Kidney-Heart Interactions: Epidemiology, Pathogenesis, and Treatment

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Heart disease accounts for approximately half of the deaths of patients with ESRD (1–3). In the past 5 yr, there has been increasing recognition of both coronary artery disease (CAD) and left ventricular hypertrophy (LVH) in ESRD patients, as these are the two typical presentations of heart disease in ESRD patients. It is also clear that many patients with chronic kidney disease (CKD) and a GFR of <60 ml/min are at risk for heart disease; many of these patients succumb to heart disease before reaching dialysis (4–6). This review includes a brief overview of the problem, a discussion of CAD and LVH, and an examination of the benefits of reducing BP and proteinuria on both the heart and the kidney and concludes with a brief section on treatment options for individuals with this disorder.

Overview of the Problem of Cardiovascular Disease in Patients with Kidney Disease

A growing awareness of heart disease in individuals with kidney disease as a major public health concern has increased sharply because of the revelation that there are millions of Americans with reduced kidney function (2). This fact, coupled with the understanding that many individuals with CKD do not reach dialysis because they die of heart disease (6), has expanded the concern about heart disease in both patients with CKD and patients with ESRD. Of interest is that whereas many superb prospective, randomized clinical trials have defined the scope of the appropriate therapy for heart disease in patients with normal kidney function, relatively few trials have addressed the issue in patients with kidney disease. Thus, the literature on cardiovascular disease (CVD) has focused on individuals without kidney disease, and guidelines regarding the management of heart disease in patients with ESRD in CKD, therefore, are largely opinion based (7).

The two clinical presentations of heart disease in patients with kidney disease are atherosclerotic vascular disease (particularly CAD) and LVH. Atherosclerosis has been known to be a particular problem in patients with ESRD for >25 yr since Lindner et al. (8) hypothesized that there was accelerated atherogenesis in patients with ESRD. In the Lindner study, 60% of a cohort who underwent dialysis for 6.5 yr died and 14 of these deaths were attributed to complications of atherosclerosis. Subsequent postmortem and angiographic studies confirmed that the prevalence of atherosclerosis is increased dramatically in patients who are on dialysis when compared with age-matched individuals without kidney impairment (9–13). Of note, this increase in atherosclerosis and, in particular, CAD is progressive over a range of reduced GFR. This has been shown by Anavekar et al. (5) and by Beddhu et al. (14). The major increase in risk for heart disease and death occurs at a GFR of <50 to 60 ml/min (5,14) (Figure 1). A major problem with the issue of coronary disease is that in many patients, there is a high burden of coronary disease despite that the patient is asymptomatic. In a recent study by deFilippi et al. (15), 44% of a large cohort of asymptomatic hemodialysis patients had significant coronary disease. This was confirmed recently by Ohtake et al. (16), who showed that coronary disease (defined as >50% stenosis in the coronary artery) was present in as many as 53% of a cohort of asymptomatic dialysis patients.

A second presentation of heart disease in this patient population is LVH. It is particularly important to understand this disorder in dialysis patients because as many as 80% of an incident dialysis population will have LVH as they begin dialysis (17). Individuals with LVH have both eccentric and concentric hypertrophy (18,19). Eccentric hypertrophy results from volume overload leading to cardiac myocyte dropout; LVH is characterized by a myocyte to arteriolar capillary mismatch. Concentric hypertrophy typically is the result of hypertension and increased afterload and is exacerbated by anemia, hyperparathyroidism, and high angiotensin II concentrations. Eccentric and concentric hypertrophy are relatively equivalent in prevalence in dialysis patients.

