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Provider Compliance With Guidelines for Management of Cardiovascular Risk in HIV-Infected Patients

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Learning Objectives

Upon completion of this activity, participants will be able to:

- Assess the association between HIV infection and cardiovascular risk factors
- Analyze the cardiovascular risk profile of patients with HIV in the current study
- Evaluate physician adherence to cardiovascular risk treatment among patients with HIV
- Distinguish variables associated with not receiving recommended treatment for cardiovascular risk factors

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PEER REVIEWED

Abstract

Introduction

Compliance with National Cholesterol Education Program Adult Treatment Panel III (NCEP) guidelines has been shown to significantly reduce incident cardiovascular events. We investigated physicians' compliance with NCEP guidelines to reduce cardiovascular disease (CVD) risk in a population infected with HIV.

Methods

We analyzed HIV Outpatient Study (HOPS) data, following eligible patients from January 1, 2002, or first HOPS visit thereafter to calculate 10-year cardiovascular risk (10yCVR), until September 30, 2009, death, or last office visit. We categorized participants into four 10yCVR strata, according to guidelines determined by NCEP, the Infectious Disease Society of America, and the Adult AIDS Clinical Trials Group. We calculated percentages of patients treated for dyslipidemia and hypertension, calculated percentages of patients who achieved recommended goals, and categorized them by 10yCVR stratum.

Results

Of 2,005 patients analyzed, 33.7% had fewer than 2 CVD risk factors. For patients who had 2 or more risk factors, 10yCVR was less than 10% for 28.2%, 10% to 20% for 18.2%, and higher than 20% for 20.0% of patients. Of patients eligible for treatment, 81% to 87% were treated for elevated low-density lipoprotein cholesterol/non-high-density lipoprotein cholesterol (LDL-C/non-HDL-C), 2% to 11% were treated for low HDL-C, 56% to 91% were treated for high triglycerides, and 46% to 69% were treated for hypertension. Patients in higher 10yCVR categories were less likely to meet treatment goals than patients in lower 10yCVR categories.

Conclusion

At least one-fifth of contemporary HOPS patients have a 10yCVR higher than 20%, yet a large percentage of at-risk patients who were eligible for pharmacologic treatment did not receive recommended interventions and did not reach recommended treatment goals. Opportunities exist for CVD prevention in the HIV-infected population.

Introduction

Compliance with guidelines established in 2001 by the National Cholesterol Education Program Adult Treatment Panel III (NCEP) for primary and secondary prevention of myocardial infarction significantly reduces incident cardiovascular events (1–5). In 2003, the Infectious Disease Society of America and the Adult AIDS Clinical Trials Group published a modification of the 2001 NCEP guidelines to address management of cardiovascular risk (CVR) in HIV-1 infected people (6). In addition to describing management of traditional CVR factors, the modification included recommendations for altering antiretroviral therapy (ART). We analyzed the degree to which the modified NCEP guidelines have been followed by physicians in the HIV Outpatient Study (HOPS) cohort and the effectiveness of these interventions in reaching NCEP treatment goals designed to reduce incident cardiovascular events.

Methods

HIV Outpatient Study cohort

The HOPS is an ongoing, prospective, observational cohort study of HIV-infected patients receiving care in 10 participating HIV clinics (4 universities, 2 public clinics, 4 private clinics) in 8 US cities (Chicago, Illinois; Denver, Colorado; Stony Brook, New York; Oakland/San Leandro, California; Philadelphia, Pennsylvania; Tampa, Florida; and Washington, DC) since March 1993. The cohort represents a convenience sample of patients receiving HIV care at the sites; enrollment varies by clinic size. Patient data, including sociodemographic characteristics, symptoms, signs, diagnoses, treatments, and laboratory values, are abstracted from medical charts and entered into an electronic database by trained staff. Data are reviewed for quality and analyzed centrally. The HOPS protocol is reviewed and

approved annually by the Centers for Disease Control and Prevention and each local site's institutional review board. This analysis used HOPS data from January 1, 2002, through September 30, 2009.

Study population

Entry criteria for participation in HOPS are documented HIV infection and informed consent to participate in the study. To assess achievement of treatment goals based on the 2001 NCEP guidelines, we included HOPS patients active on or after January 1, 2002, to allow clinicians the opportunity to respond to the 2001 guidelines. We defined "active" patients as those who had attended 2 or more office visits since January 1, 2002. The categorization of patients by their baseline 10-year CVR (10yCVR) required that eligible patients have had 2 or more blood pressure readings recorded and at least 1 fasting lipid panel obtained as baseline measurements any time between 12 months before and up to 9 months after the baseline date, and at least 1 year after baseline. Patients were followed until September 30, 2009, death, or last office visit.

We assessed whether the percentages of patients treated per modified NCEP guidelines and achieving recommended goals at any time during the follow-up period differed across progressively higher categories of baseline 10yCVR and according to key baseline demographic characteristics. We also calculated rates of incident cardiovascular events per 100 person-years of observation by 10yCVR group for descriptive purposes because of the small number of cardiovascular events.

Classification of patients into 10yCVR categories

We categorized HOPS patients by baseline 10yCVR into 4 categories per NCEP guidelines (2). These guidelines categorize risk using a combination of the Framingham 10-year absolute risk for developing coronary heart disease (CHD) and conditions defined as CHD equivalents (eg, diabetes mellitus [diabetes], prior cardiovascular event). A Framingham risk calculation was not performed in patients who had low risk (ie, ≤ 1 major risk factor) and for whom 10yCVR was less than 10%. A Framingham risk calculation was performed among patients with 2 or more major risk factors, and they were categorized as follows: those with a 10yCVR of less than 10% were at moderate risk, those with a 10yCVR of 10% to 20% were at moderately high risk, and those with either CHD, a CHD equivalent, or a 10yCVR higher than 20% were at high risk.

Major risk factors for CHD included cigarette smoking (past or current), hypertension (systolic/diastolic blood pressure $\geq 140/90$ mm Hg; or for patients diagnosed with diabetes or chronic renal insufficiency, $\geq 130/80$ mm Hg; or prescription of antihypertensive therapy with a diagnosis of hypertension, regardless of blood pressure), elevated serum low-density lipoprotein cholesterol (LDL-C), serum high-density lipoprotein cholesterol (HDL-C) less than 40 mg/dL for men and less than 50 mg/dL for women, a family history of premature CHD (first-degree male relative aged < 55 y, first-degree female relative aged < 65 y), and older age (men, ≥ 45 y; women, ≥ 55 y). CHD equivalents are conditions that pose a CHD event risk equal to the risk for having a new CHD event in people with established CHD; these include known noncardiac vascular disease (peripheral arterial disease, abdominal aortic aneurysm, and symptomatic carotid artery disease) and diabetes. We compared the 2,005 patients with those who were excluded because of missing 1 or more required lipid or blood pressure readings.

