Research Article

Serum Lipid Profile and Risk of Prostate Cancer Recurrence: Results from the SEARCH Database

Emma H. Allott1,2,3, Lauren E. Howard1,3,4, Matthew R. Cooperberg5, Christopher J. Kane6, William J. Aronson7,8, Martha K. Terris9,10, Christopher L. Amling11, and Stephen J. Freedland1,3,12

Abstract

**Background:** Evidence for an association between total cholesterol, low- and high-density lipoproteins (LDL and HDL, respectively), triglycerides, and prostate cancer is conflicting. Given that prostate cancer and dyslipidemia affect large proportions of Western society, understanding these associations has public health importance.

**Methods:** We conducted a retrospective cohort analysis of 843 radical prostatectomy (RP) patients who never used statins before surgery within the Shared Equal Access Regional Cancer Hospital (SEARCH) database. Multivariable Cox proportional hazards analysis was used to investigate the association between cholesterol, LDL, HDL, and triglycerides and biochemical recurrence risk. In secondary analysis, we explored these associations in patients with dyslipidemia, defined using National Cholesterol Education Program guidelines.

**Results:** Elevated serum triglycerides were associated with increased risk of prostate cancer recurrence ([HR per 10 mg/dl, 1.03; 95% confidence interval (CI), 1.01–1.05]) but associations between total cholesterol, LDL, and HDL, and recurrence risk were null. However, among men with dyslipidemia, each 10 mg/dl increase in cholesterol and HDL was associated with a 9% increased recurrence risk (HR, 1.09; 95% CI, 1.01–1.17) and 39% reduced recurrence risk (HR, 0.61; 95% CI, 0.41–0.91), respectively.

**Conclusions:** Elevated serum triglycerides were associated with increased risk of prostate cancer recurrence. Cholesterol, LDL, or HDL were not associated with recurrence risk among all men. However, among men with dyslipidemia, elevated cholesterol and HDL levels were associated with increased and decreased risk of recurrence, respectively.

**Impact:** These findings, coupled with evidence that statin use is associated with reduced recurrence risk, suggest that lipid levels should be explored as a modifiable risk factor for prostate cancer recurrence. *Cancer Epidemiol Biomarkers Prev; 23(11); 2349–56. ©2014 AACR.*

Introduction

Prostate cancer is the most commonly diagnosed noncutaneous cancer in U.S. males and the second most common cause of cancer-related deaths (1). Approximately two thirds of the U.S. population are overweight or obese (2), a metabolic disorder associated with increased risk of aggressive prostate cancer and prostate cancer mortality (3). Hypercholesterolemia, a condition strongly related to obesity, currently affects approximately 20% of the U.S. adult population (4). Cholesterol is hypothesized to contribute to prostate cancer progression due to its established role as a signaling molecule in prostate growth and differentiation (5), in addition to evidence from laboratory studies suggesting that cholesterol may drive prostate cancer growth via multiple biologic mechanisms, including Akt signaling (6) and de novo steroidogenesis (7). Given the high prevalence of hypercholesterolemia in Western society, understanding the potential association between this modifiable risk factor and prostate cancer progression is of great public health importance.
Although epidemiologic evidence does not support an association between serum cholesterol levels and risk of total prostate cancer (8, 9), there is a suggestion that elevated cholesterol may be associated with increased risk of aggressive disease (8, 10–12), although not all studies have reported this finding (9, 13–15). There is mixed evidence for an association between serum cholesterol levels and risk of prostate cancer progression, with some studies reporting positive associations between elevated cholesterol and risk of prostate cancer recurrence (16) and mortality (17, 18), whereas another study reported no association with risk of prostate cancer mortality (19). Fewer studies examined the association between cholesterol subfractions—low- and high-density lipoprotein (LDL and HDL, respectively)—and prostate cancer. Although there is some evidence that elevated LDL (12, 20) and low HDL (10, 21) are associated with increased risk of aggressive prostate cancer, not all studies reported these findings (13, 15, 22), and the association between cholesterol subfractions and risk of prostate cancer recurrence has not been widely studied. Finally, evidence for an association between serum triglycerides and prostate cancer recurrence is mixed (14, 23). Thus, the impact of dyslipidemia on risk of prostate cancer recurrence is not well understood.