The dominant LV physiology that accompanies heart disease in patients who are on dialysis is that of diastolic dysfunction (17,19). This physiology results in a sharp increase in LV diastolic pressure with modest increments in LV volume; the implication of the physiology is that patients have a lower threshold to pulmonary edema under these circumstances (Figure 2). Conversely, patients who undergo ultrafiltration on dialysis can experience a sharp fall in LV diastolic pressure under circumstances of a modest volume reduction, thereby risking sudden hypotension and hemodynamic instability. Patients with LVH also often have a reduction in systolic function,
Cardiac arrhythmias and cardiac deaths are common in dialysis patients of all ages but are particularly problematic in patients who are older than 65 yr (21). There are many potential reasons for the problem of sudden death: Abnormalities in the coronary microcirculation; impaired coronary reserve; reduced aortic compliance; increased activity of the sympathetic nervous system; the increased plasma concentration of angiotensin II; and sudden changes in the concentrations of potassium, calcium, and magnesium (which occur normally during dialysis) all can contribute to this problem. Another recent contributor to the problem of sudden death in this patient population may be related to the degree of fibrosis in the hearts of individuals who were on dialysis. Individuals with a higher fibrosis score had a shorter survival than individuals with a lower fibrosis score, suggesting that the presence of dense fibrosis denotes intrinsic myocardial damage and lowers the threshold for arrhythmias.

In the aggregate, then, both CAD and LVH contribute to CV mortality in the dialysis population that dramatically exceeds that of the general population. This has been demonstrated in several recent publications and represents the major challenge for physicians who treat such patients. Figure 3 depicts this increase in CV mortality that occurs in this patient group graphically.

**Causes of Coronary Disease and LVH in CKD/ESRD**

A comprehensive discussion of all of the causes of heart disease in patients with CKD and ESRD is beyond the scope of this communication. This section provides a brief overview of several current issues that are relevant to the problem.

It is known that patients with kidney disease have a high prevalence of other disorders that independently are associated with poor CV outcomes. For example, the high prevalence of diabetes in any CKD/ESRD cohort, the presence of hypertension in virtually all of the patients, the presence of LVH by electrocardiogram or echocardiography in the great majority of patients, that most patients have reduced physical activity and low exercise tolerance, that many patients have a lower-than-ideal HDL cholesterol concentration, that there is high oxidant stress in this group of individuals, and that many individuals have a high concentration of inflammatory biomarkers all contribute to the high prevalence of heart disease in this group. It should be acknowledged that patients in this group do have a lower prevalence of some traditional CV risk factors. For example, this group of patients is less typically obese, uses tobacco less frequently, and has less hypercholesterolemia (lower LDL levels) than the general population.

**Figure 1.** Kaplan-Meier estimates of the rates of death at 3 yr from cardiovascular (CV) causes, reinfarction, congestive heart failure (CHF), stroke, resuscitation after cardiac arrest, and the composite end point, according to the estimated GFR at baseline. From reference (5).

**Figure 2.** Diastolic pressure-volume relation in patients with diastolic heart failure and in control subjects. From reference (19). ---, Diastolic heart failure; —, control.

**Figure 3.** Cardiovascular mortality in general population (NCHS) and dialysis population (USRDS). Sarnak 2000 KI 58: 1758, 2000
disease in patients with kidney disease could result ultimately in LVH and CAD in patients with CKD and ESRD. A number of modifiable risk factors cited above are present in this patient group. Several uremia-related factors may also contribute to the problem; for example, anemia contributes to LVH, and a high calcium/phosphorus product will reduce aortic compliance that leads to an increased pulse wave velocity and an additional burden on the left ventricle. Vasoactive peptide concentrations (e.g., with endothelin and angiotensin II) are elevated and act as potent vasoconstrictors that exacerbate coronary vasoconstriction. The net result of all of these factors is an increased prevalence of coronary disease and LVH in this patient group.

With regard to coronary disease in patients with ESRD, it is increasingly apparent that increased oxidant stress and inflammation are present, and this is reviewed in the following section. A number of nontraditional factors for coronary disease have also been identified in patients with CKD/ESRD. Hyperhomocysteinemia is associated with poorer survival in patients with coronary disease in the dialysis population (23). Recent studies on the ability of folic acid to lower the CV event rate by lowering homocysteine concentrations in patients with ESRD have been disappointing (24). For example, a recent study that used high-dose folic acid in 510 patients and had a mean follow-up of 24 mo did not detect any effect of folic acid on event-free survival (24). New prospective studies on this subject are under way and should reveal some insight into the best management of this problem.