Clinical interventions for lipid and blood pressure management are made according to 10yCVR risk category. A switch in ART at any time during the study period from a regimen associated with lipid abnormalities or insulin resistance to a regimen less likely to have these associations was considered an appropriate intervention (7–14).

NCEP treatment goals and thresholds for medical intervention

Conditions for modification are 1) LDL-C in people with triglyceride levels ≤ 200 mg/dL and 2) non-HDL-C (total cholesterol minus HDL-C) when triglycerides exceed 200 mg/dL. Recommended values of LDL-C and non-HDL-C at which to initiate therapeutic lifestyle changes and/or drug therapy, as well as target values (goal), differ by each of the 4 NCEP-defined 10yCVR categories (Table 1). We included 3 additional conditions for modification to reduce CVR: low HDL-C, hypertriglyceridemia, and hypertension. We defined the HDL-C value below which to initiate lipid-modifying therapy and at or above which we considered goal to have been achieved as 40 mg/dL in men and 50 mg/dL in women. We also defined a triglyceride level of ≥ 500 mg/dL (the level at which triglycerides adversely affect LDL-C metabolism) as the level at which medical intervention should be initiated, with a triglyceride level of less than 500 mg/dL as goal. We considered treatment of triglycerides between 200 mg/dL and 499 mg/dL as optional. Treated hypertensive patients achieved goal if blood pressure was less than 140/90 mm Hg (or $< 130/80$ mm Hg with concurrent diabetes or chronic renal insufficiency).

We defined the presence of metabolic syndrome as having at least 3 of the following: fasting triglycerides of at least 150 mg/dL, low HDL-C, fasting blood glucose of at least 110 mg/dL, systolic blood pressure of at least 130 mm Hg or diastolic blood pressure of at least 85 mm Hg, and a body mass index (BMI) of at least 28.9 kg/m² for men or 24.9 kg/m² for women.

We defined appropriate treatment as prescription of a 3-hydroxy-3-methylglutaryl-CoA (HMG CoA) reductase inhibitor (statin), with or without ezetimibe, or bile acid sequestrants for elevated LDL-C/non-HDL-C (Table 1); of nicotinic acid, documentation of exercise, or both for low HDL-C; of fibrates or omega-3 fatty acids for hypertriglyceridemia, and of antihypertensives for hypertension. However, data on initiation of therapeutic lifestyle changes such as increased exercise were not systematically collected in this cohort. We also considered changes in ART regimens as appropriate treatment.

Statistical analyses

Summaries of descriptive data, univariate analyses, and Cochrane–Armitage tests for trend were conducted with SAS version 9.2 (SAS Institute, Inc, Cary, North Carolina). We compared the distributions of categorical variables using a likelihood ratio, continuity-adjusted χ^2 test, or Fisher exact test, and we compared the distributions of continuous variables with a Wilcoxon rank-sum test for 2-group comparisons. We used a χ^2 test and Kruskal–Wallis test for comparisons of more than 2 groups. We computed incidence rates of cardiovascular events per 100 person-years of observation for each 10yCVR category and compared them across 10yCVR categories using StatCalc (EpiInfo 2000, Centers for Disease Control and Prevention, Atlanta, Georgia). Proportion confidence intervals were calculated using mid-*P* exact confidence limits (15). Statistical associations with 2-sided *P* values less than .05 were considered significant.

Results

Our study population of 2,005 HOPS patients (baseline median age, 42 y; median CD4+ T cell count, 395 cells/mm³) was predominantly male (76%) and racially/ethnically diverse (52% non-Hispanic white, 33% non-Hispanic black, and 12% Hispanic); 55% of patients were current or former tobacco users at baseline. Seventeen percent of patients had no ART exposure, 77.7% had exposure to highly active ART (HAART) (45.3% had prior mono- or dual-antiretroviral [ARV] exposures; 32.4% had HAART only), and 5.1% were classified as having unknown or missing data on ARV exposure. Of the 2,005 HOPS patients analyzed, 675 were categorized at baseline as low risk, 565 as moderate risk, 365 as moderately high risk, and 400 as high risk. Median follow-up after baseline was approximately 5.5 years and did not differ significantly across 10yCVR categories (Table 2).

At baseline, as 10yCVR increased, the percentage of people with the following characteristics increased significantly: those with a diagnosis of either hypertension or metabolic syndrome; median serum total cholesterol and triglycerides; and those prescribed lipid-lowering agents. Median serum HDL-C decreased significantly with increasing 10yCVR categories, whereas LDL-C, the distribution by race/ethnicity, and baseline use of ART did not vary consistently (Table 2).

Excluded patients were significantly younger (median age, 40 vs 42 y); more likely to be male (82.0% vs 76.4%); more likely to be white (56.2% vs 52.3%); more likely to be privately insured (65.5% vs 57.2%); and less likely to have had an AIDS-defining illness (22.0% vs 36.3%), diabetes (4.3% vs 9.5%), hypertension (27.5% vs 51.7%), or BMI higher than 30 kg/m² (12.7% vs 19.4%); and more likely to have unknown or missing ARV history (17.0% vs 5.1%) at baseline.

As 10yCVR increased, the percentage of people with each cardiovascular condition of interest increased (Table 3). The percentage of patients treated among those patients with each condition of interest also increased; the highest percentage was treated for high LDL-C and hypertriglyceridemia. However, the percentage of patients treated according to modified NCEP guidelines and who met guideline-defined goals at any time during the follow-up period for each condition decreased across each of the 4 CVR categories, particularly in the high-risk group.

Women, nonwhites, and patients with public insurance or no insurance were treated for hypertension per guidelines more often than men, whites, or patients with private insurance, despite a lower representation of the former patients in the higher CVR categories. Pharmacologic treatment of low HDL-C was more frequent in whites and men but overall treatment rates were low. Management of LDL-C/non-HDL-C, HDL-C, and triglycerides was similar when these groups were compared (Figures 1, 2, and 3).

