

Reversal of Arterial Stiffness and Maladaptative Arterial Remodeling After Kidney Transplantation

Alexandre Karras, MD, PhD; Pierre Boutouyrie, MD, PhD; Marie Briet, MD, PhD; Erwan Bozec, PhD; Jean-Philippe Haymann, MD, PhD; Christophe Legendre, MD; Lawrence P. McMahon, MBBS, MD; Michel Delahousse, MD

Background—Chronic kidney disease is characterized by stiffening, thinning, dilatation, and increased circumferential wall stress of large arteries, associated with increased cardiovascular risk. Kidney transplantation (KT) reverses many pathological features of chronic kidney disease and improves life expectancy; however, longitudinal studies exploring the impact of KT on recipient large arteries are scarce.

Methods and Results—This study was designed to appraise arterial changes following KT. Carotid-femoral pulse wave velocity, carotid remodeling (circumferential wall stress and carotid internal diameter), and stiffness were measured in 161 consecutive recipients receiving either a living (n=49) or a deceased (n=112) donor allograft, at 3 and 12 months after transplantation. Mean pulse wave velocity decreased from 10.8 m/s (95% confidence interval, 10.5–11.2 m/s) (at month 3) to 10.1 m/s (95% confidence interval, 9.8–10.5 m/s) (at month 12) ($P<0.001$). After multivariate adjustment, pulse wave velocity reduction from month 3 to month 12 was significantly larger in the living donor allograft KT ($P<0.001$). Circumferential wall stress decreased, 70 kPa (95% confidence interval, 68–72 kPa) to 64 kPa (95% confidence interval, 62–67 kPa), as well as carotid internal diameter and carotid stiffness ($P<0.001$ for all). Reductions in circumferential wall stress, diameter, and stiffness were significantly larger in the living donor allograft KT ($P<0.001$). When deceased donor allograft patients were classified into standard and expanded criteria donors, changes in both pulse wave velocity and circumferential wall stress were blunted in expanded criteria donors. Changes were independent of graft function and blood pressure changes.

Conclusions—Large-artery stiffness and maladaptive carotid artery remodeling of chronic kidney disease is partially reversed within 12 months of KT and appears unrelated to renal function. Improvements were independently associated with live organ donation. Our data suggest that expanded criteria donors may hamper vascular recovery. (*J Am Heart Assoc.* 2017;6:e006078. DOI: 10.1161/JAHA.117.006078.)

Key Words: renal disease • stiffness • transplantation • vascular function • vascular remodeling

Chronic kidney disease (CKD) accelerates aging of the arterial vascular tree, with complex changes producing large-vessel stiffening and outward remodeling characterized as dilatation, decreased thickness, and increased wall stress.^{1–4} These changes possibly explain the high cardiovascular mortality rate observed among patients with CKD and especially with end-stage renal disease.^{5–7}

Aortic stiffness has been used for several years as a strong independent predictive marker of cardiovascular mortality, in populations with diabetes mellitus and hypertension in particular.^{8,9} Carotid to femoral pulse wave velocity (PWV) is a simple and reproducible method to measure aortic stiffness, and is highly predictive of cardiovascular mortality in patients with end-stage renal disease¹⁰ and kidney transplant and

From the Nephrology Department, Hôpital Européen Georges Pompidou (A.K.), Pharmacology Department, Hôpital Européen Georges Pompidou (P.B., M.B., E.B.), Physiology Department, Hôpital Tenon (J.-P.H.), and Renal Transplantation Department, Hôpital Necker (C.L.), Assistance Publique Hôpitaux de Paris, Paris, France; Nephrology and Renal Transplantation Department, Hôpital Foch, Suresnes, France (M.D.); Institut National de la Santé et de la Recherche Médicale U970-PARCC, Paris, France (A.K., P.B., M.B., E.B.); Université Paris Descartes, Paris, France (A.K., P.B., M.B.); Université Pierre et Marie Curie, Paris, France (J.-P.H.); Eastern Health Clinical School, Monash University, Melbourne, Australia (L.P.M.); INSERM U 1018 CESP Team 5, UVSQ, Villejuif, France (M.D.).

Accompanying Tables S1 through S5 are available at <http://jaha.ahajournals.org/content/6/9/e006078/DC1/embed/inline-supplementary-material-1.pdf>

Correspondence to: Alexandre Karras, MD, PhD, Nephrology Department, Hôpital Européen Georges Pompidou, Assistance Publique Hôpitaux de Paris, 20, rue Leblanc, 75015 Paris, France. E-mail: alexandre.karras@aphp.fr

Received March 9, 2017; accepted June 29, 2017.

© 2017 The Authors. Published on behalf of the American Heart Association, Inc., by Wiley. This is an open access article under the terms of the Creative Commons Attribution-NonCommercial-NoDerivs License, which permits use and distribution in any medium, provided the original work is properly cited, the use is non-commercial and no modifications or adaptations are made.

Clinical Perspective

What Is New?

- Kidney transplantation can reverse the modifications of large arteries induced by chronic kidney disease, such as stiffening, dilatation, decreased thickness, and increased wall stress.
- Improvement of vascular health in allograft recipients is more pronounced when transplantation is performed with living donors, independently of post-transplant renal function.
- In contrast, kidney transplantation with extended criteria deceased donors—those older than 60 or those older than 50 years with a history of cardiovascular disease—may hamper arterial recovery of the recipient.

What Are the Clinical Implications?

- In patients with chronic kidney disease with significant cardiovascular risk factors and/or complications, living donor kidney transplantation may contribute to the improvement of post-transplant cardiovascular morbidity and mortality.
- Kidney recipients of extended criteria deceased donors may not benefit from this post-transplant reversal of arterial modifications.

CKD,^{3,11} which is remarkable in the absence of predictive equations in these populations. We and others have shown that not only aortic stiffness but also internal diameter enlargement, thinning of the arterial wall, increased circumferential wall stress, and reduction of the arterial distensibility occur early during CKD and progress in parallel with the fall in renal function and influence prognosis.^{1–4,12}

Kidney transplantation (KT) reduces cardiovascular risk in patients with end-stage renal disease, with cardiovascular death rates reduced by $\approx 75\%$ in transplant recipients when compared with matched patients undergoing dialysis still on the waiting list.^{13,14} This important benefit is mainly explained by the reversal of renal dysfunction. However, the mechanism of cardiovascular risk improvement is yet to be adequately elucidated since transplant-specific diseases, especially side effects of immunosuppressive drugs, may offset the decrease in cardiovascular risk.

Several but not all studies have suggested that PWV may fall after KT,^{15,16} which may represent a global improvement in vascular health translating into improvement in graft and patient survival.^{11,17,18} However, longitudinal data relating to large-vessel vascular remodeling in a transplant population are lacking, especially concerning specific changes in vessel diameter, thickness and wall stress. We hypothesized that such changes can occur following KT. Therefore, the objective of this noninterventional, prospective study was to describe

the arterial modifications following KT and to identify the key determinants of this process.

Methods

Design and Patients

This prospective single-center study ran from January 2009 to the end of December 2012. Eligible patients were aged 18 to 70 years and recipients of a first living or cadaveric single KT. A body mass index $<35 \text{ kg/m}^2$ was required. Immunosuppression regimens had to include a calcineurin inhibitor. Estimated glomerular filtration rate (GFR) at month 3 (M3), time of the first arterial parameters evaluation, had to be $>30 \text{ mL/min per } 1.73 \text{ m}^2$. Noninclusion criteria were cardiac arrhythmia, symptomatic lower limb arteriopathy and dual-kidney or combined organ transplantation.