The aim of this study was to examine the association between serum lipid levels and risk of biochemical recurrence in a retrospective cohort of radical prostatectomy (RP) patients who never used statins before surgery, from the Shared Equal Access Regional Cancer Hospital (SEARCH) database. We hypothesized that elevated serum cholesterol, triglycerides, and LDL would be associated with increased risk of prostate cancer recurrence, with a protective association between elevated HDL and risk of recurrence.

Materials and Methods

Study sample
After obtaining Institutional Review Board approval from each institution, data from patients undergoing RP (n = 2,542) between 1999 and 2013 at six VA Medical Centers (West Los Angeles, CA; Palo Alto, CA; San Diego, CA; Durham, NC; Asheville, NC; and Augusta, GA) were combined into the SEARCH database (24). SEARCH does not include patients treated with preoperative androgen deprivation or radiation therapy. Given that preoperative serum cholesterol level was our primary exposure of interest, patients who used statins before surgery were excluded (n = 1,135). We also excluded patients with missing data for serum lipid levels (n = 482), preoperative PSA (n = 9), body mass index (BMI; n = 25), pathologic Gleason score (n = 11), pathologic features (n = 29), and PSA follow-up (n = 8), resulting in a study sample of 843 men.

Exposure assessment and definitions
Fasting total serum cholesterol, LDL, HDL, and triglyceride levels measured within the year before RP were abstracted from VA-computerized medical records. Recommended cutoffs for normal versus abnormal serum levels (all in mg/dl) of total cholesterol (<200 vs. ≥200), LDL (<130 vs. ≥130), HDL (≥40 vs. <40), and triglycerides (<150 vs. ≥150) were selected according to National Cholesterol Education Program (NCEP)-Adult Treatment Panel (ATP) III guidelines (25). NCEP-ATPIII borderline and high lipid categories were combined to have adequate numbers of patients with abnormal lipid levels for the analysis. On the basis of the NCEP-ATP III guidelines, we defined normal versus abnormal categories for each individual lipid independently of the others. For example, a patient could be included in the abnormal category for cholesterol but in the normal category for triglycerides if his cholesterol level was ≥200 mg/dl and his triglyceride level was <150 mg/dl.

Follow-up
Follow-up protocols were at the discretion of the treating physicians. Biochemical recurrence was defined as a single PSA >0.2 ng/mL, two consecutive concentrations at 0.2 ng/mL, or secondary treatment for detectable postoperative PSA. Men receiving adjuvant therapy after surgery for an undetectable PSA were considered nonrecurrent at the time of adjuvant therapy, and their follow-up was censored at that point.

Statistical analysis
Given that our primary hypothesis was to test the association between total serum cholesterol levels and risk of biochemical recurrence, analysis of cholesterol was considered primary, whereas analysis of LDL, HDL, and triglyceride levels was considered secondary. Differences in demographic, clinical, and pathologic factors between patients with normal versus abnormal total serum cholesterol (<200 vs. ≥200 mg/dl) were examined using t tests and χ² tests for continuous and categorical variables, respectively, and rank-sum tests for continuous variables not normally distributed.