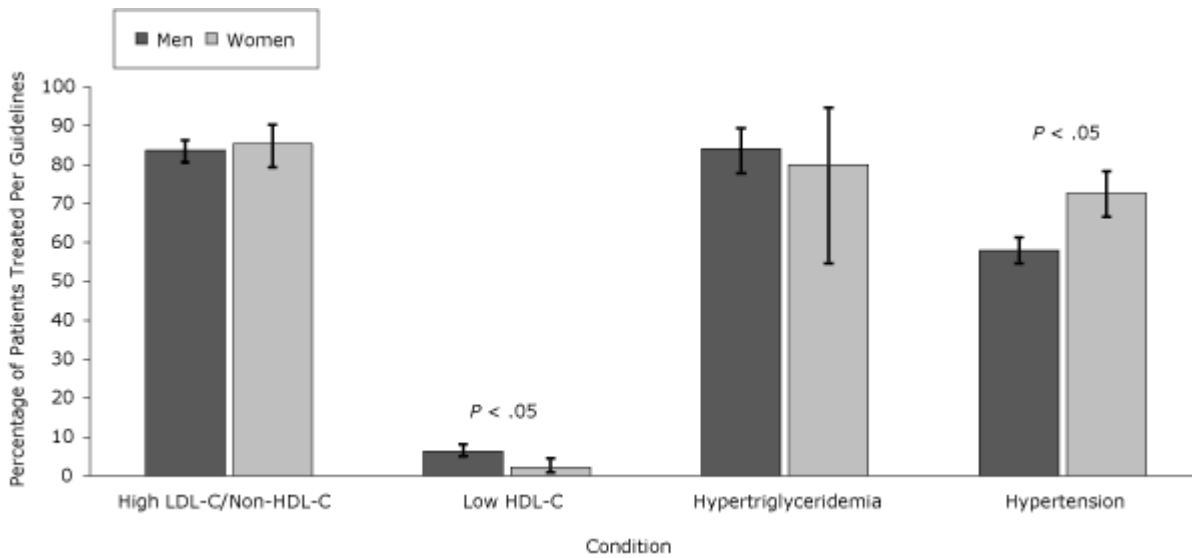


Figure 1. Percentage of patients with condition treated per National Cholesterol Education Program Adult Treatment Panel III (NCEP) guidelines, by sex, the HIV Outpatient Study, January 2002–September 2009. Error bars indicate 95% confidence intervals. Abbreviations: LDL-C, low-density lipoprotein cholesterol; HDL-C, high-density lipoprotein cholesterol. [A tabular version of this figure is also available.]

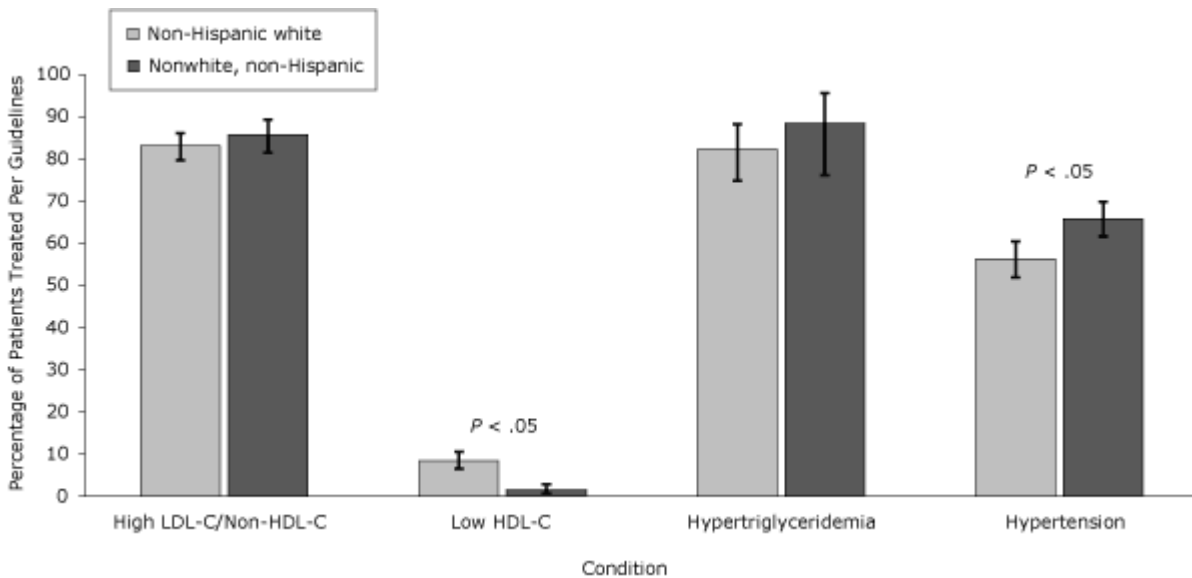


Figure 2. Percentage of patients with condition treated per National Cholesterol Education Program Adult Treatment Panel III (NCEP) Guidelines, by race/ethnicity, the HIV Outpatient Study, January 2002–September 2009. Error bars represent 95% confidence intervals. Abbreviations: LDL-C, low-density lipoprotein cholesterol; HDL-C, high-density lipoprotein cholesterol. [A tabular version of this figure is also available.]

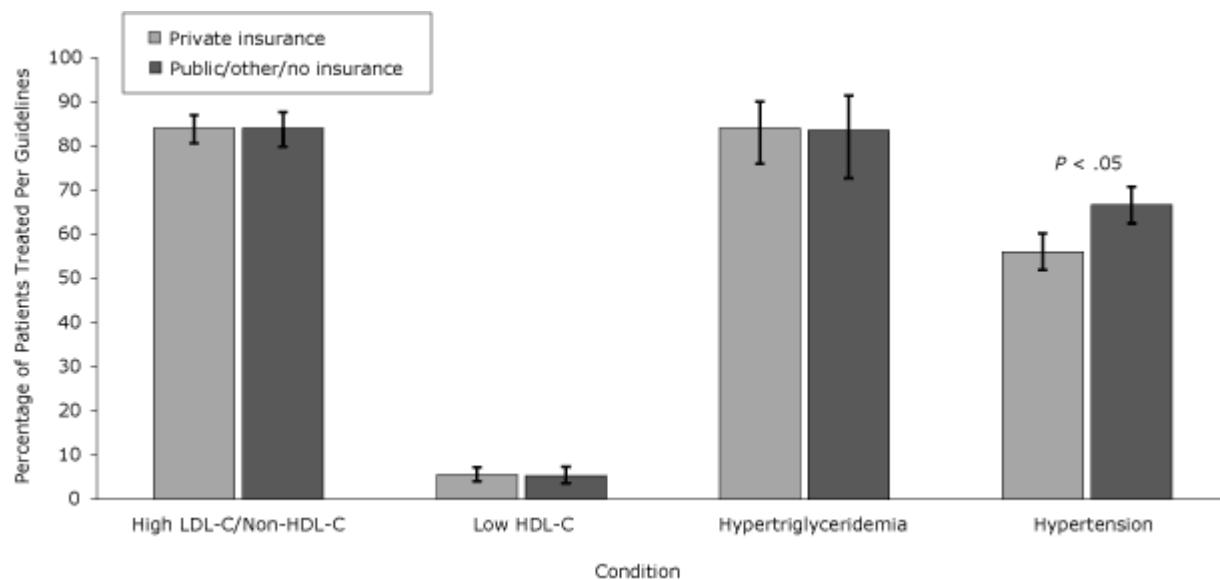


Figure 3. Percentage of patients with condition treated per National Cholesterol Education Program Adult Treatment Panel III (NCEP) Guidelines, by payer, the HIV Outpatient Study, January 2002–September 2009. Error bars represent 95% confidence intervals. Abbreviations: LDL-C, low-density lipoprotein cholesterol; HDL-C, high-density lipoprotein cholesterol. [A tabular version of this figure is also available.]