A total of 307 KTs were performed at the transplantation department of Foch Hospital during this period. A total of 232 consecutive patients met the inclusion criteria (169 deceased donor [DD] allograft KT and 63 living donor [LD] allograft KT), of whom 192 were enrolled and 161 completed the study. Of note, 31 patients did not complete the study because of graft failure ($n=12$), death ($n=2$), or personal reasons ($n=17$). Enrolled patients underwent an extended evaluation including medical interview, clinical examination, blood samples, direct measured GFR through ⁵¹Cr-EDTA urinary clearance, and study of arterial parameters (see below) at M3 and 12 months (M12) after transplantation.

Baseline clinical data included recipient's age and sex, cardiovascular risk factors and previous cardiovascular events, cause of CKD, and duration of dialysis therapy.

The main characteristics of the kidney donor were also collected, including age, sex, cardiovascular risk factors, serum creatinine, and estimated GFR before organ donation. Kidney donors were classified as LDs, standard criteria donors (SCDs), or expanded criteria donors (ECDs), using classical criteria.¹⁹

Post-transplant clinical events were collected, such as delayed graft function, biopsy-proven acute rejections episodes, and diagnosis of new-onset diabetes mellitus after transplantation. All medications, including immunosuppressant, antihypertensive, statin, and diabetic therapies were listed.

The protocol was approved by the local ethics committee, and all patients gave written informed consent before enrolment.

Arterial Parameters

Patients were studied in a quiet, temperature-controlled room ($22 \pm 1^\circ\text{C}$) room as previously described.¹ Changes

were calculated for every measured parameter between M3 and M12 (M12–M3, negative figures meaning decrease). Blood pressure was monitored with a standard oscillometric method and mean blood pressure (MBP) was derived from systolic and diastolic blood pressure using the 1/3 formula.

Aortic stiffness was estimated by the carotid to femoral PWV using the validated foot-to-foot velocity method (Sphygmocor, Atcor Medical). Briefly, pressure waveforms were obtained at the common carotid artery and the femoral artery with a pressure transducer. The transit time was measured between the feet of the 2 waveforms by intersecting tangents. The distance was measured between the 2 sites (carotid-femoral). PWV was calculated as carotid-femoral distance (m)/transit time (s), corrected (distance \times 0.8) according to published recommendations.²⁰ Measurement of carotid-femoral PWV was always performed by choosing the side opposite to renal allograft arterial anastomosis.

Common carotid artery pressure waveforms were recorded noninvasively by applanation tonometry (Sphygmocor, Atcor Medical) and local carotid artery pulse pressure was used for further calculations.

End-diastolic internal diameter, change in internal diameter between systole and diastole, and intima-media thickness (IMT) were measured on the right common carotid artery with a high-precision echotracking system (Artlab system, Esaote), as previously described and validated. Carotid internal diameter (CID) was normalized to body surface area. Circumferential wall stress (CWS), given in kPa, was calculated according to the Lamé's equation as: (MBP.CID)/2IMT. Carotid distensibility was determined from systolic-diastolic variations in arterial cross-sectional area (ΔA) and local pulse pressure (ΔP) as described earlier, assuming the lumen to be circular. Cross-sectional distensibility coefficient was calculated as $\Delta A/A\Delta P$. Carotid stiffness was calculated as (distensibility coefficient)^{-1/2}.

Biological Parameters

Blood and urine samples were collected at M3 and M12 to determine values of serum creatinine, glucose, glycated hemoglobin, lipid profile (high-density lipoprotein/low-density lipoprotein and triglyceride), C-reactive protein, calcium, and phosphorus. Proteinuria was expressed as urinary protein: creatinine ratio (uPCR).

GFR was measured by the renal clearance of ⁵¹Cr-EDTA, as mentioned. Briefly, 1.8 to 3.5 MBq of ⁵¹Cr-EDTA (GE Healthcare) was injected intravenously as a single bolus. After allowing 1 hour for distribution of the tracer in the extracellular fluid, average renal ⁵¹Cr-EDTA clearance was determined on 5 to 6 consecutive 30-minute clearance periods.

Statistical Analyses

Data are expressed as mean \pm SD, mean (95% confidence interval [CI]), or median [interquartile range] as appropriate. Group comparisons were made using 2-sample *t* test or Wilcoxon rank test as appropriate. Chi-square test was used for discrete variables. Within-group comparisons (change between M3 and M12) were made using the paired *t* test or the Wilcoxon signed rank nonparametric test as appropriate. Associations between arterial parameter changes (as dependent variable) and all other independent variables related to recipient characteristics, donor characteristics, transplantation procedure (eg, cold ischemia time), main post-transplant events (delayed graft function, acute rejection, new-onset diabetes mellitus after transplantation), and treatments (eg, renin-angiotensin system blockers, immunosuppressive agents) were tested using stepwise multiple robust regressions. All tests were 2-sided and *P* values <0.05 were deemed significant.

All statistical analyses were performed using NCSS 10 Statistical Software (2015) (NCSS, LLC).

Results

Demographics and Clinical Data

Of the 161 transplant recipients who completed the study, 49 (30.4%) received LD kidneys and 112 (69.6%) received cadaveric kidneys, 60 of whom were SCDs and 52 ECDs. The mean age was 50 \pm 11 years, with DD recipients tending to be older. A total of 68% of patients were men, with a similar preponderance in both LD and DD groups. The cause of end-stage renal disease was also similar between groups, with the most frequent causes including primary glomerulonephritis (19%), polycystic kidney disease (22%), diabetic nephropathy (10%), and hypertensive nephropathy (10%) (Table 1).

Renal function (measured GFR) was adequate at M3 (54; 95% CI, 43–64 mL/min per 1.73 m²) and did not change appreciably during the study, with no difference between the LD and DD groups at either M3 or M12. Similarly, proteinuria was minimal (uPCR, 0.03; 95% CI, 0.02–0.04 g/mmol) and did not alter during the study. None of the biological variables were independently associated with arterial changes, except for a weak association between uPCR at M12 and PWV change.

In the LD group (n=49), the median duration of dialysis treatment before transplantation was shorter compared with the DD group (*P*<0.001). There were no episodes of delayed graft function (*P*<0.001) and the cold ischemia time was markedly reduced: 1.4 hours (95% CI, 1.0–1.7) versus 16.6 hours (95% CI, 12.2–22.6) (*P*<0.001), when compared with the DD group.

