 50 and 28% from 2001 onwards (Table 1). The gradient by educational level was more pronounced for MSM compared to other groups; the proportion of AHD ranged between 45 and 26% for patients with uncompleted basic and tertiary education (Table 1). Gradients by educational level varied by cohort, but were consistently observed when data were grouped by broad geographical regions, although more noticeable in Greece, Italy and Spain (Table 1).

The gradient by educational level was maintained in multivariate analyses; compared to tertiary education, the adjusted odds ratios (aORs) for AHD were 1.72 [95% confidence interval (CI) 1.48–2.00], 1.39 (95% CI 1.24–1.56) and 1.20 (95% CI 1.08–1.34) in patients with uncompleted basic, basic and secondary completed education. In the adjusted logistic model, male sex, calendar period of diagnosis 1996–2000, non-European geographical origin, membership in the heterosexual and injecting drug use transmission groups, and age above 35

years at HIV diagnosis were significant predictors of presenting with AHD (data not shown).

Figure 1 shows the aOR for presentation with AHD and sex-stratified educational level. The gradient by level of education was more prominent in men (P value for interaction 0.082). For instance, the aOR for uncompleted basic education versus tertiary education was 1.70 (95% CI 1.41–2.04) in men and 1.31 (95% CI 0.96–1.82) in women. The trend by educational level was maintained restricting our analysis to MSM; aORs were 1.88 (95% CI 1.35–2.61), 1.67 (95% CI 1.40–1.99), and 1.27 (95% CI 1.10–1.46) compared to patients with tertiary education (Table 2) (P value 0.225 for interaction test).

There were 9486 (62%) individuals with LHD at presentation. Among patients with uncompleted basic, basic, secondary and tertiary educational level, the proportions of LHD were 73, 65, 59 and 55%. Trends in proportion of LHD and aOR by level of education were similar to the AHD analysis (Appendix 1, <http://links.lww.com/QAD/A557>).

Sensitivity analyses

Individuals with unknown level of education had similar demographic characteristics and similar proportions of AHD and LHD, but were more likely to have an unknown transmission mode and geographical origin (Appendix 2, <http://links.lww.com/QAD/A557>). Results were

Table 1. Patient characteristics at HIV diagnosis and proportion who presented with advanced HIV disease by baseline characteristics, overall and stratified by completed level of education.

Characteristics at HIV diagnosis	N	N with AHD	%	Proportion with AHD by completed level of education			
				Uncompleted basic	Basic	Secondary	Tertiary
Overall	15414	6129	40%	753 (52%)	1972 (45%)	2667 (37%)	737 (31%)
Sex							
Male	11667	4591	39%	444 (54%)	1392 (48%)	2106 (36%)	649 (30%)
Female	3737	1538	41%	309 (49%)	580 (40%)	561 (39%)	88 (41%)
Calendar year							
<2001	4763	2130	45%	321 (55%)	614 (43%)	998 (43%)	197 (44%)
≥2001	10651	3999	38%	432 (50%)	1358 (46%)	1669 (34%)	540 (28%)
Age (years)							
>35	7436	3665	49%	392 (59%)	1142 (56%)	1674 (47%)	457 (40%)
≤35	7978	2464	31%	361 (46%)	830 (36%)	993 (27%)	280 (23%)
Geographical origin							
European	11960	4691	39%	422 (52%)	1474 (46%)	2218 (37%)	577 (31%)
Non-European	2484	1093	44%	292 (52%)	349 (45%)	341 (41%)	111 (34%)
Unknown	970	345	35%	39 (43%)	149 (41%)	108 (32%)	49 (27%)
Transmission mode							
MSM	6427	1995	31%	84 (45%)	394 (39%)	1090 (31%)	427 (26%)
Heterosexual – men	3313	1675	51%	221 (57%)	624 (55%)	682 (47%)	148 (46%)
Heterosexual – women	3070	1259	41%	252 (49%)	469 (40%)	465 (39%)	73 (39%)
IDU – men	1299	584	45%	86 (54%)	274 (45%)	204 (42%)	20 (49%)
IDU – women	423	165	39%	22 (34%)	79 (40%)	61 (39%)	3 (75%)
Other/unknown – men	638	337	53%	53 (62%)	100 (62%)	130 (48%)	54 (45%)
Other/unknown – women	244	114	47%	35 (67%)	32 (41%)	35 (38%)	12 (55%)
European region of the cohort							
Austria, France and Switzerland	3930	1384	35%	133 (45%)	287 (41%)	836 (34%)	128 (28%)
Greece, Italy and Spain	8508	3449	41%	445 (54%)	1443 (47%)	1079 (36%)	482 (30%)

COHERE in EuroCoord, 1996–2011. Advanced HIV disease: a CD4⁺ cell count below 200 cells/ μ l or an AIDS-defining event in the 6 months following HIV diagnosis. AHD, advanced HIV disease; IDU, injecting drug users.