Another area of growing interest and focus has been the heightened recognition that arterial calcification in coronary vessels is very common in patients with CKD and ESRD (25,26). There is currently a debate over whether the calcification in coronary vessels that is detected by electron beam tomography (EBCT) in patients with CKD/ESRD denotes intraluminal narrowing of these vessels. Much of the calcification in these coronary vessels of patients with uremia resides in the media of the vessel, which does not result in intraluminal narrowing of the vessel. A recent study suggested that the calcification score of vessels obtained did not correlate with the severity of vessel stenosis by coronary angiography (27). However, another recent study suggested that the calcium score that is obtained by EBCT is reliable in defining the number of coronary vessels involved in an ESRD population (28).

It is clear that the process of calcification in the coronary bed and other vascular beds is complicated; one new recently proposed factor that may have a major role to play in this calcification scheme is a substance called fetuin-A, which inhibits the mineralization of vascular smooth muscle cells both in vitro and in vivo (29). Even if the calcification of coronary vessels is not linked directly to intraluminal narrowing in patients with CKD and ESRD, the increased vascular stiffness in the aorta increases pulse pressure, which, in turn, results in increased afterload and the burden of cardiac work (30). An increase in pulse wave velocity occurs as renal function decreases, suggesting that there is a continuum of an increased burden on the heart as renal function declines (30). In summary, vessel calcifications are common in patients with CKD and ESRD, particularly in the coronary vascular bed. Having coronary vessel calcifications on EBCT denotes a poorer prognosis than not having them, but there is still controversy over the degree of coronary luminal narrowing that can be ascribed to these calcifications in this patient group.

Another subject of recent interest and study with regard to the cause of coronary disease has been the role of oxidant stress. In this regard, the accumulation of asymmetric dimethyl arginine has been shown to be a potent inhibitor of nitric oxide synthesis in endothelial cells (31). More problematic is the issue of whether antioxidant therapy can reduce CV events in patients with ESRD. In this regard, a recent study in which 134 patients who were on hemodialysis were followed for 14 mo received the potent antioxidant N-acetylcysteine (NAC; 600 mg twice daily) or a placebo. The primary end point in the study was a composite of myocardial infarction, CV death, coronary angioplasty, coronary artery bypass grafting, ischemic stroke, peripheral vascular disease, or peripheral vascular disease with amputations. The result of the study was mixed; 28% of the individuals in the NAC group reached one of the end points versus 47% of the individuals in the control group (P < 0.03) (32). However, for individual end points such as cardiac death, ischemic stroke, and peripheral vascular disease, there was no difference in the groups.

Finally, there is increasing interest as to whether biomarkers can predict individuals who are at particular risk for CV death in the setting of CKD or ESRD. deFilippi et al. (15) suggested that a combination of high C-reactive protein cardiac troponin-T levels not only predicts CV death in patients with ESRD but also identifies the subpopulation of patients with ESRD and multivessel coronary disease (15) (Figure 5). Similarly, the use of N-terminal pro brain natriuretic peptide levels may have some utility in predicting outcome in patients with ESRD as well (33). As more pathophysiologic factors of importance are identified for LVH and coronary disease and more biomarkers are linked to these processes, it is likely that there will be an array of screening tests that will identify a particular patient’s particular risk for a CV death.
Can Both Heart and Renal Disease Be Affected Favorably by Lowering of BP, Inhibition of the Renin Angiotensin System, and Lowering of Urinary Protein Excretion?

Given that the prevalence of impaired kidney function continues to increase (34) in the context of higher BP and proteinuria (35,36), the question arises as to whether therapies that are directed at these abnormalities yield both a cardiac and a renal protective effect. This cardio–renal relationship is particularly evident in individuals with diabetes (37–39), a population of patients that accounts for the greatest proportion of the CKD/ESRD group (40). This section reviews the possibilities that interventions aimed at lowering BP via the inhibition of the renin-angiotensin system (RAS) and lowering protein excretion may be beneficial to both heart and kidney function.