Table 1. Clinical and Demographic Data of Study Population

	All Patients (N=161)	Living Donors (n=49)	Deceased Donors (n=112)	P Value	Standard Criteria Donors (n=60)	Expanded Criteria Donors (n=52)	P Value
Recipient age, y	50 (49–52)	48 (44–51)	51 (49–53)	0.1	47 (45–50)	56 (54–58)	<0.001
Men/women	110/51	37/49	73/112	0.1	40/60	33/52	0.7
BMI, kg/m ²	24 (24–25)	25 (23–26)	24 (23–25)	0.3	24 (23–25)	24 (23–25)	0.7
Time on dialysis, mo	30 [10–51]	7 [1–24]	37 [21–66]	<0.001	39 [23–64]	36 [19–70]	0.9
Pre-KT diabetes mellitus, %	12	10	12	0.7	7	19	0.05
Pre-KT dyslipidemia, %	47	43	51	0.5	38	60	0.03
Pre-KT former tobacco, %	22	26	24	0.9	22	23	0.9
Preemptive KT, No.	10	9	1	<0.001	2	0	0.2
Donor age, y	51 (49–53)	50 (47–53)	52 (49–54)	0.4	42 (40–45)	63 (61–65)	<0.001
DGF, %	14	0	22	<0.001	17	25	0.2
Acute rejection, %	9	7	11	0.3	15	6	0.1
NODAT, %	19	10	23	0.05	18	29	0.2
ARB or ACEI, %	67	65	69	0.6	65	73	0.005
mGFR M3, mL/min per 1.73 m ²	53 [43–64]	52 [44–60]	55 [44–66]	0.8	58 [49–70]	51 [39–61]	0.05
mGFR M12, mL/min per 1.73 m ²	54 [45–68]	57 [46–65]	52 [42–69]	0.3	60 [46–71]	47 [38–60]	0.001
uPCR M3, g/mmol	0.03 [0.02–0.04]	0.03 [0.02–0.04]	0.02 [0.02–0.04]	0.8	0.02 [0.02–0.04]	0.02 [0.02–0.05]	0.5
uPCR M12, g/mmol	0.02 [0.01–0.03]	0.02 [0.01–0.03]	0.02 [0.01–0.04]	0.9	0.02 [0.01–0.03]	0.02 [0.01–0.04]	0.4

Recipient age, donor age, and body mass index (BMI) values are expressed as mean (95% confidence interval). Time on dialysis, measured glomerular filtration rate (mGFR), and urine protein/creatinine ratio (uPCR) values are expressed as median [interquartile range]. ACEI indicates angiotensin-converting enzyme inhibitor; ARB, angiotensin receptor blocker; DGF, delayed graft function; KT, kidney transplantation; M3, month 3 after kidney transplantation; M12, month 12 after kidney transplantation; NODAT, new-onset diabetes mellitus after transplantation.

The standard immunosuppressive regimen included corticosteroids (96%), mycophenolate mofetil (100%), and tacrolimus or cyclosporin (91% and 9%, respectively). Induction immunosuppressive therapy included polyclonal antilymphocyte antibodies (Thymoglobulin) in 43% of all patients, anti-interleukin 2 receptor monoclonal antibodies in 43%, and polyvalent intravenous immunoglobulins in 27%. There was a tendency to a lower incidence of new-onset diabetes mellitus after transplantation in the LD group and a higher incidence in the ECD group.

Other medications were widely used and included angiotensin-converting enzyme inhibitors/angiotensin receptor blockers (69%), calcium channel blockers (56%), statins (43%), and β -blockers (40%). The mean number of antihypertensive agents used per patient was 1.8 ± 1.1 . No difference was identified between the LD and DD groups.

Arterial Stiffness: PWV and Carotid Stiffness

PWV decreased from 10.8 (95% CI, 10.5–11.2) to 10.2 (95% CI, 9.8–10.5) ($P < 0.001$) during the study. This occurred in parallel with a small albeit significant decrease in MBP (-3 mm Hg; 95% CI, -4 to -1). Once adjusted for changes

in MBP, changes in PWV remained virtually identical (data not shown). The fall in PWV was more marked ($P < 0.001$) in the LD group (-1.4 m/s; 95% CI, -1.8 to -1.0) than in the DD group (-0.4 m/s; 95% CI, -0.6 to -0.1). Carotid stiffness showed similar changes to PWV between M3 and M12 (-0.2 m/s; 95% CI, -0.4 to 0.0) ($P < 0.05$), with changes again greater ($P < 0.001$) in the LD group (Figure 1A, Table 2).

Carotid Remodeling: CID and IMT

The fall in CID was significant overall ($P < 0.001$), but again more marked in the LD group: -0.4 mm (95% CI, -0.5 to -0.3) versus -0.1 mm (95% CI, -0.2 to 0.0) ($P < 0.01$). IMT rose significantly during the study (17 μ m; 95% CI, 7 – 28 [$P < 0.01$]), an effect largely driven by the LD group (44 μ m; 95% CI, 20 – 66), as there was no appreciable change in the DD group (Figure 1B). Other indices of carotid stiffness were also modified. We observed a 10% fall in Young's elastic modulus (Figure 1C) (-47 kPa; 95% CI, -79 to -14 [$P < 0.01$]), with a 31% fall in the LD group ($P < 0.001$). No changes were found in the DD group. Circumferential wall stress integrates both the changes in diameter and IMT, and decreased by 9% overall ($P < 0.001$), which mainly related to a 17% reduction in the LD