Time from RP to biochemical recurrence was compared between normal versus abnormal serum cholesterol categories using Kaplan–Meier plots and the log-rank test. Cox proportional hazards analysis was used to test whether serum cholesterol levels (abnormal vs. normal, as well as continuous) independently predicted time to recurrence. Continuous lipid levels were presented in 10 mg/dl increments to facilitate interpretation of the HRs. The proportionality assumption was tested by examining the Schoenfeld residuals. Cox models were adjusted for age at surgery (continuous), race (black vs. non-black), preoperative PSA (continuous; log-transformed), year of surgery (continuous), BMI (continuous; log-transformed), pathologic Gleason score (2–6, 7 (3 + 4), 7 (4 + 3)–10), positive surgical margins (yes vs. no), extracapsular extension (yes vs. no), seminal vesicle invasion (yes vs. no), and surgical center. Models were also adjusted for post-RP statin use which was treated as a time-dependent variable to account for varying start dates and duration of post-RP statin use during follow-up, based upon our
of the entire cohort of 843 men who never used statins before RP, 325 (39%) patients had abnormal preoperative cholesterol levels (≥200 mg/dl), as defined using NCEP-ATP III guidelines (Table 1; ref. 25). As anticipated, men with normal preoperative cholesterol levels (<200 mg/dl) were significantly less likely to use statins after RP, relative to men with abnormal preoperative cholesterol (P < 0.001). However, there were no significant differences in age at surgery, race, preoperative PSA, diabetes status, BMI, or any pathologic features by cholesterol status (Table 1). With the exception of LDL and triglycerides, lipid levels were moderately intercorrelated, with the strongest correlation between cholesterol and LDL (Pearson correlation coefficient = 0.83). LDL and triglyceride levels were positively correlated with BMI, whereas HDL was negatively correlated with BMI (Supplementary Table S2).

Total serum cholesterol and risk of recurrence

A total of 293 (35%) men experienced biochemical recurrence. Median follow-up among men who did not recur was 74.3 months (Q1–Q3: 41.5–102.8). Patients with normal cholesterol levels were more recently treated than patients with abnormal cholesterol (2004 vs. 2005; P < 0.001; Table 1), but there was no significant difference in follow-up between groups (P = 0.5). Kaplan–Meier plots revealed no significant effect of cholesterol on risk of recurrence (log-rank P = 0.334; data not shown). Total serum cholesterol was not associated with risk of recurrence, either as a continuous or categorical variable, on multivariable analysis (both P ≥ 0.4; Table 2). Competing risk analysis did not alter our results (Supplementary Table S3).

Serum lipids and risk of recurrence

Similar to our null findings for total cholesterol, neither LDL nor HDL were significantly related to risk of prostate cancer recurrence, either as continuous or categorical variables, on either univariable (log-rank P = 0.824 and P = 0.339, respectively) or multivariable analysis (all P ≥ 0.339; Table 2). However, elevated triglycerides were associated with 35% increased risk of recurrence on multivariable analysis [abnormal vs. normal; HR, 1.35; 95% confidence interval (CI), 1.05–1.74; Table 2]. Furthermore, there was a significant 2% increased risk of recurrence for every 10 mg/dl increase in triglyceride level when treated as a continuous variable on both univariable (HRper 10 mg/dl, 1.02; 95% CI, 1.02–1.03) and multivariable analyses (HRper 10 mg/dl, 1.02; 95% CI, 1.02–1.04; Table 2). Given that abnormal serum triglycerides form part of the NCEP-ATPIII diagnostic panel for diabetes (25), we restricted our analysis to men without diabetes and found a similar association (abnormal vs. normal; HR, 1.46; 95% CI, 1.10–1.93). Given the strong correlation between LDL and cholesterol, we did not mutually adjust for LDL and cholesterol, but mutually adjusting for other combinations of lipids did not alter our results (data not shown). We found no significant interaction between BMI and any...
Treating death as a competing risk did not materially affect the associations between HDL, LDL, and triglycerides and recurrence (Supplementary Table S3). Given that preclinical disease may affect serum lipid levels, we repeated our analysis after excluding the first year of follow-up and found that none of the HRs were appreciably altered (data not shown).