Is Lowering BP Equally Cardio- and Renoprotective?

A large number of studies have repeatedly confirmed that treatment of increased BP decreases both cardiac and particularly cerebrovascular events in patients with diabetes (41–44) as well as those without diabetes (45,46). Such an effect of lowering BP is also one of the primary findings of a large meta-analysis that was performed by the Blood Pressure Lowering Treatment Trialist Collaboration that concluded, “Any commonly used regimen reduces the risk of total CV events and larger reductions in BP produce larger reductions in risk” (47). A similar protection afforded by decreasing BP on the progression of renal disease emerged from the analysis performed by Bakris et al. (48), who found a significant (P < 0.05) relationship between the achieved mean arterial pressure in 10 trials that were performed in patients with diabetes and the rate of decline in GFR in the same trials. The trials that demonstrated CV protection in individuals with diabetes were performed predominantly or almost exclusively in patients who had little or no underlying renal disease. Because the presence of renal disease per se, as noted above, imparts an added risk, it is unclear whether the results of those trials can be readily extrapolated to patients with impaired renal function. Furthermore, although target BP have been lowered progressively, it is not clear which component of the BP is important and whether the optimal BP is the same for the heart as it is for the kidney. In this regard, most CV trials have scant renal end points, and, conversely, renal trials are frequently devoid of CV outcome data. Recent trials in patients with overt diabetic nephropathy (49,50) monitored both renal and CV events, allowing perhaps for the first time the assessment of the effect of BP reduction on the CV and renal systems in the same patient population.

An analysis of the effect of achieved BP in the Irbesartan Diabetic Nephropathy Trial (IDNT) (51) sheds some light on the question at hand. This trial is a double blind, randomized, placebo-controlled trial performed in 209 clinics worldwide in patients with overt diabetic nephropathy (urinary protein >900 mg/24 h) and a serum creatinine concentration up to 3.0 mg/dl. Of the 1715 patients enrolled in the trial, 1590 had at least a 6-mo follow-up. In this group of patients, over a mean follow-up period of 2.6 yr, an average of eight BP were available per patient. As is depicted in Figure 6, the relative risk (RR) for reaching a renal end point (doubling of serum creatinine or ESRD, grouped by 10-mmHg increments) reveals that the risk progressively decreases as achieved systolic BP (SBP) decreases from 180 to 120 mmHg. The group below 120 mmHg did not have a risk substantially different from that of the 121- to 130-mmHg cohort. A 20-mmHg decrement in SBP was associated with a 47% risk reduction for developing a renal end point. Furthermore, achieved SBP is an independent risk factor for adverse renal outcomes, independent of the baseline SBP (51). These effects of SBP to slow progression of renal disease were also described in a trial of individuals without diabetes (52) but were not observed in the African American Study of Kidney Disease and Hypertension (AASK) trial of black individuals.
The IDNT revealed no significant relation between diastolic BP (DBP) and renal outcomes (51).

An analysis of achieved BP and all-cause mortality/composite CV mortality in this same patient population provides a somewhat different picture (Figure 7). As is evident from the examination of Figure 7, the risk for both all-cause mortality (Figure 7A) and CV mortality (Figure 7B) rose in the subgroup of patients whose SBP was <120 mmHg by a RR of 3.05 and 4.06, respectively (both \( P < 0.001 \)). There were only 53 patients in this subgroup, and an examination as to how these patients differed from the other 1537 patients revealed that they had a much higher prestudy history of heart disease and congestive heart failure (54). It is of note that when the effect of different levels of SBP are analyzed in only those patients with an SBP >120 mmHg, there is a highly significant protection for lowering SBP to this level for all-cause mortality, CV mortality, and congestive heart failure but not for risk for myocardial infarction or stroke (54).