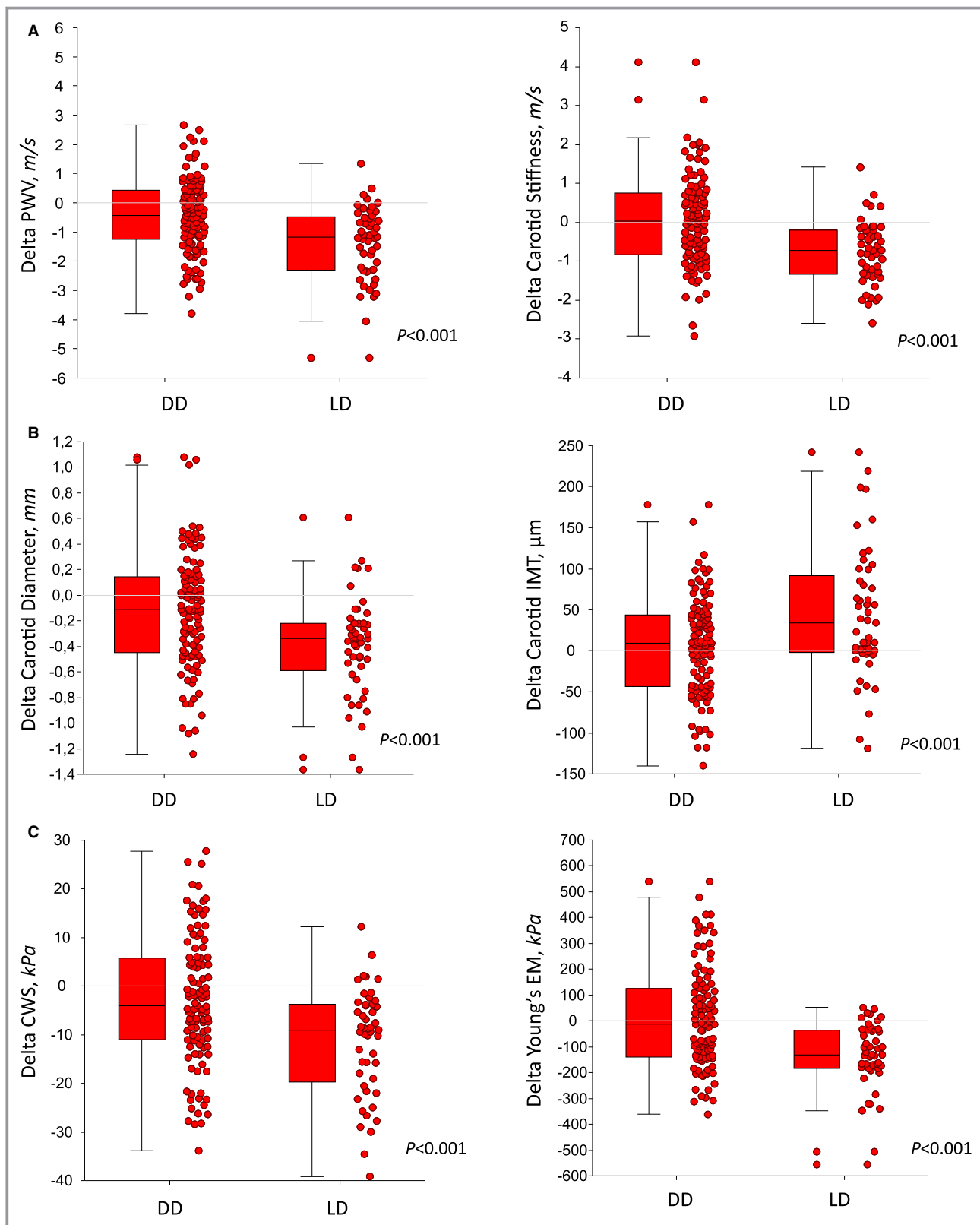


Figure 1. Changes in measured and calculated parameters between 3 and 12 months post transplant in 161 recipients according to donor source (deceased donor [DD] vs live donor [LD]). A, Pulse wave velocity (PWV) and carotid stiffness. B, Carotid internal diameter and intima-media thickness. C, Circumferential wall stress (CWS) and Young's elastic modulus (EM). Box plots represent median, interquartile range, and adjacent values. Comparisons are for between-group changes.

Downloaded from <http://ahajournals.org> by on August 23, 2018

Table 2. Baseline Values and Changes of Arterial Parameters During the Study

Variable	All Patients (N=161)	LDs (n=49)	DDs (n=112)	P Value	ECDs (N=52)	SCDs (n=60)	P Value
Baseline arterial parameters (at M3)							
MBP, mm Hg	98 (97–100)	99 (96–102)	98 (96–100)	0.5	99 (96–102)	97 (94–100)	0.4
PWV, m/s	10.8 (10.5–11.2)	11.2 (10.5–11.9)	10.7 (10.3–11.0)	0.4	10.6 (10.1–11.2)	10.7 (10.2–11.2)	0.7
Carotid stiffness, m/s	6.6 (6.4–6.8)	6.7 (6.3–7.0)	6.6 (6.3–6.9)	0.5	6.8 (6.3–7.2)	6.4 (6.0–6.8)	0.2
Carotid diameter, mm	6.4 (6.2–6.5)	6.2 (6.0–6.5)	6.4 (6.3–6.4)	0.2	6.6 (6.4–6.8)	6.3 (6.1–6.5)	<0.05
IMT, μ m	601 (586–616)	583 (552–613)	610 (593–627)	0.1	622 (598–646)	599 (575–623)	0.2
CWS, kPa	70 (68–72)	71 (66–75)	70 (67–73)	0.6	71 (67–75)	69 (64–73)	0.4
Young's EM, kPa	476 (441–512)	466 (410–522)	481 (436–527)	0.8	508 (436–579)	457 (398–517)	0.4
Variable	All Patients	LDs	DDs	P Value	ECDs	SCDs	P Value
Change in arterial parameters from M3 to M12							
MBP, mm Hg	-3 (-4 to -1) [†]	-4 (-7 to -1) [*]	-2 (-4 to 0) [*]	0.3	-3 (-0.6 to 0) [*]	-1 (-4 to 1)	0.3
PWV, m/s	-0.7 (-0.9 to -0.5) [‡]	-1.4 (-1.8 to -1) [‡]	-0.4 (-0.6 to 0.1) [†]	<0.001	0.1 (-0.4 to 0.4)	-0.7 (-1 to -0.4) [‡]	<0.01
Carotid stiffness, m/s	-0.2 (-0.4 to 0) [*]	-0.9 (-1.1 to -0.6) [‡]	0.1 (-0.2 to 0.3)	<0.001	0.1 (-0.2 to 0.4)	0.1 (-0.3 to 0.4)	0.7
Carotid diameter, mm	-0.2 (-0.3 to -0.1) [‡]	-0.4 (-0.5 to -0.3) [‡]	-0.1 (-0.2 to 0) [†]	<0.001	-0.1 (-0.2 to 0.1)	-0.2 (-0.3 to -0.1) [‡]	0.1
IMT, μ m	17 (7–28) [†]	44 (20–66) [‡]	6 (-6 to 17)	<0.001	1 (-13 to 16)	10 (-8 to 27)	0.5
CWS, kPa	-6 (-8 to -4) [‡]	-11 (-15 to -11) [‡]	-3 (-6 to -1) [†]	<0.001	-3 (-7 to 1)	-4 (-7 to 0) [†]	0.8
Young's EM, kPa	-47 (-79 to -14) [†]	-145 (-187 to -102) [‡]	-2 (-43 to 38)	<0.001	6 (-52 to 65)	-10 (-68 to 48)	0.7

Values are expressed as mean (95% confidence interval). CWS indicates circumferential wall stress; DDs, deceased donors; ECDs, extended criteria donors; EM, elastic modulus; IMT, intima-media thickness; LDs, living donors; M3, month 3 after kidney transplantation; M12, month 12 after kidney transplantation; MBP, mean arterial blood pressure; PWV, pulse wave velocity; SCDs, standard criteria donors. P, statistical difference between groups (LD:DD; ECD:SCD); * P<0.05, † P<0.01, and ‡ P<0.001 for change within groups.