Table 1. Demographic, clinical, and pathologic characteristics of patients by serum cholesterol status

<table>
<thead>
<tr>
<th></th>
<th>Cholesterol &lt; 200 mg/dL (n = 518; 61%)</th>
<th>Cholesterol &gt; 200 mg/dL (n = 325; 39%)</th>
<th>P</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age, mean ± SD</td>
<td>60.4 ± 6.3</td>
<td>60.6 ± 6.4</td>
<td>0.664a</td>
</tr>
<tr>
<td>Race, n (%)</td>
<td>Black 222 (43)</td>
<td>121 (37)</td>
<td>0.106b</td>
</tr>
<tr>
<td></td>
<td>Non-black 296 (57)</td>
<td>204 (63)</td>
<td></td>
</tr>
<tr>
<td>Follow-up, median (Q1–Q3)</td>
<td>74.7 (41.3–99.0)</td>
<td>73.4 (41.5–108.1)</td>
<td>0.501c</td>
</tr>
<tr>
<td>PSA, median (Q1–Q3)</td>
<td>6.3 (4.8–9.4)</td>
<td>7.2 (5.0–10.5)</td>
<td>0.071c</td>
</tr>
<tr>
<td>Statin use, n (%)</td>
<td>Never 308 (59)</td>
<td>108 (33)</td>
<td></td>
</tr>
<tr>
<td></td>
<td>Started after RP 210 (41)</td>
<td>217 (67)</td>
<td></td>
</tr>
<tr>
<td>BMI, median (Q1–Q3)</td>
<td>27.4 (24.7–30.5)</td>
<td>27.5 (24.9–30.6)</td>
<td>0.936d</td>
</tr>
<tr>
<td>Diabetes, n (%)</td>
<td>84 (21)</td>
<td>48 (20)</td>
<td>0.687c</td>
</tr>
<tr>
<td>LDL, mean (SD)</td>
<td>101.9 (21.6)</td>
<td>141.9 (25.4)</td>
<td>&lt;0.0001</td>
</tr>
<tr>
<td>HDL, mean (SD)</td>
<td>43.8 (13.7)</td>
<td>51.2 (20.3)</td>
<td>&lt;0.0001</td>
</tr>
<tr>
<td>Triglycerides, mean (SD)</td>
<td>126.0 (83.2)</td>
<td>150.0 (83.9)</td>
<td>0.0001</td>
</tr>
<tr>
<td>Pathologic Gleason score, n (%)</td>
<td>2–6 177 (34)</td>
<td>125 (39)</td>
<td>0.276h</td>
</tr>
<tr>
<td></td>
<td>7 (3+4) 214 (41)</td>
<td>134 (41)</td>
<td></td>
</tr>
<tr>
<td></td>
<td>7 (4+) 127 (25)</td>
<td>66 (20)</td>
<td></td>
</tr>
<tr>
<td>Positive margins, n (%)</td>
<td>228 (44)</td>
<td>141 (43)</td>
<td>0.857c</td>
</tr>
<tr>
<td>Extracapsular extension, n (%)</td>
<td>87 (17)</td>
<td>66 (20)</td>
<td>0.198c</td>
</tr>
<tr>
<td>Seminal vesicle invasion, n (%)</td>
<td>44 (8)</td>
<td>30 (9)</td>
<td>0.713c</td>
</tr>
<tr>
<td>Positive lymph nodes, n (%)</td>
<td>12 (2)</td>
<td>9 (3)</td>
<td>0.919f</td>
</tr>
</tbody>
</table>

NOTE: P values calculated by a t test, b z test, or c Wilcoxon rank-sum test. Abbreviations: Q1 = 25th percentile; Q3 = 75th percentile.

