

META-ANALYSIS

The prognosis of kidney transplant recipients with aorto-iliac calcification: a systematic review and meta-analysis

Elsaline Rijkse¹ , Jacob L. van Dam¹ , Joke I. Roodnat² , Hendrikus J. A. N. Kimenai¹ , Jan N. M. IJzermans¹  & Robert C. Minnee¹ 

¹ Division of HPB and Transplant Surgery, Department of Surgery, Erasmus MC University Medical Center, Rotterdam, The Netherlands
² Division of Nephrology, Department of Internal Medicine, Erasmus MC University Medical Center, Rotterdam, The Netherlands

Correspondence

Robert C. Minnee MD, PhD,
 Department of Surgery RG-234,
 Erasmus Medical Center, Postbus
 2040, 3000 CA Rotterdam, The
 Netherlands.
 Tel.: +31107031810;
 fax: +31107032396;
 e-mail: r.minnee@erasmusmc.nl

SUMMARY

The prognosis of kidney transplant recipients (KTR) with vascular calcification (VC) in the aorto-iliac arteries is unclear. We performed a systematic review and meta-analysis to investigate their survival outcomes. Studies from January 1st, 2000 until March 5th, 2019 were included. Outcomes for meta-analysis were patient survival, (death-censored) graft survival and delayed graft function (DGF). Twenty-one studies were identified, eight provided data for meta-analysis. KTR with VC had a significantly increased mortality risk [1-year: risk ratio (RR) 2.19 (1.39–3.44), 5-year: RR 2.28 (1.86–2.79)]. The risk of 1-year graft loss was three times higher in recipients with VC [RR 3.15 (1.30–7.64)]. The risk of graft loss censored for death [1-year: RR 2.26 (0.58–2.73), 3-year: RR 2.19 (0.49–9.82)] and the risk of DGF (RR 1.24, 95% CI 0.98–1.58) were not statistically different. The quality of the evidence was rated as very low. To conclude, the presence of VC was associated with an increased mortality risk and risk of graft loss. In this small sample size, no statistical significant association between VC and DGF or risk of death-censored graft loss could be demonstrated. For interpretation of the outcomes, the quality and sample size of the evidence should be taken into consideration.

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Key words

atherosclerosis, graft survival, kidney transplantation, meta-analysis, systematic review

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Introduction

As kidney transplant recipients (KTR) are becoming older and vascular disease is more prevalent, the challenge of transplanting a kidney onto atherosclerotic aorto-iliac arteries is likely to become more common. Nearly 25% of all kidney transplant candidates have vascular calcification (VC) in the aorto-iliac arteries on lumbar X-ray [1]. Risk factors for VC are common risk factors for vascular disease such as diabetes,

smoking, hypertension and dyslipidemia [2,3]. Patients suffering from end-stage renal disease (ESRD) are at higher risk to develop vascular disease due to added risk factors like chronic uremia, use of calcium-based phosphate binders and, most importantly, dialysis treatment [4]. Severe VC in the aorto-iliac arteries has been considered as a relative contra-indication for kidney transplantation (KTx). It has been found that 43% of all transplant candidates who are considered ineligible is due to VC [5].

There are several reasons why KTx in patients with VC in the aorto-iliac arteries can be problematic. First, the vascular anastomosis itself may be technically challenging. Over the years, transplant surgeons found various ways to overcome this issue. A Fogarty catheter can be used in case of an unclampable iliac artery [6]. In case of compromised blood flow in the external iliac artery (EIA) caused by VC, a percutaneous transluminal angioplasty (PTA), endarterectomy or vascular bypass can be performed in a staged or simultaneous procedure [7]. Second, vascular complications like steal syndrome or trash foot may be a threat to the vascular challenging transplant candidate [8–10]. As a third reason, patients with VC are considered to have a limited life expectancy due to cardiovascular comorbidities, leading to a high perioperative mortality risk and limiting 5-year patient survival to 35% [11,12]. Due to organ shortage, it might not be ethical to perform a deceased donor KTx in patients with such a limited life expectancy. On the other hand, the quality of life is significantly improved if a transplant can be performed successfully [13]. Therefore, proper recipient evaluation is of paramount importance.