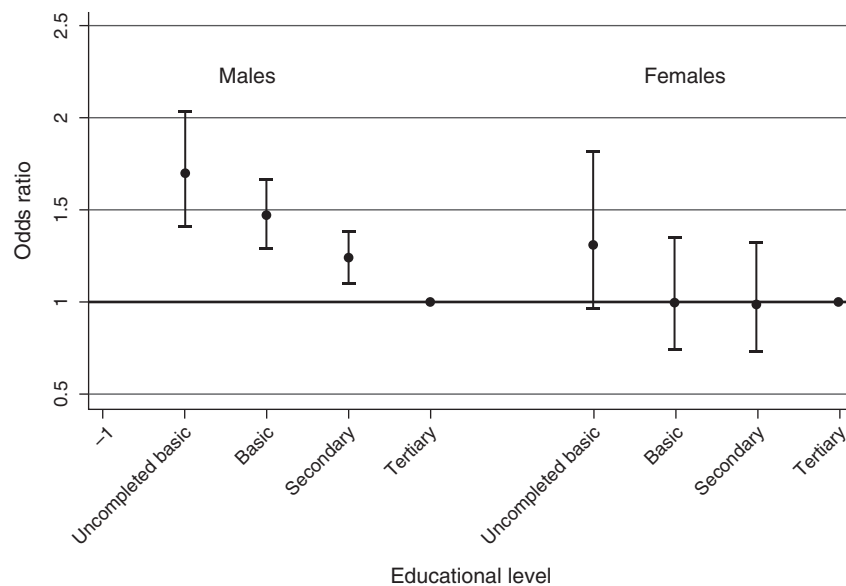


Fig. 1. Multivariate analyses of presentation with advanced HIV disease by sex. Estimates from multivariable logistic models adjusted by transmission mode, geographical origin, age at HIV diagnosis, calendar period and cohort. Test for interaction: $P=0.082$. COHERE in EuroCoord, 1996–2011.

consistent in all sensitivity analyses (Appendices 3 and 4, <http://links.lww.com/QAD/A557>).

Delayed initiation of antiretroviral therapy

Of the 15414 individuals included in the previous analysis, 72% (11 035) initiated cART. These patients were mostly men (78%), MSM, had initiated cART after a median (IQR) of 2 (1–7) months after HIV diagnosis with a median (IQR) CD4⁺ cell count of 218 (90–331) cells/ μ l and age 36 (31–42) years. Overall, results from the median regression analyses indicate a strong association between CD4⁺ cell count distribution at cART initiation and educational level ($P<0.001$ heterogeneity test). Notably, patients with uncompleted basic, basic and secondary education had estimated median CD4⁺ cell counts of 49, 27 and 19 cells/ μ l lower than patients with tertiary education at the time of cART initiation (Table 3). When the analyses were limited to the 5906 individuals who did not present with AHD, we found a trend of lower CD4⁺ cell count with lower educational level, but it was not statistically significant (Table 4). The logistic regression analysis for late cART initiation, defined as CD4⁺ cell count below 350 cells/ μ l,

showed a statistically significant association between CD4⁺ cell count distribution at cART initiation and educational level.

Discussion

We found that in Europe in the cART era, individuals with lower educational level were substantially more likely to present with AHD and LHD, even after taking into account individual characteristics that are traditionally associated with delayed HIV diagnosis. This gradient was more marked for men than for women. Among patients who initiated cART, lower level of education was independently associated with lower median CD4⁺ cell count at cART initiation.

The mechanisms underlying the observed associations are likely to implicate a number of material and psychosocial pathways through which education influences attitudes toward HIV testing and cART initiation. As a surrogate of socioeconomic status, individuals with higher education have better material resources such as employment and

Table 2. Multivariate analyses of presentation with advanced HIV disease by transmission category.