Table 2. HRs for serum lipid levels predicting risk of biochemical recurrence after radical prostatectomy

<table>
<thead>
<tr>
<th></th>
<th>Total cholesterol</th>
<th>LDL</th>
<th>HDL</th>
<th>Triglycerides</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>&lt; 200 mg/dL</td>
<td>≥ 130 mg/dL</td>
<td>≥ 40 mg/dL</td>
<td>&lt; 150 mg/dL</td>
</tr>
<tr>
<td></td>
<td>Unadjusted</td>
<td>Adjustedb</td>
<td>Unadjusted</td>
<td>Adjustedb</td>
</tr>
<tr>
<td></td>
<td>1.13 (0.90–1.43), P = 0.294</td>
<td>1.09 (0.85–1.39), P = 0.483</td>
<td>1.03 (0.81–1.31), P = 0.824</td>
<td>1.11 (0.87–1.41), P = 0.402</td>
</tr>
</tbody>
</table>

Note: Cells display HR (95% CI), P value.
bHRs are for every 10 mg/dL increase.
HRs are adjusted for age, race, preoperative PSA, year of surgery, BMI, surgical center, postoperative statin use, pathologic Gleason score, prostate weight, positive surgical margins, extracapsular extension, and seminal vesicle invasion.
Risk of recurrence among men with abnormal lipid levels

Given these null overall findings for cholesterol, LDL, and HDL, we further explored the association between serum lipid levels and risk of prostate cancer recurrence by modeling these relationships using LOWESS plots (Fig. 1). Because these plots suggested a possible relationship between lipid levels and risk of recurrence when lipid values were in the abnormal range, we performed secondary analyses restricted to patients with abnormal lipid levels. On multivariable analysis of men with abnormal lipid levels, there was a significant association between higher cholesterol levels and risk of recurrence, with each 10 mg/dl increase in cholesterol above 200 mg/dl associated with 9% increased risk of recurrence (HR\textsubscript{per 10 mg/dl} 1.09; 95% CI, 1.01–1.17; Table 3). Furthermore, increasing HDL within the abnormal range was significantly protective, with each 10 mg/dL increase associated with 39% reduced risk of recurrence (HR\textsubscript{per 10 mg/dL} 0.61; 95% CI, 0.41–0.91), while the association between LDL and risk of recurrence was null (HR\textsubscript{per 10 mg/dL} 1.05; 95% CI, 0.94–1.17). In line with our analysis among all men, elevated triglycerides within the abnormal range remained associated with increased risk of recurrence (HR\textsubscript{per 10 mg/dL} 1.03; 95% CI, 1.01–1.05). There were no significant interactions between abnormal levels of any lipid and postoperative statin use in predicting risk of recurrence (data not shown). Treating death as a competing risk did not materially affect the associations between HDL, LDL, and triglycerides and recurrence (Supplementary Table S4). Among men with normal lipid levels, total cholesterol, LDL, HDL and triglycerides were not associated with risk of recurrence (data not shown).

Discussion

Although obesity is an established risk factor for aggressive prostate cancer and prostate cancer mortality (3), the mechanisms contributing to this obesity–prostate cancer link are not well understood. On the basis of biologic evidence supporting an important role for cholesterol in
prostate cancer, we hypothesized that serum lipid levels, a potentially modifiable factor, may influence risk of prostate cancer recurrence. In contrast to this hypothesis, we found a null association between total cholesterol, LDL and HDL, and risk of recurrence. However, each 10 mg/dl increase in serum triglyceride levels was associated with 2% increased risk of prostate cancer recurrence, with elevated serum triglyceride levels, a potentially modifiable factor, may influence risk of recurrence. These findings, which require confirmation in other studies, may highlight the importance of controlling total cholesterol and HDL levels in patients with prostate cancer with dyslipidemia, not only for cardiovascular disease risk reduction but also for potential advantage to prostate cancer progression. Cholesterol promotes prostate cancer cell line growth both in vitro and in xenograft models via lipid raft-mediated Akt signaling (6). Moreover, reduction of serum cholesterol, the preceptor for sex steroid synthesis, has been demonstrated to lower tumor androgen levels and slow tumor growth in xenograft models of human prostate cancer (7). However, epidemiologic evidence for an association between cholesterol and prostate cancer progression is mixed. One retrospective study reported a protective association between low cholesterol and risk of recurrence in radiation-treated patients with prostate cancer (16), and two large studies found that elevated cholesterol levels were associated with increased risk of prostate cancer mortality (17, 18). In contrast, another study reported a null association between cholesterol and prostate cancer mortality, although this study was conducted in a predominantly Asian population (19). Of note, in contrast to total cholesterol, we found no association between LDL and recurrence even among men with abnormal LDL levels. Although LDL levels are important for estimating cardiovascular disease risk, it is unknown which cholesterol subfractions are most important for tumor growth, and this requires further study. Although the primary mechanism by which HDL reduces cardiovascular disease risk is via reverse cholesterol transport (28), it is not known whether this same mechanism may affect prostate cancer growth. HDL has also been demonstrated to have antiinflammatory, antiinflammatory, and antioxidant properties (29), which may slow prostate cancer growth and progression, although epidemiologic evidence has been inconclusive (22). Finally, high triglycerides have been associated with elevated levels of reactive oxygen species and oxidative stress, in addition to development of insulin resistance, all of which have been associated with prostate tumorigenesis (30). Thus, the possible association between dyslipidemia and increased risk of prostate cancer recurrence is supported by these multiple biologic pathways by which serum lipids may affect prostate cancer growth and progression.