To date, there is scant information about the prognosis of KTR with VC in the aorto-iliac arteries. Most published studies are too small to provide definite conclusions about survival outcomes. We performed a systematic review and meta-analysis concerning clinical outcomes after KTx in patients with VC. The primary objective of this systematic review and meta-analysis was to evaluate the risk of mortality and graft loss in KTR with VC in comparison to KTR without VC. As secondary outcomes, we investigated the risk of DGF and 1-year kidney function.

Methods

This systematic review and meta-analysis was performed according to the guidelines for observational studies as described in the Preferred Reporting Items for Systematic review and Meta-Analysis Protocols (PRISMA) guidelines [14].

Search strategy

Together with the help from a clinical librarian, we searched Embase, Medline, Web of Science, Cochrane and Google Scholar database. A search for the Embase database was created and the search terms for other databases were derived from this one. The search included the following terms: kidney/renal transplantation,

atherosclerosis, iliac artery. The first search was performed on August 2nd 2017 and the last search on March 5th, 2019. Detailed search strategies are included in Table S1.

Study selection

The studies were firstly screened on title and abstract by two independent reviewers (ER and JLD). Eligible study designs were cross-sectional studies, cohort studies and case-control studies. Studies were included if they reported either patient survival, uncensored graft survival, death-censored graft survival, DGF or kidney function. Studies were included in meta-analysis if they compared clinical outcomes between KTR with any degree of VC in the aorto-iliac arteries and KTR without VC. Also, studies describing KTx outcomes after treatment of VC [through PTA, endarterectomy (EAT) or vascular bypass] were eligible for systematic review. For both the meta-analysis part as well as the systematic review part, the following exclusion criteria were used: conference abstracts, systematic or narrative reviews, studies published not in the English language, studies including multi-organ transplantation and studies published before January 1st, 2000. Disagreements were discussed between both reviewers and, if necessary, consulted with a third party (RCM). References were manually checked for relevant studies.

Quality assessment

The quality of the evidence was assessed using the GRADE tool for prognosis studies [15,16]. For the GRADE tool, risk of bias, heterogeneity, directness of the evidence, precision of effect estimates and risk of publication bias were assessed for every outcome. The Newcastle-Ottawa scale for cohort studies was adopted to assess the quality of each individual study [17]. Studies were graded according to selection of study groups, comparability and ascertainment of exposure and outcomes. ER and JLD assessed the studies independently.

Data collection and extraction

Data extraction was completed by two independent authors. The following items were extracted from included studies that did not provide data for meta-analysis: study design, sample size per group, donor type, age, dialysis treatment, and treatment for aorto-iliac calcification. For studies included in meta-analysis, the following data was extracted: study design, sample size per group, and possible confounding factors such as recipient age,

sex, smoking history, hypertension, diabetes mellitus, hypercholesterolemia, hemodialysis treatment, donor type, history with myocardial infarction or cerebrovascular accident (CVA)/transient ischemic attack (TIA). The following outcomes were considered for meta-analysis when compared between patients with and without VC: patient survival, uncensored graft survival, death-censored graft survival, DGF, and kidney function. Events for survival outcomes were deduced from Kaplan-Meier survival curves using DataThief software and from numbers and percentages described in the results section [18].

Statistical analysis

For meta-analysis, pooled risk ratios (RR) with 95% confidence intervals (CI) were calculated at fixed time spans based on the number of events per group as described in the individual studies. Because of the expected observational designs of included studies resulting in high between-study variance, a random-effect model was used as described by DerSimonian and Laird [19]. The Mantel-Haenszel analysis method was used with calculation of the overall effect using the Z-test. To investigate potential confounders, baseline characteristics were collected of

included patients without VC and with any VC. For continuous variables, the group mean weighed for number of included patients was reported with pooled standard error, if the included study reported the mean. Normality of the means of the included studies was assumed because of the sample size, according to the central limit theorem. Therefore, baseline characteristics were compared with the unpaired *T*-test in case of continuous variables and with chi-square test for categorical variables using MED-CALC software (version 16.2). Statistical heterogeneity was visually assessed by judging overlap in the 95% confidence intervals and with I^2 . Publication bias was assessed using funnel plots of the logarithm of RR versus their standard errors, which are included in the Supplemental Digital Content [20]. A *P*-value below 0.05 was considered statistically significant. The program used for meta-analysis was REVIEW MANAGER 5.3 [21].