Incidence of cardiovascular events (per 100 person-years) increased with increasing 10yCVR category: 0.38 (14 events) in the low-risk group compared with 0.89 (27 events) in the moderate-risk group, 2.24 (45 events) in the moderately high-risk group, and 2.98 (62 events) in the high-risk group (all $P < .01$ compared with the low-risk group).

Discussion

HIV-infected patients are at higher risk and have a higher incidence of cardiovascular events than the general public (16–22). In our cohort, 38.2% of patients were in either the moderately high-risk or high-risk categories, of whom 77.9% were current or former smokers, 74.2% had hypertension, 71.5% had baseline elevated LDL-C levels, 70.5% had low HDL-C levels, and 35.8% had metabolic syndrome. The percentage of patients with high LDL-C/non-HDL-C and triglyceride levels, low HDL-C levels, and hypertension generally increased with increasing 10yCVR category. Although most affected patients were prescribed recommended therapy for high LDL-C/non-HDL-C and hypertriglyceridemia, only half were treated for hypertension and very few were treated pharmacologically for low HDL-C. We also observed lower frequency of hypertension treatment among men, people who were white, and people who were privately insured; these lower frequencies were not explained by collinearity or differences in 10yCVR distribution.

Several studies have demonstrated increases in total cholesterol, LDL-C, HDL-C and triglycerides after initiating ART in antiretroviral-naïve patients (23–32). In untreated advanced HIV infection, levels of total cholesterol, LDL-C, and HDL-C are typically low. It is unclear to what extent the lipid elevations observed after initiating treatment of HIV infection were an adverse effect of ART versus a return to health (33). In light of the increased CVR faced by HIV-infected patients, focusing efforts on improving lipid profiles and controlling hypertension as recommended in the modified NCEP recommendations seems prudent.

HDL-C deserves additional attention. Normalizing HDL-C levels is a tertiary goal of the NCEP guidelines (34). HIV-infected patients characteristically have low HDL-C levels, and although levels rise with ART, they rarely normalize (33). We also found a high percentage of people (62% men, 65% women) in our cohort with low HDL-C measurements compared with the 1999–2002 National Health and Nutrition Examination Survey (NHANES III) population (33% men, 20% women) (35). Only 5.4% of patients in our study received pharmacologic treatment of HDL-C, of whom only 26.5% achieved target levels. Possible explanations for low compliance may include concern about side effects, pill burden, designation of HDL-C as a tertiary target, and pharmaceutical marketing emphasis on triglyceride elevations.

Although much attention has focused on the effect of ART on elevating triglyceride levels, only 8.3% of patients in our cohort had levels exceeding 500 mg/dL at baseline. Of those patients, 83.8% received triglyceride-lowering therapy and 78.6% of those patients had reductions of triglycerides to less than 500 mg/dL. Elevated triglycerides directly contribute little to increased CVR. Triglycerides exceeding 500 mg/dL interfere with the release of LDL-C from small dense atherogenic particles, resulting in deposition of these particles into atheromatous plaques (36). Statins contribute to expression of LDL-C receptors with consequent reduction of the circulating LDL-C substrate. Reducing triglycerides to less than 500 mg/dL enhances the effectiveness of statin therapy via this mechanism.

Although other cohort studies of HIV-infected patients have evaluated CVD risk factors, including 10yCVR, few to our knowledge have assessed physician adherence to guidelines for management of these conditions (18,37–39). We found that 38.2% of our study population had a 10yCVR at or higher than 10% (moderately high risk and high risk) compared with 22% to 26% in the Data Collection on Adverse Events of Anti-HIV Drugs (D:A:D) study (40). In an analysis of the Multicenter AIDS Cohort Study (MACS) and the Women’s Interagency HIV Study (WIHS), Kaplan et al defined moderate CVD risk as a 10yCVR of 15% to 25% and high CVD risk as a 10yrCVR of 25% or higher or having a diagnosis of diabetes. They found that 2% of men and 1% of women had a moderate predicted CVD risk and 17% of men and 12% of women had a high predicted CVD risk (18).

Comparing NHANES III data with those from the HOPS cohort, 30.0% versus 41.5% of people had elevated baseline LDL-C. When considering only moderately high-risk and high-risk groups, 68.0% versus 71.5% of patients had high LDL-C, of whom 25.0% versus 83.7% received treatment. Of those patients treated, 39.1% versus 59.3% in the moderately high-risk group and 22.3% versus 33.5% in the high-risk group met NCEP goals. Although comparisons of NHANES III and HOPS data are complicated by differences in study design, population sociodemographic characteristics, laboratory measurements of lipids, and ascertainment of treatment interventions, findings suggest that there is still a significant opportunity for improvement in CVD risk management in HOPS patients.

This study, which used routinely collected medical abstraction data from HIV outpatients, had several limitations. Our inclusion criteria, which required certain baseline examinations, may have enriched our analysis cohort in people whom clinicians considered to be at higher risk of CVD and who had warranted those examinations. Whether the excluded patients were considered lower risk by their physicians or had unrecognized risk factors necessarily limits the overall findings in our study. We also cannot infer reasons for lack of intervention when apparently indicated. Failure to initiate recommended interventions may have resulted from patients deferring or declining therapy, contraindications to these therapies not recorded in HOPS chart abstractions, or clinician oversight. CVD risk factor data were not systematically charted, precluding our ability to assess whether interventions resulted in sustained responses over time. We did not have adequate information on family history of cardiovascular events in first-degree relatives, a major CHD risk factor. The absence of these data would likely lead us to underestimate the true prevalence of CVR in our population. Furthermore, we did not have information on waist size for most patients. We substituted BMI at or above 28.9 kg/m² for men and at or above 24.9 kg/m² for women for waist size greater than 40 inches in men and greater than 35 inches in women, when measured, for our estimation of metabolic syndrome, guided by findings from prior studies in HIV cohorts (33,41). The data on pharmacotherapeutic treatments may have been recorded incompletely, and patients may have received treatments outside of the HOPS clinics (eg, cardiology specialist consultations) that were also not captured by chart abstraction. Finally, we could not account for therapeutic lifestyle change interventions such as intentional smoking cessation or weight loss, improved diet, or increased exercise; these data are not collected systematically in the HOPS.

Our findings suggest that HOPS participants who comprise a heterogeneous convenience sample of HIV-infected patients seen at 10 HIV specialty clinics in the United States have significantly greater CVD risk than both the general US population and participants in other HIV cohorts. A large percentage of at-risk patients who were eligible for pharmacologic treatment did not receive recommended interventions and did not reach recommended treatment goals. As HIV-infected patients live longer, it is imperative that clinicians minimize CVD risk by optimizing patients’ lipid profiles and blood pressure and helping patients who use tobacco products to quit. Intensified educational efforts are needed and further research is warranted to understand the barriers to initiating recommended interventions and to achieving recommended treatment goals that are unique to HIV-infected patients.