Our study has several limitations which should be considered. First, all serum lipid measurements were obtained within the year before RP and thus may potentially be affected by the presence of preclinical disease (31). To address this, we explored the impact of excluding recurrences during the first year of follow-up, and found

Table 3. HRs for abnormal lipid levels predicting risk of biochemical recurrence after radical prostatectomy

<table>
<thead>
<tr>
<th>Total cholesterol</th>
<th>Continuous; ≥ 200 mg/dL</th>
</tr>
</thead>
<tbody>
<tr>
<td>Unadjusted</td>
<td>1.06 (0.97–1.15), P = 0.206</td>
</tr>
<tr>
<td>Adjusteda</td>
<td>1.09 (1.01–1.19), P = 0.027</td>
</tr>
<tr>
<td>LDL</td>
<td>Continuous; ≥ 130 mg/dL</td>
</tr>
<tr>
<td>Unadjusted</td>
<td>1.01 (0.90–1.13), P = 0.860</td>
</tr>
<tr>
<td>Adjusteda</td>
<td>1.05 (0.94–1.17), P = 0.356</td>
</tr>
<tr>
<td>HDL</td>
<td>Continuous; &lt; 40 mg/dL</td>
</tr>
<tr>
<td>Unadjusted</td>
<td>0.73 (0.51–1.04), P = 0.085</td>
</tr>
<tr>
<td>Adjusteda</td>
<td>0.61 (0.41–0.91), P = 0.016</td>
</tr>
<tr>
<td>Triglycerides</td>
<td>Continuous; ≥ 150 mg/dL</td>
</tr>
<tr>
<td>Unadjusted</td>
<td>1.04 (1.02–1.06), P &lt; 0.001</td>
</tr>
<tr>
<td>Adjusteda</td>
<td>1.03 (1.01–1.05), P = 0.004</td>
</tr>
</tbody>
</table>

NOTE: Cells display HR (95% CI), P value.
All HRs are for every 10 mg/dl increase.