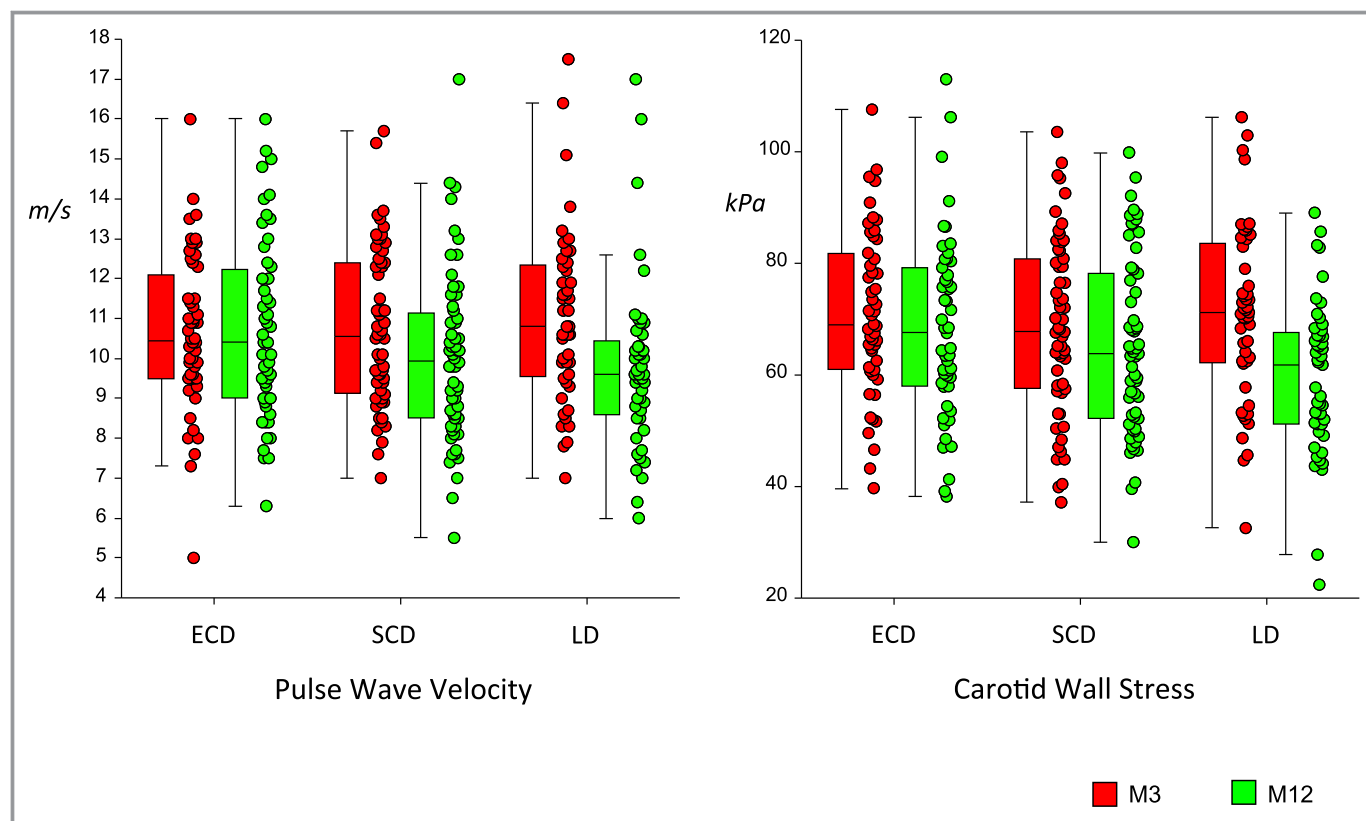


Figure 2. Measured pulse wave velocity and circumferential wall stress at 3 months (red) and at 12 months (green) post transplant in 161 recipients according to donor source. The deceased donor group is further subdivided into standard criteria donor (SCD) and expanded criteria donor (ECD) groups. Box plots represent median, interquartile range, and adjacent values. Comparisons are for within-group differences. LD indicates living donor; M3, month 3; M12, month 12.

group, compared with a 4% reduction in the DD group ($P<0.001$ for difference) (Table 2).

LD Versus SCD Versus ECD

When the DD group was further divided into SCDs and ECDs, it became evident that most of the improvement found in this group was related to changes in the SCD subgroup. There was no significant difference between the ECD and SCD groups for sex balance, time on dialysis, rate of delayed graft function, rate of new-onset diabetes mellitus after transplantation, acute rejection rate, and BMI either at M3 or M12. However, and by construction, the ECD group was older (56 ± 11 years versus 47 ± 11 years, $P=0.0001$) and received kidneys from older donors (56 ± 9 versus 42 ± 10 , $P=0.0001$) than the SCD group. GFR was significantly lower in the ECD than in the LD and SCD groups at M3 and M12 ($P<0.01$). Patients receiving SCD kidneys had significant reductions for PWV ($P<0.001$), CID ($P<0.001$), and CWS ($P<0.05$), and favorable trends in carotid stiffness, IMT, or Young's elastic modulus. By contrast, no parameter improved in the ECD group, despite a significant fall in MBP ($P<0.05$). When compared with the LD group, the

improvement in PWV, carotid diameter, IMT, and CWS was observed to be “stepped,” suggesting that the improvement in the SCD group was midway between the LD and ECD groups (Figure 2). These differences were unaltered by adjustment for potential confounding factors (Tables 1 and 2).

Correlations

Five variables consistently and highly significantly correlated with changes in the 5 parameters studied (PWV, carotid stiffness, CID, IMT, and CWS, $N=161$) during the study. These variables were: (1) basal (M3) measurements, (2) donor source (LD/DD) or (3) characteristics of the transplanted kidney (LD/SCD/ECD), (4) cold ischemia time, and (5) time on dialysis. Basal values at M3 were similarly correlated, with the direction of change supportive of a greater change in patients with worse initial pathology. Reduction in MBP correlated with improvement of PWV ($P<0.001$), carotid stiffness ($P<0.001$), CID ($P<0.05$), and Young's elastic modulus ($P<0.001$) (Table 3, Tables S1 through S5).

Multivariate analysis including those covariates identified specifically and consistently that donor source (LD versus DD,

Table 3. Multivariate Analysis of Changes in Measured Parameters According to Donor Source (LDs or DDs)

Independent Variable	Beta Coefficient (95% CI)	R (Standardized Coefficient)	P Value	Partial R ²
PWV (R²=0.33)				
Donor source (LDs/DDs)	−0.85 (−0.24 to −0.46)	−0.36	<0.001	10.5
Basal PWV	−0.24 (−0.33 to −0.15)	−0.35	<0.001	16.2
Δ MBP	0.03 (0.01–0.05)	0.24	0.001	6.3
Log uProt/uCreat	0.24 (0.06–0.43)	0.14	0.011	4.1
Carotid CWS (R²=0.29)				
Donor source (LDs/DDs)	−7.1 (−11.1 to −3.1)	−0.27	<0.001	9.6
Basal CWS	−0.35 (−0.47 to −0.23)	−0.45	<0.001	22.5
Carotid IMT (R²=0.09)				
Donor source (LDs/DDs)	26.9 (6.2–47.6)	0.20	0.02	4.1
Basal IMT	−0.11 (−0.21 to −0.00)	−0.16	<0.05	2.6
Carotid stiffness (R²=0.38)				
Donor source (LDs/DDs)	−0.73 (−1.06 to −0.41)	−0.29	<0.001	11.8
Basal carotid stiffness	−0.30 (−0.41 to −0.20)	−0.37	<0.001	18.1
Δ MBP	0.03 (0.02–0.05)	0.29	<0.001	11.5
Δ BMI	0.11 (0.02–0.19)	0.17	0.02	4.2
Carotid diameter (R²=0.26)				
Donor source (LDs/DDs)	−0.27 (−0.41 to −0.14)	−0.31	<0.001	11.3
Δ BMI	0.06 (0.03–0.09)	0.26	0.001	8.3
Basal carotid diameter	−0.11 (−0.18 to −0.04)	−0.23	<0.005	6.4
Δ MBP	0.01 (0.00–0.01)	0.20	0.01	5.2

Δ indicates change; BMI, body mass index; CWS, circumferential wall stress; DDs, deceased donors; IMT, intima-media thickness; LDs, living donors; MBP, mean arterial blood pressure; PWV, pulse wave velocity; uProt/uCreat, urine protein/urine creatinine ratio.

Table 3), or transplanted kidney characteristics (LD/SCD/ECD, Table 4), and basal values at M3 were the primary independent determinants of isobaric change among the significant variables. Donor source and transplanted kidney characteristics were highly associated with improvement ($P<0.001$) of all variables including increase in IMT ($P<0.05$) (Tables 3 and 4).

Discussion

This study demonstrates improved aortic stiffness and reversal of maladaptive vascular remodeling in transplant recipients in the first year following KT. We demonstrated that: (1) both large-artery stiffening and maladaptive outward remodeling of CKD can significantly improve within the first year after KT, and (2) the donor source (living versus cadaveric) and kidney characteristics (living, standard criteria, or expanded criteria) are the main independent determinants of changes in arterial properties.

These changes have to be interpreted in the context of large-artery changes in CKD. We have previously shown that

early stages in CKD (2–5) are marked by maladaptive remodeling of large arteries associating increased stiffness, dilatation, and thinning, resulting in increased wall stress. We have further shown that the intensity of this maladaptive remodeling was associated with the evolution of CKD (notably wall stress) and morbid cardiovascular events (notably aortic stiffness and carotid dilatation).^{1–4} Although previous studies have identified significant and potentially favorably prognostic modifications in aortic stiffness^{16,21} and/or carotid IMT²² within several months of KT, these studies have been either cross-sectional or have primarily addressed the relationship between carotid IMT and inflammation.