In contrast to the neutral effect of changes in DBP on renal outcomes, decrements in DBP were associated with an opposing effect on the rate of myocardial infarction and strokes. Thus, whereas the former, as depicted in Figure 8, increased markedly at a DBP <85 mmHg per 10 mmHg (RR 1.61; 95% confidence interval 1.26 to 2.02; \( P < 0.001 \)), the risk for strokes decreased (RR 0.65; 95% confidence interval 0.48 to 0.88; \( P < 0.005 \)) (54).

The increased risk for acute coronary events again raises the old controversy of whether there is a J-point effect. Its existence has been championed by some (55) but questioned by others (56). Cruickshank (55) argued that at a DBP <85 mmHg, there may be a J-point effect in patients with underlying ischemic disease, as the coronary circulation is particularly sensitive to a decrease in DBP. The population studied in IDNT of elderly patients with type 2 diabetes and overt nephropathy is very likely to be enriched by such patients. Furthermore, it is of interest to note that the INVEST trial that studied patients with hypertension and CAD (57) also found a distinct association of lower DBP with myocardial infarctions (58). Likewise, a meta-analysis of seven large, randomized trials also revealed a J point at a DBP of 84 mmHg on all-cause mortality (59). Because decreasing DBP is associated with increasing pulse pressure, the possibility that an increment in pulse pressure may be culpable for these findings has been suggested. The findings in the Framingham cohort that revealed that an increase in CV events occurred in patients with a low DBP and an SBP >140 mmHg (60) supports such a contention. Although the effect of low DBP and increased pulse pressure are difficult to dissociate, it is of interest that in both INVEST and IDNT, the decrement of low DBP was not associated with an increase in strokes, suggesting that there is no overall vasculotoxic effect of an...
increased pulse pressure and that the consequences of a low DBP are specific to the coronary circulation.

In summary, therefore, as pertains to BP, it seems that the progressive lowering of SBP to 120 mmHg is associated with improved renal protection and a decrement in mortality and CV events. However, at an SBP <120 mmHg, there is continued renal protection but an increase in all-cause mortality, CV mortality, and congestive heart failure in patients who have more severe underlying cardiac disease. When examined in populations that are enriched with patients with CAD, a J-curve effect is observed at a DBP <85 mmHg for acute myocardial infarction but not for renal progression. The effect of low DBP cannot be readily dissociated from the increase in pulse pressure; the failure to observe an increment in cerebrovascular events suggests an independent effect that most likely is mediated by impairment in coronary artery perfusion.

**Does Inhibition of RAS Protect Both Heart and Kidney?**

Numerous studies that were published in the 1980s and 1990s consistently showed that the inhibition of angiotensin-converting enzyme (ACE) reduces the risk for death and CV events after myocardial infarction (61–65) and increases survival in patients with decreased LV function (66–68). Such cardioprotective effects, particularly in hospitalizations related to congestive heart failure, were observed in both the Reduction of Endpoint in NIDDM with the Angiotensin II Antagonist Losartan (RENAAL) (50) and IDNT (69) studies, which use angiotensin receptor blockers (ARB). In fact, these two classes of drugs (ACE inhibitors and ARB) seem to provide a comparable beneficial effect on CV outcomes both in patients with heart failure (70,71) and in patients with postmyocardial infarction (72,73). Likewise, on the background of a large body of experimental data suggesting a salutary effect of aldosterone on progressive cardiac fibrosis (74), the use of aldosterone inhibitors such as with spironolactone (75) or eplerenone (76) have been found to decrease significantly mortality in patients with advanced heart failure.

The renoprotective effect of RAS inhibition on the progression of renal disease has also been documented amply in individuals both with (77) and without diabetes (78). In fact, RAS inhibition may retard the progression of diabetic renal disease at every stage. Specifically, ACE inhibitors but not calcium channel blockers decrease the risk for conversion from normoalbuminuria to microalbuminuria (79). Similarly, both ACE inhibitors (80) and ARB (81) significantly decrease the rate of progression from microalbuminuria to overt proteinuria, but the two classes do not seem to differ in the rate at which renal function is lost in these patients (82). Finally, in patients with overt proteinuria, two independent studies have demonstrated a positive effect of ARB on the rate of loss of GFR (49,50).