	MSM	IDU	Heterosexual	Other/unknown
Level of education	<i>N</i> = 6427	<i>N</i> = 1722	<i>N</i> = 6383	<i>N</i> = 881
Uncompleted basic	1.88 (1.35, 2.61)	1.20 (0.61, 2.34)	1.39 (1.11, 1.74)	1.64 (0.97, 2.81)
Basic	1.67 (1.40, 1.99)	0.93 (0.50, 1.74)	1.13 (0.92, 1.38)	1.02 (0.64, 1.63)
Secondary	1.27 (1.10, 1.46)	0.99 (0.52, 1.87)	1.24 (1.10, 1.39)	0.89 (0.58, 1.36)
Tertiary	1	1	1	1
	$P<0.001$	$P=0.472$	$P=0.003$	$P=0.073$

Estimates from multivariable logistic models adjusted by sex, geographical origin, age at HIV diagnosis, calendar period and cohort. Test for interaction: $P=0.225$. COHERE in EuroCoord, 1996–2011. AHD, advanced HIV disease; IDU, injecting drug users.

Table 3. Distribution of CD4⁺ at combination antiretroviral therapy initiation and multivariable analyses.

	N	Median (IQR) CD4 ⁺ cell count at cART initiation	Proportion with CD4 ⁺ <350 at cART	Multivariable analysis		
				Median regression		Logistic regression
				Median CD4 ⁺ count (95% CI)	P	Odds ratios for CD4 ⁺ <350 at cART (95% CI) P
Overall	11035	218 (90 331)	78%	Baseline 231 (216 248)		
Level of education					<0.001	<0.001
No basic	1036	173 (55 294)	86%	−49 (−67, −30)		1.71 (1.36, 2.14)
Basic	3288	198 (70 316)	82%	−27 (−41, −12)		1.23 (1.05, 1.45)
Secondary	5117	238 (101 342)	77%	−19 (−32, −6)		1.07 (0.93, 1.24)
Tertiary	1594	251 (126 345)	77%	–		1

Estimates from multivariable median regression and logistic regression models adjusted by calendar period, sex, transmission mode, geographical origin, age at cART initiation and cohort. COHERE in EuroCoord, 1996–2011. cART, combination antiretroviral therapy; CI, confidence interval; IQR, interquartile range.

higher-paying occupations [26], which imply easier access to healthcare facilities. People with higher education are more likely to practice health-promoting behaviours, including timely healthcare check-ups and screenings [27] and, therefore, might be more likely to test for HIV when they perceive themselves at increased risk. Higher education increases people’s health literacy and cognitive skills, enabling them to make better informed health-related choices [4,27], including the importance of appropriate HIV testing and timely initiation of cART with better access to websites and community resources. Finally, education is linked with social and psychological factors, including sense of control, social standing and social support [26], and individuals with higher education may face fewer barriers to access HIV care and be more resilient to stigma [28].

Interestingly, the association between lower education and lower CD4⁺ cell count at cART initiation was substantially reduced though did not disappear, when the analyses were restricted to individuals with timely HIV diagnosis. Therefore, the observed association between

delayed cART initiation and lower educational level could be largely, but not solely, attributed to patterns of delayed HIV diagnosis by educational level. This highlights the existence of additional socioeconomic barriers that deter access to cART after HIV diagnosis.

The observed differences concord with previous evidence from Spain and Italy indicating a higher frequency of delayed HIV diagnosis and cART initiation among individuals of lower educational level [19,20], and build on previous work conducted within COHERE showing very high levels of delayed HIV diagnoses in Europe [11]. We show in this study that not only presentation with AHD and LHD is common across all educational level groups, but that it exhibits an increasing trend with decreasing educational levels. The gradient of the association between level of education and delayed diagnosis is more remarkable for AHD than for LHD. Thus, socioeconomic inequalities are particularly visible in patients with very low CD4⁺ cell count levels, usually associated with high risk of AIDS and mortality [29]. These results are compatible with inequity in access to

Table 4. Distribution of CD4⁺ cell count at combination antiretroviral therapy initiation and multivariable analyses restricting to individuals who did not present with advanced HIV diagnosis.