The major finding of this study is that KT reverses the maladaptive remodeling of large arteries particularly if the transplanted kidney comes from an LD. These patients showed a 13% improvement in both PWV and carotid stiffness, 6% increase in ICD, 7% increase in IMT, and a 17% reduction in CWS. These changes are important since they correspond grossly to 10/15 years of normal vascular aging.^{23–25} This improvement persisted unchanged after adjustment for dialysis duration and other potential

Table 4. Multivariate Analysis of Changes in Measured Parameters According to Transplanted Kidney Characteristics (LDs, SCDs, ECDs)

Independent Variable	Beta Coefficient (95% CI)	R (Standardized Coefficient)	P Value	Partial R ² , %
PWV ($R^2=0.37$)				
ECDs vs SCDs	0.73 (0.48–0.98)	0.44	<0.001	17.3
LDs vs SCDs	−0.63 (−0.89 to −0.38)	−0.37	<0.001	13.2
Basal PWV	−0.21 (−0.30 to −0.13)	−0.33	<0.001	14.2
Δ MBP	0.04 (0.02–0.06)	0.27	<0.001	10.4
Carotid CWS ($R^2=0.29$)				
ECDs vs SCDs	3.33 (1.01–5.66)	0.23	0.012	4.9
LDs vs SCDs	−0.63 (−0.89 to −0.37)	−0.33	<0.001	9.7
Basal CWS	−0.32 (−0.42 to −0.21)	−0.46	<0.001	22.7
Carotid IMT ($R^2=0.09$)				
ECDs vs SCDs	−10.55 (−23.93 to 2.84)	−0.14	0.2	1.5
LDs vs SCDs	17.95 (4.14–31.75)	0.24	<0.05	4.1
Basal IMT	−0.36 (−0.46 to −0.25)	−0.18	<0.05	3.2
Carotid stiffness ($R^2=0.36$)				
ECDs vs SCDs	0.43 (0.22–0.64)	0.31	<0.001	9.6
LDs vs SCDs	−0.63 (−0.89 to −0.37)	−0.37	<0.001	13.1
Basal carotid stiffness	−0.32 (−0.42 to −0.21)	−0.41	<0.001	19.7
Δ MBP	0.03 (0.02–0.05)	0.27	<0.001	10.0
Carotid diameter ($R^2=0.28$)				
ECDs vs SCDs	0.19 (0.11–0.27)	0.38	<0.001	12.1
LDs vs SCDs	−0.19 (−0.27 to −0.11)	−0.38	<0.001	12.4
Basal carotid diameter	−0.13 (−0.20 to −0.06)	−0.26	0.001	8.5
Δ MBP	0.01 (0–0.01)	0.21	0.007	5.6
Δ BMI	0.05 (0.02–0.08)	0.25	0.001	8.0

Δ indicates change; BMI, body mass index; CI, confidence interval; CWS, circumferential wall stress; ECDs, extended criteria donors; IMT, intima-media thickness; LDs, living donors; MBP, mean arterial blood pressure; PWV, pulse wave velocity; SCDs, standard criteria donors.

confounders. In addition, when the DD data were further split into SCD and ECD, it was evident that all effective improvement in vascular compliance and remodeling in the DD group were observed in the SCD group and not in the ECD group. The lack of change in this latter group is notable, since ECD patients were similar to their SCD counterparts in respect to time on dialysis, rate of delayed graft function, and episodes of acute rejection. Measured GFR was lower in the ECD group (Table 1) but the difference in arterial changes between groups was still significant after adjustment for measured graft function. The patients in the ECD group were both older and received older kidneys. Vascular health deteriorates with age, and it is possible that the greater age of ECD recipients reduced the capacity for vessel changes. In addition, we have previously shown that donor age has an important impact on recipient aortic stiffness in the post-transplant period,²⁶ which, together with the deleterious effect of brain death and

cold ischemia on kidney function, can clearly influence the capacity for vascular aging reversal and, potentially, long-term renal and cardiovascular prognosis. Two recent, large epidemiological studies demonstrated that: (1) kidney allografts from live donors or young cadaveric donors were associated with better overall post-transplant mortality,^{27,28} and (2) ECD allografts correlated with a higher risk of graft failure.²⁹

The contrasting findings between the 2 groups (ECD versus SCD) in this study raise the issue of optimization of kidney allocation. In the context of a shortage of organ supply, a growing number of patients receive kidney allografts from donors aged older than 60 years and/or with vascular comorbidities. Although this strategy does offer a significant benefit for quality of life and survival compared with long-term dialysis therapy,³⁰ as shown here it is associated with suboptimal correction of major cardiovascular risk markers. It is not possible to address this issue with any certainty from

the current data; however, our results do support the promotion of live donation for KT, and also suggest that an awareness of long-term cardiovascular prognosis is a relevant consideration when accepting kidneys for donation.

The fact that renal function did not relate to vascular stiffness has been observed in several trials, especially when GFR was measured directly, rather than by equation.¹ Renal function was stable during the study, although patients with clear graft failure were excluded, thus minimizing any potential effect on GFR. The duration of observation might also be too short, and longer-term analysis of this cohort may shed light on whether relatively mild renal dysfunction influences the return of vascular properties to normal, and whether one donor subgroup is more likely to experience clinical sequelae than another.

Study Limitations

The main limitations to this study are that it is observational by nature and thus causality cannot be ascertained. The difference in waiting time on dialysis between the LD and DD groups was a potential confounder, although such a difference was not evident between the SCD and ECD groups, and results persisted after adjustment for this variable. More specific biological markers of CKD such as mineral metabolism parameters or vasculotoxic uremic toxins might have a specific influence on vascular changes. However, most are strongly linked with renal function³¹ and it is not clear whether any have a significant independent direct effect on arterial wall properties. Finally, we must underline the fact that our study population was characterized by a particularly low (12%) prevalence of pretransplant diabetes mellitus, when compared with the usual kidney transplant populations. In other words, the results of this study concern an intermediate cardiovascular risk population but the fact that the basal values of arterial stiffness were correlated with the magnitude of the post-transplant improvement suggest that our results may also apply to patients with more severe vascular disease.

Conclusions

Our data demonstrate that the maladaptive vascular remodeling observed in patients with CKD can improve within 12 months of KT. This improvement is independent of BP and renal function and appears most evident with live donation.

Sources of Funding

This study was supported by a grant from the French Ministry of Health (PHRC 2010—AOM10008—MIN 00-08).

Disclosures

None.