Whether such protection is also afforded by ACE inhibitors is not entirely known, as the results in small series have yielded variable results (83). Because the large trials did not have an ACE inhibitor component, this question may well remain unanswered. In analogy with the observation described above in cardiac protection, there is increasing interest in the possibility that aldosterone, independent of the RAS inhibition, could be implicated in the progression of renal disease. This has support in studies performed in experimental animals (84,85), but there are studies that suggest an antiproteinuric effect associated with aldosterone inhibition (86,87).

Thus Bianchi et al. (87) observed in an uncontrolled pilot study a reversible decrease in protein excretion when 42 patients with CKD received 25 mg of Aldactone for 8 wk. These observations clearly call for the study of the effect of such agents on the progression of kidney disease with more established clinical end points.

The question that arises is whether there is an interaction between lowering BP and assignment to a RAS inhibitor. This interaction was examined in recent reports for both reduction in the rate of loss of renal function (51) and the risk for congestive heart failure (54). For the former, the analysis is depicted in Figure 9. It is of note that at any quartile of achieved SBP, the patients who were placed on the ARB had better renal outcomes than those who were assigned to the alternative treatment arms. In fact, assignment to the ARB resulted in a 33% risk reduction ($P < 0.001$) of reaching a renal end point beyond that achieved by lowering the SBP. The effects of lowering SBP and treatment with the ARB were completely independent and additive (51). A similar independent effect was observed for lowering SBP and assignment to the ARB in the decrement of congestive heart failure hospitalizations. Thus, while being on the ARB lowered the risk for heart failure by 29% and lowering SBP by 20 mmHg did so by 25%, the combination of both lowered the risk by 53% (54).

In summary, inhibition of the RAS with either an ACE inhibitor or an ARB provides both cardiac and renal protection. The protection is most marked for congestive heart failure. Inhibition of aldosterone provides survival advantage in patients with low LV ejection fraction and may also have antiproteinuric effects. The heart failure and renoprotective effects of RAS inhibition are independent and additive to the benefit obtained from lowering BP.

![Figure 9. Simultaneous impact of quartile of achieved SBP and treatment modality on the RR of reaching a renal end point. Reprinted from reference (51), with permission.](image-url)
Is Lowering Protein Excretion Equally Reno- and Cardioprotective?

Almost every multifactorial analysis of risk factors that predict the progression of kidney disease consistently points to the baseline level of proteinuria as an important predictor of such an event. This was observed in a study in individuals with type 1 diabetes (88), in the Gruppo Italiano di Studi Epidemiologici in Nefrologia study that also involves individuals without diabetes (89), and in the aforementioned RENAAL (90) and IDNT trials (91). Likewise, baseline proteinuria also has been found to be a predictor of CV events in these last two trials (92,93), as well as in other studies, such as Losartan Intervention for Endpoint Reduction in Hypertension (LIFE) (94), Heart Outcomes Prevention Evaluation (HOPE) (95) and Framingham (96). Thus, there is little question that proteinuria is a marker or even a surrogate for enhanced renal and cardiac risk. What remains to be proved is whether lowering protein excretion per se reduces the risk. This was examined in a subanalysis of the aforementioned trials. Specifically, in RENAAL, a >30% reduction in protein excretion at 6 mo was associated with a significantly decreased risk for reaching a renal end point than those who did not do so (92). In fact, the analysis suggests that much of the renoprotective effect of the ARB antagonist that was used was mediated by lowering protein excretion. The reduction in protein excretion was also associated with a decrement in CV events such that halving of the protein excretion resulted in halving of the CV risk (92). In regard to renal protection, very similar data emerged from the analysis of IDNT (91), as a 50% reduction in proteinuria at 12 mo was associated with an approximate 50% reduction in reaching a renal end point. Although further data on this issue would be welcome, it seems that lowering protein excretion clearly is beneficial in protecting the kidney and probably the heart as well.