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References

1. Cannon CP, Braunwald E, McCabe CH, Rader DJ, Rouleau JL, Belder R, et al. Intensive versus moderate lipid lowering with statins after acute coronary syndromes. *N Engl J Med* 2004;350(15):1495-504. Erratum in: *N Engl J Med* 2006;354(7):778.
2. Expert Panel on Detection, Evaluation, and Treatment of High Blood Cholesterol in Adults. Executive summary of the Third Report of the National Cholesterol Education Program (NCEP) Expert Panel on Detection, Evaluation, and Treatment of High Blood Cholesterol in Adults (Adult Treatment Panel III). *JAMA* 2001;285(19):2486-97. CrossRef [↗](#) PubMed [↗](#)
3. Heart Protection Study Collaborative Group. MRC/BHF Heart Protection Study of cholesterol lowering with simvastatin in 20,536 high-risk individuals: a randomised placebo-controlled trial [summary for patients in *Curr Cardiol Rep* 2002;4(6):486-7]. *Lancet* 2002;360(9326):7-22. CrossRef [↗](#) PubMed [↗](#)
4. Sever PS, Dahlöf B, Poulter NR, Wedel H, Beevers G, Caulfield M, et al. Prevention of coronary and stroke events with atorvastatin in hypertensive patients who have average or lower-than-average cholesterol concentrations, in the Anglo-Scandinavian Cardiac Outcomes Trial-Lipid Lowering Arm (ASCOT-LLA): a multicentre randomised controlled trial. *Lancet* 2003;361(9364):1149-58. CrossRef [↗](#) PubMed [↗](#)
5. Shepherd J, Blauw GJ, Murphy MB, Bollen EL, Buckley BM, Cobbe SM, et al. Pravastatin in elderly individuals at risk of vascular disease (PROSPER): a randomised controlled trial. *Lancet* 2002;360(9346):1623-30. CrossRef [↗](#) PubMed [↗](#)
6. Dubé MP, Stein JH, Aberg JA, Fichtenbaum CJ, Gerber JG, Tashima KT, et al. Guidelines for the evaluation and management of dyslipidemia in human immunodeficiency virus (HIV)-infected adults receiving antiretroviral therapy: recommendations of the HIV Medical Association of the Infectious Disease Society of America and the Adult AIDS Clinical Trials Group. *Clin Infect Dis* 2003;37(5):613-27. CrossRef [↗](#) PubMed [↗](#)
7. Gallant JE, Staszewski S, Pozniak AL, DeJesus E, Suleiman JM, Miller MD, et al. Efficacy and safety of tenofovir DF vs stavudine in combination therapy in antiretroviral-naïve patients: a 3-year randomized trial. *JAMA* 2004;292(2):191-201. CrossRef [↗](#) PubMed [↗](#)
8. Johnson M, Grinsztejn B, Rodriguez C, Coco J, DeJesus E, Lazzarin A, et al. Atazanavir plus ritonavir or saquinavir and lopinavir/ritonavir in patients experiencing multiple virological failures. *AIDS* 2005;19(7):685-94. CrossRef [↗](#) PubMed [↗](#)
9. Mills AM, Nelson M, Jayaweera D, Ruxrungtham K, Cassetti I, Girard PM, et al. Once-daily darunavir/ritonavir vs. lopinavir/ritonavir in treatment-naïve, HIV-1-infected patients: 96-week analysis. *AIDS* 2009;23(13):1679-88. CrossRef [↗](#) PubMed [↗](#)
10. Molina JM, Andrade-Villanueva J, Echevarria J, Chetchotisakd P, Corral J, David N, et al. Once-daily atazanavir/ritonavir compared with twice-daily lopinavir/ritonavir, each in combination with tenofovir and emtricitabine, for management of antiretroviral-naïve HIV-1-infected patients: 96-week efficacy and safety results of the CASTLE Study. *J Acquir Immune Defic Syndr* 2010;53(3):323-32. CrossRef [↗](#) PubMed [↗](#)
11. Podzamczar D, Ferrer E, Sanchez P, Gatell JM, Crespo M, Fisac C, et al. Less lipoatrophy and better lipid profile with abacavir as compared to stavudine: 96-week results of a randomized study. *J Acquir Immune Defic Syndr* 2007;44(2):139-47. CrossRef [↗](#) PubMed [↗](#)