	N	Median (IQR) CD4 ⁺ count at cART initiation	Proportion with CD4 ⁺ <350 at cART	Multivariable analysis		
				Median regression		Logistic regression
				Median CD4 ⁺ cell count (95% CI)	P	Odds ratios for CD4 ⁺ <350 at cART (95% CI) P
Overall	5906	314 (250 410)	62%	Baseline 312 (297 326)		
Level of education					0.132	0.006
No basic	441	305 (245 399)	42%	−18 (−34, −2)		1.44 (1.11, 1.88)
Basic	1517	310 (246 419)	36%	−9 (−21, 3)		1.12 (0.93, 1.35)
Secondary	2923	319 (250 430)	38%	−4 (−14, 6)		0.99 (0.84, 1.16)
Tertiary	965	315 (256 408)	37%	–		1

Estimates from multivariable regression and logistic regression models adjusted by calendar period, sex, transmission mode, geographical origin, age at cART initiation and cohort. COHERE in EuroCoord, 1996–2011. cART, combination antiretroviral therapy; CI, confidence interval; IQR, interquartile range.

HIV testing and are worrisome considering that the study was conducted in six European countries with universal public health systems.

The study shows that the association of delayed HIV diagnosis with educational level differed between men and women in absolute and relative terms. The milder gradient in AHD by educational level observed in women might be explained by universal HIV testing offered to all pregnant women in European countries for prevention of mother-to-child HIV transmission [30]. These findings, however, need to be put in the context of the literature that underlies that economic position may be a poorer predictor for a number of health outcomes in women [31]. The proportions of AHD for women from all four educational levels are lower than those of the heterosexual men but, compared to MSM, women with secondary and tertiary education exhibit higher levels of AHD. MSM, one of the key target groups for HIV screening in most European countries [6], still had a steep decline in AHD by educational level. These results indicate that universal HIV screening approaches such as those aiming at all pregnant women, as an alternative to targeted HIV testing policies based on HIV risk perception such as those aimed at MSM, could have the additional benefit to decrease socioeconomic inequities in accessing HIV testing. Whereas the former approach would offer a voluntary HIV test to all pregnant women, for MSM to be offered an HIV test, disclosure of unprotected sex with other men and/or gay identity becomes a prerequisite which is likely to be influenced by socioeconomic position [28,32].

For a long time, the interest to collect patient's socioeconomic information in HIV cohorts has been rather limited, though some groups have strongly promoted it [33]. The advantages of this study which harmonized socioeconomic variables across several cohorts in European countries are its large sample size, allowing exploring interactions by sex and transmission group, the inclusion of patients under routine care and its standardised definition of the exposure variable across European countries. The pan-European perspective of the study suggests that the observed gradient of increased risk of late HIV diagnosis and late cART initiation with lower educational level is present across European countries with different HIV healthcare and education systems. The study also has some limitations. Not all cohorts in COHERE collect data on educational level, so this study was based on data from only six European countries. Results thus might not be generalizable to settings with different social systems such as those from northern and eastern Europe. Some bias may have been introduced through misclassification since an individual's educational level might not always reflect their socioeconomic status and whether this error is different for men and women. In smaller-sized cohort studies, combined indicators such as living on welfare, unstable housing and unemployment were found to be associated

with non-adherence, whereas no association could be found with each component alone [34]. Protopopescu *et al.* have recently described how an indicator combining low educational level and unemployment was found associated with higher rates of mortality in the APROCO cohort while none of each separate variable had a significant effect [35]. However, educational level was the only variable collected by the six participating cohorts and it is important to note that education is one of the most used measures of socioeconomic status in healthcare research because of its influence on future occupational opportunities and earning potential [14]. Foreigners unfamiliar with the educational system of the host country may have misreported their educational level. However, these forms of misclassification are likely to be non-differential, resulting in an underestimate of the relationship between socioeconomic status and delayed HIV diagnosis and cART initiation. Despite this potential bias, we found significant trends and a clear gradient by educational level, and we believe this strengthens, rather than weakens, our conclusions about the existence of socioeconomic inequalities. Finally, a number of individuals were excluded from the analyses because their educational level was unknown or no CD4⁺ cell count measurement was available at the time of HIV diagnosis. Our estimates might be biased if the included and excluded patients had a different distribution of educational level and CD4⁺ cell count at HIV diagnosis. Our conclusions were, however, robust to a set of sensitivity analyses exploring different scenarios of missing data mechanisms.