In summary, in answering the initially posed question as to whether what is good for the kidney is good for the heart, it is clear that lowering BP protects the kidney, but there may be levels of SBP and DBP below which some increase in risk emerges. A BP of 120/85 mmHg seems to be a “rational” target as of now. Inhibiting the RAS clearly is beneficial to both organs, but the protection that is afforded by aldosterone inhibition remains to be defined convincingly for the kidney. Finally, lowering urinary protein seems to slow the progression of kidney disease and also may be cardioprotective, but this latter observation will require further confirmatory data.

Treatment Options for LVH and CAD

As stated at the outset, relatively few studies have examined prospectively in randomized trials one therapeutic strategy for overt heart disease versus another in patients with CKD or ESRD. With regard to the management of LVH, two studies have probed the potential benefit of a β blocker in this setting (97,98). In the first of these studies, the β blocker carvedilol was used in ESRD patients with classes II and III New York Heart Association heart failure. During a 12-mo period, there was some improvement in the carvedilol group as compared with the placebo-treated group (97). In a follow-up of this patient group over several years, there was superior survival and fewer hospitalizations in the β blocker–treated group as compared with placebo (98). Taken together, these studies suggest that there may be a beneficial role for β blockers in the setting of advanced heart failure in dialysis patients. It should be stressed that in these trials, the use of a β blocker was not accomplished easily, as fully 21% of individuals who were exposed to the carvedilol therapy were forced to drop out of the study.

With regard to the management of CAD, there is an ongoing debate of the relative advantage of coronary artery bypass grafting versus percutaneous interventions. Several years ago, one study suggested that coronary artery bypass grafts were associated with an improved survival compared with percutaneous procedures (99). There are several reasons for why such a result might have been obtained, including that many patients with more complicated medical problems (and who therefore possess a very high risk for death in the periprocedure period) are excluded from coronary artery bypass grafting surgery. In addition, percutaneous procedures have improved in recent years. The clinical and angiographic restenosis rate in patients who undergo percutaneous interventions has been reduced dramatically recently (100). This field is moving rapidly as reports of improved patency with drug-eluting stents in this group of patients is now being reported. For example, in one such report of 10 consecutive patients who were on dialysis, 10 of 10 patients had a very good result that was durable for >1 yr without any deaths, recurrent myocardial infarction, or the need for target lesion revascularization (100).

Our current strategy in the population of patients with CKD and ESRD is to use a combination of low-dose aspirin and β blockers in patients with known coronary disease. The potential benefit of β blockers in all patients with stage 4/5 CKD and ESRD must await a randomized, prospective study. As mentioned above, there is no consensus on whether antioxidants are beneficial, and new studies are planned to answer this question. A recent study of the cholesterol-lowering agent atorvastatin in 1255 dialysis patients yielded mixed results over an average follow-up period of 4 yr (101). On the one hand, cardiac events were somewhat reduced in the atorvastatin-treated patients, but the RR of fatal stroke was increased in this group. This mixed result led to no significant effect on the composite end point of the study. These findings will require additional analysis and study, but our practice is to use statins to reduce LDL cholesterol concentrations in concordance with current guidelines (7). For patients with diabetes, tight glucose control is important, as is evident from the discussion on BP and protein excretion; the antihypertensive agent of choice in the patient population is a drug that interrupts the RAS axis, either an ACE inhibitor or an ARB. Extracellular fluid volume control via longer dialysis sessions or extra dialysis sessions is currently also becoming more usual. Whether nocturnal dialysis or quotidian dialysis will be effective and affordable in treating patients in the United States is under active study at present. Strict management of calcium and phosphorus is currently accomplished through a combination of phosphate binders, including non–calcium-containing phosphate binders. Guidelines for the management of the hematocrit and dyslipidemia now suggest that the hematocrit should be in the middle to upper 30s as a
minimum and the LDL cholesterol should be lowered to <100 mg/dl in dialysis patients and <70 mg/dl in patients with known coronary disease.

References


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This In-Depth Review by Berl and Henrich should provide background for the paper by Khan et al. in JASN (pages 244–253), which relates mortality from heart failure (systolic) to kidney function.