12. Smith KY, Weinberg WG, DeJesus E, Fischl MA, Liao Q, Ross LL, et al. Fosamprenavir or atazanavir once daily boosted with ritonavir 100 mg, plus tenofovir/emtricitabine, for the initial treatment of HIV infection: 48-week results of ALERT. *AIDS Res Ther* 2008;5(5):1–10. PubMed [↗](#)
13. van Leth F, Phanuphak P, Stroes E, Gazzard B, Cahn P, Raffi F, et al. Nevirapine and efavirenz elicit different changes in lipid profiles in antiretroviral-therapy-naïve patients infected with HIV-1. *PLoS Med* 2004;1(1):e19. Erratum in: *PLoS Med* 2004;1(3):e73. CrossRef [↗](#) PubMed [↗](#)
14. Walmsley S, Avihingsanon A, Slim J, Ward DJ, Ruxrungtham K, Brunetta J, et al. Gemini: a noninferiority study of saquinavir/ritonavir versus lopinavir/ritonavir as initial HIV-1 therapy in adults. *J Acquir Immune Defic Syndr* 2009;50(4):367–74. CrossRef [↗](#) PubMed [↗](#)
15. Dean AG, Sullivan KM, Soe MM. OpenEpi: open source epidemiologic statistics for public health, version 2.3.1. www.OpenEpi.com. Accessed September 4, 2012.
16. Currier JS, Kendall MA, Zackin R, Henry WK, Alston-Smith B, Torriani FJ, et al. Carotid artery intima-media thickness and HIV infection: traditional risk factors overshadow impact of protease inhibitor exposure. *AIDS* 2005;19(9):927–33.
17. Henry K, Melroe H, Huebsch J, Hermundson J, Levine C, Swensen L, et al. Severe premature coronary artery disease with protease inhibitors. *Lancet* 1998;351(9112):1328. CrossRef [↗](#) PubMed [↗](#)
18. Kaplan RC, Kingsley LA, Sharrett AR, Li X, Lazar J, Tien PC, et al. Ten-year predicted coronary heart disease risk in HIV-infected men and women. *Clin Infect Dis* 2007;45(8):1074–81. CrossRef [↗](#) PubMed [↗](#)
19. Lai S, Fishman EK, Lai H, Moore R, Cofrancesco JJ Jr, Pannu H, et al. Long-term cocaine use and antiretroviral therapy are associated with silent coronary artery disease in African Americans with HIV infection who have no cardiovascular symptoms. *Clin Infect Dis* 2008;46:600–10. CrossRef [↗](#) PubMed [↗](#)
20. Savès M, Chène G, Ducimetière P, Lepout C, Le Moal G, Amouyel P, et al. Risk factors for coronary heart disease in patients treated for human immunodeficiency virus infection compared with the general population. *Clin Infect Dis* 2003;37(2):292–8. CrossRef [↗](#) PubMed [↗](#)
21. Triant VA, Lee H, Hadigan C, Grinspoon SK. Increased acute myocardial infarction rates and cardiovascular risk factors among patients with human immunodeficiency virus disease. *J Clin Endocrinol Metab* 2007;92(7):2506–12. CrossRef [↗](#) PubMed [↗](#)
22. Wand H, Calmy A, Carey DL, Samaras K, Carr A, Law MG, et al. Metabolic syndrome, cardiovascular disease and type 2 diabetes mellitus after initiation of antiretroviral therapy in HIV infection. *AIDS* 2007;21(18):2445–53. CrossRef [↗](#) PubMed [↗](#)
23. Carr A, Samaras K, Burton S, Law M, Freund J, Chisholm DJ, et al. A syndrome of peripheral lipodystrophy, hyperlipidaemia and insulin resistance in patients receiving HIV protease inhibitors. *AIDS* 1998;12(7):F51–8. CrossRef [↗](#) PubMed [↗](#)
24. Carr A, Samaras K, Thorisdottir A, Kaufmann GR, Chisholm DJ, Cooper DA. Diagnosis, prediction, and natural course of HIV-1 protease-inhibitor-associated lipodystrophy, hyperlipidaemia, and diabetes mellitus: a cohort study. *Lancet* 1999;353(9170):2093–9. CrossRef [↗](#) PubMed [↗](#)
25. Eron J, Yeni P, Gathe J Jr, Estrada V, DeJesus E, Staszewski S, et al. The KLEAN study of fosamprenavir-ritonavir versus lopinavir-ritonavir, each in combination with abacavir-lamivudine, for initial treatment of HIV infection over 48 weeks: a randomised non-inferiority trial. *Lancet* 2006;368(9534):476–82. CrossRef [↗](#) PubMed [↗](#)
26. Gutiérrez F, Padilla S, Navarro A, Masiá M, Hernández I, Ramos J, et al. Lopinavir plasma concentrations and changes in lipid levels during salvage therapy with lopinavir/ritonavir-containing regimens. *J Acquir Immune Defic Syndr* 2003;33(5):594–600. CrossRef [↗](#) PubMed [↗](#)
27. Johnson M, Grinsztejn B, Rodriguez C, Coco J, DeJesus E, Lazzarin A, et al. Atazanavir plus ritonavir or saquinavir, and lopinavir/ritonavir in patients experiencing multiple virological failures [republished from *AIDS* 2005;19(2):153–62]. *AIDS* 2005;19(7):685–94. CrossRef [↗](#) PubMed [↗](#)
28. Martínez E, Domingo P, Galindo MJ, Milinkovic A, Arroyo JA, Baldovi F, et al. Risk of metabolic abnormalities in patients infected with HIV receiving antiretroviral therapy that contains lopinavir-ritonavir. *Clin Infect Dis* 2004;38(7):1017–23. CrossRef [↗](#) PubMed [↗](#)
29. Mary-Krause M, Cotte L, Simon A, Partisani M, Costagliola D; Clinical Epidemiology Group from the French Hospital Database. Increased risk of myocardial infarction with duration of protease inhibitor therapy in HIV-infected men. *AIDS* 2003;17(17):2479–86. CrossRef [↗](#) PubMed [↗](#)
30. Montes ML, Pulido F, Barros C, Condes E, Rubio R, Cepeda C, et al. Lipid disorders in antiretroviral-naïve patients treated with lopinavir/ritonavir-based HAART: frequency, characterization and risk factors. *J Antimicrob Chemother* 2005;55(5):800–4. CrossRef [↗](#) PubMed [↗](#)

31. Mulligan K, Grunfeld C, Tai VW, Algren H, Pang M, Chernoff DN, et al. Hyperlipidemia and insulin resistance are induced by protease inhibitors independent of changes in body composition in patients with HIV infection. *J Acquir Immune Defic Syndr* 2000;23(1):35–43. PubMed [↗](#)
32. Simpson KN, Jones WJ, Rajagopalan R, Dietz B. Cost effectiveness of lopinavir/ritonavir tablets compared with atazanavir plus ritonavir in antiretroviral-experienced patients in the UK, France, Italy and Spain. *Clin Drug Investig* 2007;27(12):807–17. CrossRef [↗](#) PubMed [↗](#)
33. Riddler SA, Smit E, Cole SR, Li R, Chmiel JS, Dobs A, et al. Impact of HIV infection and HAART on serum lipids in men. *JAMA* 2003;289(22):2978–82. CrossRef [↗](#) PubMed [↗](#)
34. Gordon T, Castelli WP, Hjortland MC, Kannel WB, Dawber TR. High-density lipoprotein as a protective factor against coronary heart disease. The Framingham Study. *Am J Med* 1977;62(5):707–14. CrossRef [↗](#) PubMed [↗](#)
35. Jacobson TA, Case CC, Roberts S, Buckley A, Murtaugh KM, Sung JC, et al. Characteristics of US adults with the metabolic syndrome and therapeutic implications. *Diabetes Obes Metab* 2004;6(5):353–62. CrossRef [↗](#) PubMed [↗](#)
36. Grundy SM, Cleeman JI, Merz CN, Brewer HP Jr, Clark LT, Hunninghake DB, et al. Implications of recent clinical trials for the National Cholesterol Education Program Adult Treatment Panel III Guidelines. *J Am Coll Cardiol* 2004;44(3):720–32. CrossRef [↗](#) PubMed [↗](#)
37. Persell SD, Lloyd-Jones DM, Baker DW. Implications of changing national cholesterol education program goals for the treatment and control of hypercholesterolemia. *J Gen Intern Med* 2006;21(2):171–6. PubMed [↗](#)
38. Data Collection on Adverse Events of Anti-HIV Drugs Study Group; Sabin CA, d'Arminio Monforte A, Friis-Moller N, Weber R, El-Sadr WM, Reiss P, et al. . Changes over time in risk factors for cardiovascular disease and use of lipid-lowering drugs in HIV-infected individuals and impact on myocardial infarction. *Clin Infect Dis* 2008;46(7):1101–10. CrossRef [↗](#) PubMed [↗](#)
39. Reinsch N, Neuhaus K, Esser S, Potthoff A, Hower M, Mostardt S, et al. Are HIV patients undertreated? Cardiovascular risk factors in HIV: results of the HIV-HEART study. *Eur J Prev Cardiol* 2012;19(2):267–74. CrossRef [↗](#) PubMed [↗](#)
40. D:A:D Study Group; Sabin CA, Worm SW, Weber R, Reiss P, El-Sadr W, Dabis F, et al. . Use of nucleoside reverse transcriptase inhibitors and risk of myocardial infarction in HIV-infected patients enrolled in the D:A:D study: a multi-cohort collaboration. *Lancet* 2008;371(9622):1417–26. CrossRef [↗](#) PubMed [↗](#)
41. Sobieszczyk ME, Hoover DR, Anastos K, Mulligan K, Tan T, Shi Q, et al. Prevalence and predictors of metabolic syndrome among HIV-infected and HIV-uninfected women in the Women's Interagency HIV Study. *J Acquir Immune Defic Syndr* 2008;48(3):272–80. CrossRef [↗](#) PubMed [↗](#)