References

- Briet M, Bozec E, Laurent S, Fassot C, London GM, Jacquot C, Froissart M, Houillier P, Boutouyrie P. Arterial stiffness and enlargement in mild-to-moderate chronic kidney disease. *Kidney Int.* 2006;69:350–357.
- Briet M, Collin C, Karras A, Laurent S, Bozec E, Jacquot C, Stengel B, Houillier P, Froissart M, Boutouyrie P; Nephrotest Study Group. Arterial remodeling associates with CKD progression. *J Am Soc Nephrol.* 2011;22:967–974.
- Karras A, Haymann JP, Bozec E, Metzger M, Jacquot C, Maruani G, Houillier P, Froissart M, Stengel B, Guardiola P, Laurent S, Boutouyrie P, Briet M; Nephrotest Study Group. Large artery stiffening and remodeling are independently associated with all-cause mortality and cardiovascular events in chronic kidney disease. *Hypertension.* 2012;60:1451–1457.
- Briet M, Boutouyrie P, Laurent S, London GM. Arterial stiffness and pulse pressure in CKD and ESRD. *Kidney Int.* 2012;82:388–400.
- Muntner P, Bowling CB, Gao L, Rizk D, Judd S, Tanner RM, McClellan W, Warnock DG. Age-specific association of reduced estimated glomerular filtration rate and albuminuria with all-cause mortality. *Clin J Am Soc Nephrol.* 2011;6:2200–2207.
- Go AS, Chertow GM, Fan D, McCulloch CE, Hsu CY. Chronic kidney disease and the risks of death, cardiovascular events, and hospitalization. *N Engl J Med.* 2004;351:1296–1305.
- van der Velde M, Matsushita K, Coresh J, Astor BC, Woodward M, Levey A, de Jong P, Gansevoort RT; Chronic Kidney Disease Prognosis Consortium. Lower estimated glomerular filtration rate and higher albuminuria are associated with all-cause and cardiovascular mortality. A collaborative meta-analysis of high-risk population cohorts. *Kidney Int.* 2011;79:1341–1352.
- Laurent S, Cockcroft J, Van Bortel L, Boutouyrie P, Giannattasio C, Hayoz D, Pannier B, Vlachopoulos C, Wilkinson I, Struijker-Boudier H; European Network for Non-invasive Investigation of Large Arteries. Expert consensus document on arterial stiffness: methodological issues and clinical applications. *Eur Heart J.* 2006;27:2588–2605.
- Cruickshank K, Riste L, Anderson SG, Wright JS, Dunn G, Gosling RG. Aortic pulse-wave velocity and its relationship to mortality in diabetes and glucose intolerance: an integrated index of vascular function. *Circulation.* 2002;106:2085–2090.
- Guerin AP, Blacher J, Pannier B, Marchais SJ, Safar ME, London GM. Impact of aortic stiffness attenuation on survival of patients in end-stage renal failure. *Circulation.* 2001;103:987–992.
- Verbeke F, Marechal C, Van LS, Van BW, Devuyst O, Van Bortel LM, Jadoul M, Vanholder R. Aortic stiffness and central wave reflections predict outcome in renal transplant recipients. *Hypertension.* 2011;58:833–838.
- Townsend RR, Wimmer NJ, Chirinos JA, Parsa A, Weir M, Perumal K, Lash JP, Chen J, Steigerwalt SP, Flack J, Go AS, Rafeq M, Rahman M, Sheridan A, Gadegebeku CA, Robinson NA, Joffe M. Aortic PWV in chronic kidney disease: a CRIC ancillary study. *Am J Hypertens.* 2010;23:282–289.
- Wolfe RA, Ashby VB, Milford EL, Ojo AO, Ettenger RE, Agodoa LY, Held PJ, Port FK. Comparison of mortality in all patients on dialysis, patients on dialysis awaiting transplantation, and recipients of a first cadaveric transplant. *N Engl J Med.* 1999;341:1725–1730.
- Ojo AO, Hanson JA, Meier-Kriesche H, Okechukwu CN, Wolfe RA, Leichtman AB, Agodoa LY, Kaplan B, Port FK. Survival in recipients of marginal cadaveric donor kidneys compared with other recipients and wait-listed transplant candidates. *J Am Soc Nephrol.* 2001;12:589–597.
- Zoungas S, Kerr PG, Chadban S, Muske C, Ristevski S, Atkins RC, McNeil JJ, McGrath BP. Arterial function after successful renal transplantation. *Kidney Int.* 2004;65:1882–1889.
- Covic A, Goldsmith DJ, Gusbeth-Tatomir P, Buhaescu I, Covic M. Successful renal transplantation decreases aortic stiffness and increases vascular reactivity in dialysis patients. *Transplantation.* 2003;76:1573–1577.
- Bahous SA, Stephan A, Blacher J, Safar M. Cardiovascular and renal outcome in recipients of kidney grafts from living donors: role of aortic stiffness. *Nephrol Dial Transplant.* 2012;27:2095–2100.
- Dahle DO, Eide IA, Asberg A, Leivestad T, Holdaas H, Jenssen TG, Fagerland MW, Pihlstrom H, Mjoen G, Hartmann A. Aortic stiffness in a mortality risk calculator for kidney transplant recipients. *Transplantation.* 2015;99:1730–1737.
- Kasiske BL, Zeier MG, Chapman JR, Craig JC, Ekberg H, Garvey CA, Green MD, Jha V, Josephson MA, Kiberd BA, Kreis HA, McDonald RA, Newmann JM, Obrador GT, Vincenti FG, Cheung M, Earley A, Raman G, Abariga S, Wagner M,

- Balk EML. KDIGO clinical practice guideline for the care of kidney transplant recipients: a summary. *Kidney Int.* 2010;77:299–311.
20. Van Bortel LM, Laurent S, Boutouyrie P, Chowienczyk P, Cruickshank JK, De Backer T, Filipovsky J, Huybrechts S, Mattace-Raso FU, Protogerou AD, Schillaci G, Segers P, Vermeersch S, Weber T; Artery Society, European Society of Hypertension Working Group on Vascular Structure and Function, European Network for Noninvasive Investigation of Large Arteries. Expert consensus document on the measurement of aortic stiffness in daily practice using carotid-femoral pulse wave velocity. *J Hypertens.* 2012;30:445–448.
 21. Verbeke F, Van BW, Honkanen E, Wikstrom B, Jensen PB, Krzesinski JM, Rasmussen M, Vanholder R, Rensma PL. Prognostic value of aortic stiffness and calcification for cardiovascular events and mortality in dialysis patients: outcome of the calcification outcome in renal disease (CORD) study. *Clin J Am Soc Nephrol.* 2011;6:153–159.
 22. Yilmaz MI, Sonmez A, Saglam M, Cayci T, Kilic S, Unal HU, Karaman M, Cetinkaya H, Eyileten T, Gok M, Oguz Y, Vural A, Mallamaci F, Zoccali C. A longitudinal study of inflammation, CKD-mineral bone disorder, and carotid atherosclerosis after renal transplantation. *Clin J Am Soc Nephrol.* 2015;10:471–479.
 23. Reference Values for Arterial Stiffness' Collaboration. Determinants of pulse wave velocity in healthy people and in the presence of cardiovascular risk factors: 'establishing normal and reference values'. *Eur Heart J.* 2010;31:2338–2350.
 24. Engelen L, Ferreira I, Stehouwer CD, Boutouyrie P, Laurent S; Reference Values for Arterial Measurements Collaboration. Reference intervals for common carotid intima-media thickness measured with echotracking: relation with risk factors. *Eur Heart J.* 2013;34:2368–2380.
 25. Engelen L, Bossuyt J, Ferreira I, van Bortel LM, Reesink KD, Segers P, Stehouwer CD, Laurent S, Boutouyrie P; Reference Values for Arterial Measurements Collaboration. Reference values for local arterial stiffness. Part A: carotid artery. *J Hypertens.* 2015;33:1981–1996.
 26. Delahousse M, Chaignon M, Mesnard L, Boutouyrie P, Safar ME, Lebre T, Pastural-Thaumat M, Tricot L, Kolko-Labadens A, Karras A, Haymann JP. Aortic stiffness of kidney transplant recipients correlates with donor age. *J Am Soc Nephrol.* 2008;19:798–805.
 27. Molnar MZ, Streja E, Kovesdy CP, Shah A, Huang E, Bunnapradist S, Krishnan M, Kopple JD, Kalantar-Zadeh K. Age and the associations of living donor and expanded criteria donor kidneys with kidney transplant outcomes. *Am J Kidney Dis.* 2012;59:841–848.
 28. Chavalitdhamrong D, Gill J, Takemoto S, Madhira BR, Cho YW, Shah T, Bunnapradist S. Patient and graft outcomes from deceased kidney donors age 70 years and older: an analysis of the Organ Procurement Transplant Network/United Network of Organ Sharing database. *Transplantation.* 2008;85:1573–1579.
 29. Pascual J, Zamora J, Pirsch JD. A systematic review of kidney transplantation from expanded criteria donors. *Am J Kidney Dis.* 2008;52:553–586.
 30. Merion RM, Ashby VB, Wolfe RA, Distant DA, Hulbert-Shearon TE, Metzger RA, Ojo AO, Port FK. Deceased-donor characteristics and the survival benefit of kidney transplantation. *JAMA.* 2005;294:2726–2733.
 31. Neiryck N, Eloit S, Glorieux G, Barreto DV, Barreto FC, Liabeuf S, Lenglet A, Lemke HD, Massy ZA, Vanholder R. Estimated glomerular filtration rate is a poor predictor of the concentration of middle molecular weight uremic solutes in chronic kidney disease. *PLoS One.* 2012;7:e44201.