aHRs are adjusted for age, race, preoperative PSA, year of surgery, BMI, surgical center, postoperative statin use, pathologic Gleason score, positive surgical margins, extracapsular extension, and seminal vesicle invasion.
that this did not alter our results. In addition, bias due to reverse causation is less likely in screened populations such as ours where prostate cancer is diagnosed early in the natural history of the disease. Second, we lacked sufficient numbers to explore the impact of abnormal levels of all four lipids simultaneously on prostate cancer recurrence. Third, we did not have access to hypertension data and therefore could not assess the association between metabolic syndrome and risk of recurrence. Neither could we assess the impact of abnormal lipid levels within the context of metabolic syndrome. However, given the use of different definitions of the metabolic syndrome across epidemiologic studies (30), in addition to recent lack of certainty regarding its pathogenesis (32), it may be equally or more informative to estimate the effect of individual components of the metabolic syndrome on risk of prostate cancer recurrence. Finally, an important limitation of all observational biomarker studies is that causality cannot be inferred from these associations. These limitations are balanced by an important strength of this study. We excluded all men who were taking statins at the time of serum lipid measurement. Although exclusion of preoperative statin users limits the generalizability of our findings to men who do not use statins, we believe that this approach strengthens our exposure assessment as the preoperative lipid levels of statin users may not reflect the environment that their tumor developed in. Furthermore, we adjusted our multivariable models for post-RP statin use as a time-dependent variable. Thus, our serum lipid level measurements were obtained in the absence of any cholesterol-lowering medications and we were able to assess the association between serum lipid levels and risk of recurrence separately from statin use. Although we found no evidence for interaction between postoperative statin use and abnormal lipid levels in predicting risk of recurrence, future studies should assess whether beginning statin therapy after prostate cancer treatment could attenuate associations between abnormal lipid levels and risk of recurrence.

In summary, we found no strong evidence to support a link between serum lipid levels and risk of prostate cancer recurrence across the entire sample. However, among men with dyslipidemia, our findings suggest that normalization of serum lipid levels may be beneficial not only for cardiovascular disease prevention but also for prostate cancer recurrence risk reduction, though these results require confirmation in future studies. Although it cannot be determined from this study if these observed associations are causal, given the biologic evidence supporting an important role of cholesterol in prostate cancer growth in addition to epidemiologic data demonstrating that statin use is associated with reduced risk of recurrence, we believe that serum lipid levels should be explored further as a risk factor for prostate cancer recurrence. Although the association between obesity and increased risk of prostate cancer recurrence is likely to be multifactorial (3), these findings suggest that dyslipidemia may be one of the mechanisms underlying this association. Given that 45% of deaths worldwide can be attributed to cardiovascular disease and cancer (15), with prostate cancer the second most common cause of male cancer deaths (1), understanding the role of dyslipidemia as a shared, modifiable risk factor for both of these common causes of mortality is of great importance.

Disclosure of Potential Conflicts of Interest
No potential conflicts of interest were disclosed.

Authors’ Contributions
Conception and design: E.H. Allott, W.J. Aronson, C.L. Amling, S.J. Freedland
Acquisition of data (provided animals, acquired and managed patients, provided facilities, etc.): C.J. Kane, M.K. Terris, C.L. Amling, S.J. Freedland
Analysis and interpretation of data (e.g., statistical analysis, biostatistics, computational analysis): E.H. Allott, L.E. Howard, W.J. Aronson, C.L. Amling, S.J. Freedland
Writing, review, and/or revision of the manuscript: E.H. Allott, L.E. Howard, M.R. Cooperberg, C.J. Kane, W.J. Aronson, C.L. Amling, S.J. Freedland
Administrative, technical, or material support (i.e., reporting or organizing data, constructing databases): S.J. Freedland
Study supervision: M.K. Terris, S.J. Freedland

Grant Support
This study was supported by grants from the NCI (5R25-CA126938-03; to E.H. Allott) and NIH (1R01-CA131235-01A1 and 1K24-CA160653; to S.J. Freedland).

The costs of publication of this article were defrayed in part by the payment of page charges. This article must therefore be hereby marked advertisement in accordance with 18 U.S.C. Section 1734 solely to indicate this fact.

Received April 24, 2014; revised July 29, 2014; accepted August 6, 2014; published OnlineFirst October 10, 2014.

References


Serum Lipid Profile and Risk of Prostate Cancer Recurrence: Results from the SEARCH Database

Emma H. Allott, Lauren E. Howard, Matthew R. Cooperberg, et al.

Cancer Epidemiol Biomarkers Prev 2014;23:2349-2356. Published OnlineFirst October 10, 2014.