Results

Study selection and characteristics

A total number of 1523 potentially relevant, observational studies were identified. None of the studies describing iliac

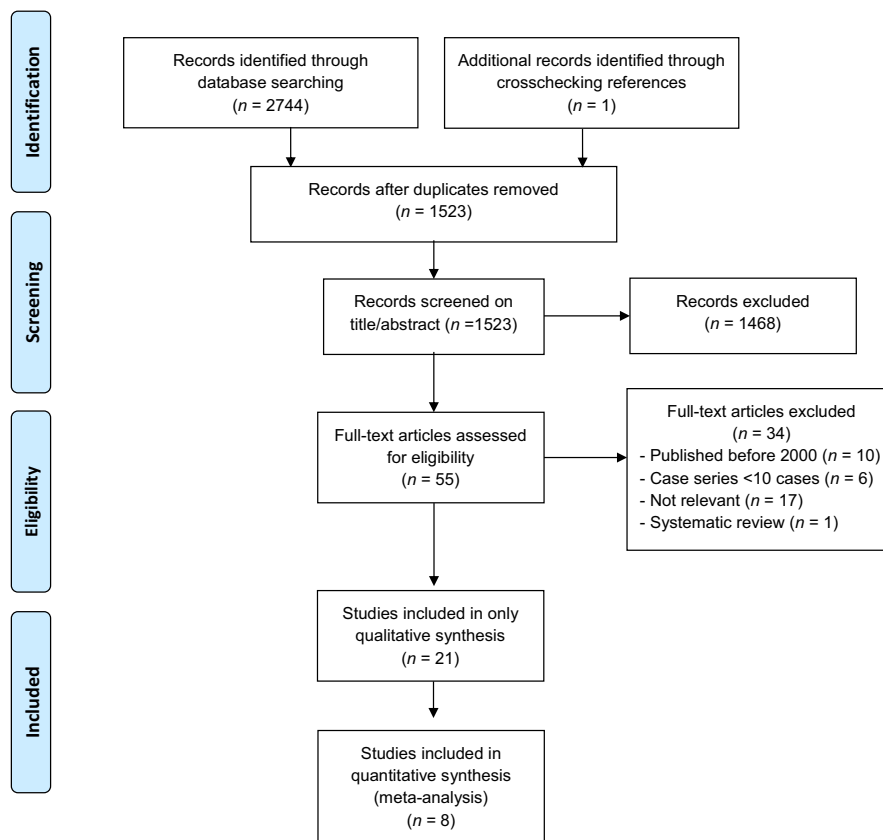


Figure 1 Preferred reporting items for systematic reviews and meta-analyses flow diagram of the systematic literature search.

Table 1. Studies providing data for meta-analysis ($n = 8$).

Study	Year	Design	nVC	VC	Diagnosis of VC	Severity VC	Location	Newcastle-Ottawa scale				Quality
								Selection	Comparability	Outcome	Outcome	
Hernandez Spain	2005	R OBS	844	273	Pelvic X-ray	No VC Any VC	IA DA	1,5	4	2	2 Median FU 49 months, lost to FU unknown	Good
Droupy France	2006	P OBS	1001	69	Palpable during transplantation	No VC Any VC	IA	1,3,4	4	0	0 FU unknown, lost to FU unknown	Poor
Aalten Netherlands	2011	P OBS	74	35	Palpable during transplantation	No VC Any VC	IA	1,2,4,5	4	0	0 FU 1 year, lost to FU unknown	Poor
Aitken Scotland	2012	P OBS	61	32	Pelvic X-ray	Minimal VC Moderate/severe VC	IA	2,4,5	4	0	3 Complete follow-up of 5 years for all patients	Poor
Munguia Spain	2015	R OBS	69	50	Pelvic X-ray, usage of Kauppila index, L4-S1	No VC (KI 0–2) Any VC (KI 3–24)	DA	1,2,3,4,5	4	0	0 FU unknown, lost to FU unknown	Poor
Davis United states	2016	R OBS	41	90	CT-scan with calcium score	No VC Any VC	IA	1	4	0	0 Mean FU 34.1 months, lost to FU unknown	Poor
Benjamins Netherlands	2018	R OBS	434	267	DXA L1-L4 with score using Schousboe method	No VC Any VC	DA	1	4	2	3 Median FU 5.4 years, lost to FU unknown	Good
Disthabanchong Thailand	2018	P OBS	108	26	Pelvic X-ray, usage of Kauppila index	VC ≤ 1 VC > 1	IA	1	4	2	2 Median FU 62.5 months, lost to FU unknown	Good
Total			2632	842	* Outcomes: 1, patient survival; 2, uncensored graft survival; 3, death-censored graft survival; 4, delayed graft function; 5, kidney function							

Outcomes: 1, patient survival; 2, uncensored graft survival; 3, death-censored graft survival; 4, delayed graft function; 5, kidney function.