In conclusion, this study shows that inequalities by educational level, a proxy of a socioeconomic status, in HIV testing and initiation of cART are present in European countries with universal healthcare systems, and thus, individuals with lower educational level will not equally benefit from the effectiveness of cART. Policies and interventions that target socioeconomic determinants leading to delays in HIV diagnosis and cART initiation are needed. Whether the observed inequalities are all avoidable, and thus amendable, is a discussion to be urgently advanced within the equity policy framework for Europe Health 2020 [36] and more deeply taken into account in clinical and epidemiological research.

Socio-economic inequalities and HIV Writing Group

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References

1. Dray-Spira R, Lert F. **Social health inequalities during the course of chronic HIV disease in the era of highly active antiretroviral therapy.** *AIDS* 2003; **17**:283–290.
2. Marmot M, Friel S, Bell R, Houweling TA, Taylor S. **Closing the gap in a generation: health equity through action on the social determinants of health.** *Lancet* 2008; **372**:1661–1669.
3. Marmot M, Allen J, Bell R, Bloomer E, Goldblatt P. **WHO European review of social determinants of health and the health divide.** *Lancet* 2012; **380**:1011–1029.
4. Cutler DM, Lleras-Muney A. **Understanding differences in health behaviors by education.** *J Health Econ* 2010; **29**:1–28.
5. van Doorslaer E, Koolman X, Jones AM. **Explaining income-related inequalities in doctor utilisation in Europe.** *Health Econ* 2004; **13**:629–647.
6. European Centre for Disease Prevention and Control. *HIV testing: increasing uptake and effectiveness in the European Union.* Stockholm: ECDC; 2010.
7. Hamers FF, Downs AM. **The changing face of the HIV epidemic in western Europe: what are the implications for public health policies?** *Lancet* 2004; **364**:83–94.
8. European Centre for Disease Control/World Health Organisation. *HIV/AIDS surveillance in Europe.* Stockholm: ECDC/WHO; 2010.
9. European Centre for Disease Prevention and Control. *Thematic report: HIV treatment, care and support. Monitoring implementation of the Dublin Declaration on Partnership to Fight HIV/AIDS in Europe and Central Asia: 2012 progress report.* Stockholm: ECDC; 2013.
10. Royo-Bordonada MA, Diez-Cornell M, Llorente JM. **Health-care access for migrants in Europe: the case of Spain.** *Lancet* 2013; **382**:393–394.
11. Mocroft A, Lundgren JD, Sabin ML, Monforte A, Brockmeyer N, Casabona J, et al. **Risk factors and outcomes for late presentation for HIV-positive persons in Europe: results from the Collaboration of Observational HIV Epidemiological Research Europe Study (COHERE).** *PLoS Med* 2013; **10**:e1001510.
12. Marks G, Crepaz N, Senterfitt JW, Janssen RS. **Meta-analysis of high-risk sexual behavior in persons aware and unaware they are infected with HIV in the United States: implications for HIV prevention programs.** *J Acquir Immune Defic Syndr* 2005; **39**:446–453.
13. Moreno S, Mocroft A, Monforte A. **Medical and societal consequences of late presentation.** *Antivir Ther* 2010; **15** (Suppl 1): 9–15.
14. Adler NE, Newman K. **Socioeconomic disparities in health: pathways and policies.** *Health Aff (Millwood)* 2002; **21**:60–76.
15. Shavers VL. **Measurement of socioeconomic status in health disparities research.** *J Natl Med Assoc* 2007; **99**:1013–1023.
16. Davey Smith G, Hart C, Hole D, MacKinnon P, Gillis C, Watt G, et al. **Education and occupational social class: which is the more important indicator of mortality risk?** *J Epidemiol Commun Health* 1998; **52**:153–160.
17. Regidor E, De Mateo S, Calle ME, Dominguez V. **Educational level and mortality from infectious diseases.** *J Epidemiol Commun Health* 2002; **56**:682–683.
18. Wolbers M, Bucher HC, Furrer H, Rickenbach M, Cavassini M, Weber R, et al. **Delayed diagnosis of HIV infection and late initiation of antiretroviral therapy in the Swiss HIV Cohort Study.** *HIV Med* 2008; **9**:397–405.
19. Sobrino-Vegas P, Rodriguez-Urrego J, Berenguer J, Caro-Murillo AM, Blanco JR, Viciano P, et al. **Educational gradient in HIV diagnosis delay, mortality, antiretroviral treatment initiation and response in a country with universal healthcare.** *Antivir Ther* 2012; **17**:1–8.