SUPPLEMENTAL MATERIAL

Table S1. Univariate analysis of Pulse Wave Velocity (PWV) change

Independent Variable	R	p
Recipient-related parameters		
Recipient age	0.13	0.126
Sex (male=1)	-0.04	0.616
Diabetes (1/0)	0.06	0.428
Previous dyslipidemia (1/0)	0.03	0.736
Smoking (past) (1/0)	0.05	0.561
Smoking (active) (1/0)	0.03	0.654
Previous MCE (1/0)	-0.04	0.602
Time on dialysis (all recipients)	0.18	0.025
Time on dialysis (DD)	0.07	0.447
M3 PWV	-0.33	0.000
Delta MBP	0.24	0.000
Donor-related parameters		
Donor source (LD=1/DD=0)	-0.34	0.000
Transplanted Kidney characteristics (ECD/SCD/LD)	-0.39	0.000
Transplantation-related parameters		
Cold ischemia time (All recipients)	0.297	0.000
Cold ischemia time (DD)	0.089	0.352
Graft function parameters		
mGFR M12	-0.079	0.317
Log Prot/CreatU M12	0.25	0.022
Main post-transplant events		
DGF (1/0)	0.065	0.415
BPAR (1/0)	0.067	0.398
Delta BMI	0.093	0.244
NODAT (1/0)	0.001	0.989

Table S2. Univariate analysis of Carotid Stiffness change

Independent Variable	R	p
Recipient-related parameters		
Recipient age	-0.00	0.869
Sex (male=1)	-0.18	0.025
Diabetes (1/0)	0.03	0.682
Previous dyslipidemia (1/0)	-0.03	0.705
Smoking (past) (1/0)	-0.09	0.213
Smoking (active) (1/0)	0.05	0.556
Previous MCE (1/0)	0.03	0.741
Time on dialysis (all recipients)	0.21	0.008
Time on dialysis (DD)	0.07	0.447
M3 carotid stiffness	-0.37	0.000
Delta MBP	0.30	0.000
Donor-related parameters		
Donor source (LD=1/DD=0)	-0.35	0.000
Transplanted Kidney characteristics (ECD/SCD/LD)	-0.30	0.000
Transplantation-related parameters		
Cold ischemia time (All recipients)	0.27	0.000
Cold ischemia time (cadaveric donor kidney recipients)	0.05	0.636
Graft function parameters		
mGFR M12	0.04	0.662
Log Prot/CreatU M12	0.13	0.114
Main post-transplant events		
DGF (1/0)	0.03	0.750
BPAR (1/0)	0.03	0.673
Delta BMI	0.24	0.00
NODM (1/0)	0.145	0.068

Table S3. Univariate analysis of Circumferential Wall Stress (CWS) change

Independent Variable	R	p
Recipient-related parameters		
Recipient age	0.04	0.653
Sex (male=1)	0.03	0.701
Diabetes (1/0)	0.11	0.161
Previous dyslipidemia (1/0)	-0.07	0.408
Smoking (past) (1/0)	-0.06	0.432
Smoking (active) (1/0)	0.03	0.689
Previous MCE (1/0)	0.03	0.692
Time on dialysis (all recipients)	0.24	0.003
Time on dialysis (DD)	0.13	0.186
M3 cws	-0.41	0.000
Donor-related parameters		
Donor source (LD=1/DD=0)	-0.29	0.000
Transplanted Kidney characteristics (ECD/SCD/LD)	-0.26	0.000
Transplantation-related parameters		
Cold ischemia time (All recipients)	0.19	0.019
Cold ischemia time (DD)	-0.03	0.792
Graft function parameters		
mGFR M12	-0.03	0.691
Log Prot/CreatU M12	0.07	0.397
Main post-transplant events		
DGF (1/0)	-0.02	0.780
BPAR (1/0)	-0.07	0.364
Delta BMI	0.05	0.504
NODM (1/0)	0.17	0.036

Table S4. Univariate analysis of Carotid IMT change

Independent Variable	R	p
Recipient-related parameters		
Recipient age	0.09	0.271
Sex (male=1)	-0.11	0.152
Diabetes (1/0)	0.00	0.993
Previous dyslipidemia (1/0)	0.07	0.368
Smoking (past) (1/0)	0.08	0.346
Smoking (active) (1/0)	0.06	0.481
Previous MCE (1/0)	0.03	0.702
Time on dialysis (all recipients)	-0.19	0.017
Time on dialysis (DD)	-0.07	0.445
M3 carotid IMT	-0.18	0.025
Donor-related parameters		
Donor source (LD=1/DD=0)	0.26	0.001
Transplanted Kidney characteristics (ECD/SCD/LD)	0.25	0.002
Transplantation-related parameters		
Cold ischemia time (All recipients)	-0.22	0.006
Cold ischemia time (DD)	-0.10	0.299
Graft function parameters		
mGFR M12	0.03	0.747
Log Prot/CreatUM12	0.06	0.474
Main post-transplant events		
DGF (1/0)	0.01	0.939
BPAR (1/0)	-0.06	0.431
Delta BMI	-0.03	0.675
NODM (1/0)	-0.117	0.141

Table S5. Univariate analysis of Carotid Diameter change

Independent Variable	R	p
Recipient-related parameters		
Recipient age	0.05	0.569
Sex (male=1)	0.02	0.832
Diabetes (1/0)	0.18	0.021
Previous dyslipidemia (1/0)	0.03	0.678
Smoking (past) (1/0)	-0.05	0.547
Smoking (active) (1/0)	0.04	0.584
Previous MCE (1/0)	0.06	0.468
Time on dialysis (all recipients)	0.11	0.151
Time on dialysis (DD)	0.02	0.873
M3 carotid diameter	-0.16	0.043
Donor-related parameters		
Donor source (LD=1/DD=0)	-0.28	0.000
Transplanted Kidney characteristics (ECD/SCD/LD)	-0.31	0.000
Transplantation-related parameters		
Cold ischemia time (All recipients)	0.24	0.004
Cold ischemia time (DD)	0.09	0.343
Graft function parameters		
mGFR M12	-0.14	0.072
Log Prot/CreatU M12	0.02	0.762
Main post-transplant events		
DGF (1/0)	0.01	0.939
BPAR (1/0)	-0.02	0.800
Delta BMI	0.24	0.003
NODM (1/0)	0.03	0.700