AAAC, abdominal aortic calcification; CT, computed tomography; DA, distal aorta; DXA, dual-energy X-ray absorptiometry; FU, follow-up; IA, iliac arteries; KI, Kauppila index; MR, magnetic resonance angiography; nVC, no vascular calcification; P OBS, prospective observational; R OBS, retrospective observational; VC, vascular calcification.

AAC, abdominal aortic calcification; CT, computed tomography; DA, distal aorta; DXA, dual-energy X-ray absorptiometry; FU, follow-up; IA, iliac arteries; KI, Kauppila index; MRA, magnetic resonance angiography; nVC, no vascular calcification; P OBS, prospective observational; R OBS, retrospective observational; VC, vascular calcification. Outcomes: 1, patient survival; 2, uncensored graft survival; 3, death-censored graft survival; 4, delayed graft function; 5, kidney function.

calcifications distinguished between calcifications of the common iliac arteries or external iliac arteries. Figure 1 presents the PRISMA flow diagram. Twenty-one studies met the inclusion criteria from which one was added after manual reference check. Eight studies provided data for meta-analysis. Characteristics of studies included in meta-analysis are presented in Table 1 and from the studies that did not provide data for meta-analysis in Table 2. The baseline characteristics of the patients included in meta-analysis are shown in Table 3. Patients with any degree of VC were older [no vascular calcification (nVC: 42.0 ± 12.4 , any VC: 54.3 ± 10.8 , $P < 0.001$), were more frequently suffering from hypertension (nVC 75.8%, any VC 86.0%, $P < 0.001$), diabetes mellitus (nVC 10.5%, any VC 32.4%, $P < 0.001$) and hypercholesterolemia (nVC 27.9%, any VC 44.3%, $P < 0.001$). Patients with any VC had more often a history with a myocardial infarction (nVC 6.7%, any VC 14.2%, $P < 0.001$) and received more frequently a living donor kidney transplant (nVC 2.1%, any VC 5.3%, $P < 0.001$). Patients with any VC were more often preemptively transplanted (nVC 7.3%, any VC 16.3%, $P < 0.001$).

Patient survival

Seven studies provided data for meta-analysis. The forest plots of the comparisons made for patient survival are shown in Fig. 2. All seven studies investigated 1-year patient survival and showed an increased mortality risk in patients with any VC with a pooled risk ratio (RR) of 2.19 ($n = 3381$; 95% CI 1.39–3.44; $P < 0.001$) [1,22–27]. Six studies also noted the risk of 3-year mortality which was also increased in recipients with any VC ($n = 3272$; RR: 2.17; 95% CI 1.66–2.83; $P < 0.001$) [1,23–27]. Five studies mentioned 5-year mortality risk. Similar to the pooled results of 1-year and 3-year survival, a significantly increased 5-year mortality risk was shown in patients with any degree of VC ($n = 3153$; RR: 2.28; 95% CI 1.86–2.79; $P < 0.001$) [1,23–25,27]. Two eligible studies were not suitable for meta-analysis. Chavent et al. divided patients into four quartiles, based on a CT-scan based calcification score. No difference was found for patient survival between quartiles after a mean follow-up of 4.18 ± 1.64 years [28]. Rijkse et al. [29] investigated the impact of aorto-iliac stenosis classified with the TASC II classification within a retrospective cohort study. They found a significantly decreased patient survival in patients with TASC II C/D lesions with a 5-year survival of 66% in the TASC II A/B group, 26% in the TASC II C/D group and 72% in the control group without stenosis (log-rank test TASC II A/B: $P = 0.078$, TASC II C/D: $P < 0.001$). After