Tables

Table 1. Low-Density Lipoprotein Cholesterol (LDL-C) Goals, by Cardiovascular Risk (CVR) Category, According to the 2001 National Cholesterol Education Program Adult Treatment Panel III (NCEP) Guidelines When Serum Triglycerides Values Are <200 mg/dL^a



CVR Category	LDL-C Level at Which to Initiate Therapeutic Lifestyle Changes, mg/dL	LDL-C Level at Which to Consider Lipid-Lowering Therapy, mg/dL	LDL-C Goal, mg/dL
Low risk, ≤1 major risk factor (10yCVR <10%)	≥160	≥190	<160
Moderate risk, ≥2 major risk factors (10yCVR <10%)	≥130	≥160	<130
Moderately high risk, ≥2 major risk factors (10yCVR 10%–20%)	≥130	≥130	<130
High risk, ≥2 major risk factors or CHD equivalent or (10yCVR >20%)	≥100	≥130	<100

Abbreviations: CHD, coronary heart disease (coronary heart disease or coronary heart disease equivalent per NCEP guidelines); HDL-C, high-density lipoprotein cholesterol.

^a When serum triglycerides are ≥ 200 mg/dL, non-HDL-C goals are used instead of LDL-C goals. Non-HDL-C goals are 30 mg/dL above the LDL-C goals shown in the table.

Table 2. Characteristics of Patients (N = 2,005), by 10-Year Cardiovascular Risk (CVR) Category, HIV Outpatient Study (HOPS), January 2002–September 2009



Characteristic ^a	10-Year CVR Category					P Value ^b
	Overall (N = 2,005)	Low Risk (<10% Risk), ≤ 1 Risk Factor (n = 675)	Moderate Risk (<10% Risk), ≥ 2 Risk Factors (n = 565)	Moderately High Risk (10%–20% Risk), ≥ 2 Risk Factors (n = 365)	High Risk (>20% Risk), ≥ 2 Risk Factors (n = 400)	
Baseline						
Age, y, median	42	38	41	47	48	<.001
Male sex	1,532 (76)	451 (67)	405 (72)	350 (96)	326 (82)	<.001
Race/ethnicity						
White, non-Hispanic	1,048 (52)	304 (45)	276 (49)	252 (69)	216 (54)	<.001
Black, non-Hispanic	657 (33)	251 (37)	199 (35)	73 (20)	134 (34)	
Hispanic	239 (12)	93 (14)	71 (13)	31 (8)	44 (11)	
Other	61 (3)	27 (4)	19 (3)	9 (2)	6 (2)	
Year of HOPS entry						
1998 or earlier	856 (43)	247 (37)	210 (37)	176 (48)	223 (56)	<.001
1999–2005	1,149 (57)	428 (63)	355 (63)	189 (52)	177 (44)	
Private insurance	1,147 (57)	420 (62)	293 (52)	233 (64)	201 (50)	<.001
History of IDU	227 (11)	37 (5)	90 (16)	38 (10)	62 (16)	<.001
Prior AIDS-defining illness	728 (36)	209 (31)	204 (36)	139 (38)	176 (44)	<.001
Cell count/viral load						
Nadir CD4+ T cell count <200 cells/mm ³	1,016 (51)	314 (47)	296 (52)	191 (52)	215 (54)	.06
Nadir CD4+ T cell count, cells/mm ³ , median	197	218	187	190	180	.27
CD4+ T cell count, cells/mm ³ , median	395	396	358	401	415	.002
Peak viral load, copies/mL, median	5,985	7,790	19,712	1,911	2,384	<.001
Viral load, copies/mL, median	419	745	908	200	200	<.001
Viral load <400 copies/mL	988 (49)	305 (45)	244 (43)	211 (58)	228 (57)	<.001

Characteristic ^a	10-Year CVR Category					P Value ^b
	Overall (N = 2,005)	Low Risk (<10% Risk), ≤1 Risk Factor (n = 675)	Moderate Risk (<10% Risk), ≥2 Risk Factors (n = 565)	Moderately High Risk (10%–20% Risk), ≥2 Risk Factors (n = 365)	High Risk (>20% Risk), ≥2 Risk Factors (n = 400)	
Baseline						
Alcohol use, drinks/week						
Missing information	127 (6)	49 (7)	32 (6)	18 (5)	28 (7)	<.001
0	977 (49)	322 (48)	284 (50)	146 (40)	225 (56)	
<7	689 (34)	242 (36)	192 (34)	144 (39)	111 (28)	
7–14	124 (6)	40 (6)	30 (5)	31 (8)	23 (6)	
>14	88 (4)	22 (3)	27 (5)	26 (7)	13 (3)	
BMI, kg/m²						
Missing information	120 (6)	47 (7)	36 (6)	21 (6)	16 (4)	.005
≤25	909 (45)	331 (49)	261 (46)	160 (44)	157 (39)	
>25	976 (49)	297 (44)	268 (47)	184 (50)	227 (57)	
Hypertension	1,036 (52)	112 (17)	356 (63)	230 (63)	338 (85)	<.001
Diabetes mellitus	190 (9)	0	0	0	190 (48)	<.001
History of tobacco use	1,108 (55)	164 (24)	348 (62)	296 (81)	300 (75)	<.001
Blood lipid panel measurement						
Total cholesterol, mg/dL, median	204	197	181	226	238	<.001
LDL cholesterol, mg/dL, median	105	108	94	114	107	<.001
HDL cholesterol, mg/dL, median	37	44	35	35	35	<.001
Non-HDL cholesterol, mg/dL, median ^c	164	151	142	186	199	<.001
Triglycerides, mg/dL, median	163	126	162	200	225	<.001
Metabolic syndrome ^d	516 (26)	85 (13)	157 (28)	103 (28)	171 (43)	<.001
Antiretroviral therapy						
Efavirenz	452 (23)	157 (23)	136 (24)	66 (18)	93 (23)	.14
Nevirapine	213 (11)	65 (10)	63 (11)	38 (10)	47 (12)	.70
Protease inhibitor, unboosted	222 (11)	62 (9)	72 (13)	41 (11)	47 (12)	.23
Protease inhibitor, boosted	626 (31)	205 (30)	171 (30)	126 (35)	124 (31)	.51
Zidovudine or Stavudine	775 (39)	233 (35)	230 (41)	143 (39)	169 (42)	.043
Didanosine	273 (14)	78 (12)	72 (13)	62 (17)	61 (15)	.07
Abacavir	443 (22)	139 (21)	132 (23)	81 (22)	91 (23)	.68