20. Girardi E, Aloisi MS, Arici C, Pezzotti P, Serraino D, Balzano R, et al. **Delayed presentation and late testing for HIV: demographic and behavioral risk factors in a multicenter study in Italy.** *J Acquir Immune Defic Syndr* 2004; **36**:951–959.
21. Karanikolos M, Mladovsky P, Cylus J, Thomson S, Basu S, Stuckler D, et al. **Financial crisis, austerity, and health in Europe.** *Lancet* 2013; **381**:1323–1331.
22. Antinori A, Coenen T, Costagliola D, Dedes N, Ellefson M, Gatell J, et al. **Late presentation of HIV infection: a consensus definition.** *HIV Med* 2011; **12**:61–64.
23. Royston P, Ambler G, Sauerbrei W. **The use of fractional polynomials to model continuous risk variables in epidemiology.** *Int J Epidemiol* 1999; **28**:964–974.
24. Sterne JA, White IR, Carlin JB, Spratt M, Royston P, Kenward MG, et al. **Multiple imputation for missing data in epidemiological and clinical research: potential and pitfalls.** *Br Med J* 2009; **338**:b2393.
25. Royston P. **Multiple imputation of missing values: further update of ice, with an emphasis on interval censoring.** *Stata J* 2007; **7**:445–464.
26. Ross C, Wu C. **The links between education and health.** *Am Sociol Rev* 1995; **60**:719–745.
27. Cutler D, Lleras-Muney A. **Education and health: evaluating theories and evidence.** National Bureau of Economic Research. Bethesda, MD: BNER; 2006.
28. McGarrity LA, Huebner DM. **Is being out about sexual orientation uniformly healthy? The moderating role of socioeconomic status in a prospective study of gay and bisexual men.** *Ann Behav Med* 2013; **47**:28–38.
29. Phillips A, Pezzotti P. **Short-term risk of AIDS according to current CD4 cell count and viral load in antiretroviral drug-naïve individuals and those treated in the monotherapy era.** *AIDS* 2004; **18**:51–58.
30. Deblonde J, Claeys P, Temmerman M. **Antenatal HIV screening in Europe: a review of policies.** *Eur J Public Health* 2007; **17**:414–418.
31. Mustard CA, Etches J. **Gender differences in socioeconomic inequality in mortality.** *J Epidemiol Commun Health* 2003; **57**:974–980.
32. White D, Stephenson R. **Identity formation, outness, and sexual risk among gay and bisexual men.** *Am J Mens Health* 2013; **8**:98–109.
33. Lewden C, Raffi F, Cuzin L, Cailleton V, Vilde JL, Chene G, et al. **Factors associated with mortality in human immunodeficiency virus type 1-infected adults initiating protease inhibitor-containing therapy: role of education level and of early transaminase level elevation (APROCO-ANRS EP11 study).** The Antiproteases Cohorte Agence Nationale de Recherches sur le SIDA EP 11 study. *J Infect Dis* 2002; **186**:710–714.
34. Bouhnik AD, Chesney M, Carrieri P, Gallais H, Moreau J, Moatti JP, et al. **Nonadherence among HIV-infected injecting drug users: the impact of social instability.** *J Acquir Immune Defic Syndr* 2002; **31** (Suppl 3):S149–153.
35. Protopopescu C, Carrieri MP, Le Moing V, Reboud P, Piroth L, Cuzin L, et al. **Socio-behavioural determinants of mortality in HIV-infected patients receiving combined antiretroviral treatment (cART): results from the ANRS CO8 APROCO-COPILOTE cohort.** 7th IAS Conference on HIV Pathogenesis, Treatment and Prevention, Kuala Lumpur; 2013.
36. World Health Organization. **Regional Office for Europe Health 2020: a European policy framework supporting action across government and society for health and well being.** Malta, 10-13 September 2012.