Characteristic ^a	10-Year CVR Category					P Value ^b
	Overall (N = 2,005)	Low Risk (<10% Risk), ≤1 Risk Factor (n = 675)	Moderate Risk (<10% Risk), ≥2 Risk Factors (n = 565)	Moderately High Risk (10%–20% Risk), ≥2 Risk Factors (n = 365)	High Risk (>20% Risk), ≥2 Risk Factors (n = 400)	
Baseline						
Tenofovir	561 (28)	174 (26)	181 (32)	92 (25)	114 (29)	.055
Lipid-lowering agents						
Statin drug/ezetimibe	245 (12)	42 (6)	31 (5)	49 (13)	123 (31)	<.001
Fibrate	116 (6)	13 (2)	16 (3)	26 (7)	61 (15)	<.001
Fish oil	17 (1)	4 (1)	3 (1)	2 (1)	8 (2)	.09
Follow-Up						
Follow-up after baseline, y, median	5.5	5.5	5.5	5.8	5.1	.32
Lipid-lowering agents						
Statin drug/ezetimibe	562 (28)	119 (18)	109 (19)	125 (34)	209 (52)	<.001
Fibrate	235 (12)	39 (6)	53 (9)	54 (15)	89 (22)	<.001
Fish oil	139 (7)	25 (4)	42 (7)	33 (9)	39 (10)	<.001
ARV regimen changes^e	1,353 (67)	442 (65)	380 (67)	260 (71)	271 (68)	.30
Antihypertensives	633 (32)	52 (8)	212 (38)	136 (37)	233 (58)	<.001
Incident CVD	148 (7)	14 (2)	27 (5)	45 (12)	62 (16)	<.001

Abbreviations: IDU, intravenous drug use; AIDS, acquired immunodeficiency syndrome; BMI, body mass index; LDL, low-density lipoprotein; HDL, high-density lipoprotein; ARV, antiretroviral.

^a Values presented as n (%) unless otherwise indicated.

^b χ^2 test for binary variables or Kruskal-Wallis test for continuous variables.

^c Non-HDL cholesterol equals total cholesterol minus HDL cholesterol.

^d Defined as the presence of any 3 of the following 5 risk factors: hypertension (≥ 130 mm Hg over ≥ 85 mm Hg), elevated triglycerides (≥ 150 mg/dL), low HDL cholesterol (< 40 mg/dL for men, < 50 mg/dL for women), fasting glucose ≥ 110 mg/dL, abdominal obesity (BMI ≥ 28.9 kg/m² for men and ≥ 24.9 kg/m² for women).

^e Treatment of either high LDL-cholesterol/non-HDL-cholesterol or hypertriglyceridemia can include the following ARV changes: Zidovudine/Stavudine to Abacavir or Tenofovir; Abacavir to Tenofovir; boosted protease inhibitor to unboosted protease inhibitor; any protease inhibitor to Efavirenz or Nevirapine; Efavirenz to Nevirapine; boosted Indinavir or Lopinavir to boosted Darunavir, Atazanavir, fos-Amprnavir, or Saquinavir/Fortovase.

Table 3. Treatment of Patients (N = 2,005) for High LDL-C/Non-HDL-C, Low HDL-C, Hypertriglyceridemia, and Hypertension, by 10-Year Cardiovascular Risk (CVR) Category, HIV Outpatient Study, January 2002–September 2009



Condition/CVR Category ^a	% (n/N) With Condition	% (n/N) Treated Per Guidelines ^b	% (n/N) Treated Who Met NCEP Goals ^c
High LDL-C/non-HDL-C	$P < .001^d$	$P = .68$	$P < .001$
Low risk	15.9 (107/675)	81.3 (87/107)	81.6 (71/87)
Moderate risk	31.5 (178/565)	86.5 (154/178)	62.3 (96/154)
Moderately high risk	62.2 (227/365)	83.3 (189/227)	59.3 (112/189)
High risk	80.0 (320/400)	84.1 (269/320)	33.5 (90/269)

Condition/CVR Category ^a	% (n/N) With Condition	% (n/N) Treated Per Guidelines ^b	% (n/N) Treated Who Met NCEP Goals ^c
Low HDL-C	<i>P</i> < .001	<i>P</i> < .001	<i>P</i> = .42
Low risk	41.6 (281/675)	2.9 (8/281)	50.0 (4/8)
Moderate risk	78.4 (443/565)	1.6 (7/443)	14.3 (1/7)
Moderately high risk	67.7 (247/365)	8.1 (20/247)	20.0 (4/20)
High risk	73.0 (292/400)	11.3 (33/292)	27.3 (9/33)
Hypertriglyceridemia	<i>P</i> < .001	<i>P</i> = .03	<i>P</i> = .84
Low risk	2.4 (16/675)	56.3 (9/16)	88.9 (8/9)
Moderate risk	5.8 (33/565)	90.9 (30/33)	80.0 (24/30)
Moderately high risk	12.3 (45/365)	82.2 (37/45)	78.4 (29/37)
High risk	18.3 (73/400)	87.7 (64/73)	76.6 (49/64)
Hypertension	<i>P</i> < .001	<i>P</i> < .001	<i>P</i> < .001
Low risk	16.6 (112/675)	46.4 (52/112)	67.3 (35/52)
Moderate risk	63.0 (356/565)	59.6 (212/356)	66.0 (140/212)
Moderately high risk	63.0 (230/365)	59.1 (136/230)	64.0 (87/136)
High risk	84.5 (338/400)	68.9 (233/338)	46.4 (108/233)

Abbreviations: LDL-C, low-density lipoprotein cholesterol; HDL-C, high-density lipoprotein cholesterol; NCEP, National Cholesterol Education Program Adult Treatment Panel III.

^a See text for definitions of 10-year cardiovascular disease risk categories and definitions of high LDL-C/non-HDL-C, low HDL-C, hypertriglyceridemia, and hypertension.

^b Eligible patients treated per modified NCEP guidelines within 2 years of baseline date. Treatment of either high LDL-C/non-HDL-C or hypertriglyceridemia can include the following ARV changes: Zidovudine/Stavudine to Abacavir or Tenofovir; Abacavir to Tenofovir; boosted protease inhibitor to unboosted protease inhibitor; any protease inhibitor to Efavirenz or Nevirapine; Efavirenz to Nevirapine; boosted Indinavir or Lopinavir to boosted Darunavir, Atazanavir, fos-Amprenavir, or Saquinavir/Fortovase.

^c Treated patients who met modified NCEP goals after starting treatment and before the end of observation in the study.

^d Likelihood ratio χ^2 test.


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