adjustment for various confounders, having a TASC II C/D lesion was a strong, independent risk factor for mortality (HR 3.25; 95% CI 1.87–5.67; $P < 0.001$) [29]. Four studies also investigated causes of death between patients with any degree of VC and no VC. Droupy et al. found that death from cardiovascular cause was more frequent in patients with any VC (nVC: 2.7%, any VC: 27%, $P < 0.001$) [25]. Also, Hernandez et al. found that death from a cardiovascular cause was more frequent among recipients with any VC (nVC: 3.1%, any VC: 9.5%) [1]. Munguia et al. investigated the combined outcome of a Major Cardiovascular Event (MACE) and cardiovascular death. They found a statistical significant difference with an incidence of 6.5% in KTR without VC and 21.7% in KTR with any VC ($P = 0.035$) [26]. Rijkse et al. found that, among deceased patients, death from a cardiovascular cause was more frequent in patients with any TASC II lesion (any TASC II lesion: 35.4%, no TASC II lesion 19.1%, $P = 0.035$) [29].

Uncensored graft survival

Four studies, from which three provided data for meta-analysis, reported uncensored graft survival. Figure 3 shows the pooled results of those studies. The risk of one-year graft loss was three times higher in KTR with any VC ($n = 321$; RR: 3.15; 95% CI 1.30–7.64; $P = 0.01$) [22,26,30]. The risk of 3-year graft loss was investigated in two studies. The pooled RR showed no significant higher risk of graft failure in KTR with VC ($n = 212$; RR: 3.41; 95% CI 0.97–11.96; $P = 0.05$) [26,30]. Rijkse et al. [29] also found a significant graft survival difference between KTR with a TASC II C/D lesion in comparison to KTR without any TASC II lesions (5-year graft survival: no TASC II lesions: 60%, TASC II C/D lesion: 22%, log-rank test: $P < 0.001$).

Death-censored graft survival

Five studies reported death-censored graft survival in recipients with any VC without an additional vascular procedure. Only two provided data for meta-analysis. Figure 4 shows the forest plots with pooled RR. The risk of death-censored graft loss was not significantly different between KTR with and without VC with a RR of 2.26 ($n = 1189$; 95% CI 0.58–8.82; $P = 0.24$) [25,26]. Also, the risk of 3-year death-censored graft loss was statistically similar ($n = 1189$; RR: 2.19; 95% CI 0.49–9.82; $P = 0.31$) [25,26]. Three eligible studies were not suitable for meta-analysis. Chavent et al. reports no difference in overall death-censored graft survival after a

Table 2. Included studies not suitable for meta-analysis ($n = 13$).

Study	Year	Design	Study size	Study population	Treatment	Outcome
Galazka Poland	2002	R OBS	$N = 128$ (54 pairs): Group A: $N = 54$ with VC Group B: $N = 54$ without VC	Pairs of recipients who received a kidney from the same donor (1 recipient with VC, 1 without VC) Donor: 0% LD, age: 48 Recipients with TASC II C/D lesions Donor: 0% LD, age: 54 \pm 9 Recipients who received a kidney transplant on a vascular bypass Donor: 91% LD, age: 57.7 \pm 13.9 Recipients who received aortiliac surgery simultaneously with KTx Age: 55 (43–65)	Group A: $N = 36$ EAT, $N = 16$ anastomosis to EIA, $N = 2$ anastomosis to CIA Group B: Anastomosis on AII	1,2
Tozzi Italy	2013	P OBS	$N = 21$		$N = 15$ simultaneous EAT $N = 4$ aorto-bi-iliac bypass $N = 9$: reconstruction with ileo-femoral bypass, $N = 1$ femoroiliac crossover bypass, $N = 1$ aortofemoral bypass	1,2,4,6
Ozcelik, Germany	2007	R OBS	$N = 11$		$N = 15$ EAT, $N = 2$ aorto-iliac bypass, $N = 5$ aorto-bi-iliac bypass, $N = 1$ aorto-bifemoral bypass, $N = 4$ arterioplasty, $N = 3$ iliac-iliac bypass	1,2,4,6
Tsivian Italy	2009	R OBS	$N = 30$ ($N = 19$ stenotic lesions, $N = 11$ aneurysms)		$N = 27$ with angioplasty ($N = 2$ PTFE reconstruction, $N = 25$ EAT)	1,2,5
Patrono, Belgium	2013	R OBS	$N = 27$ $N = 32$ from literature	Recipients who received a kidney transplant on a prosthetic graft Donor: 7.4% LD, age: 56 (35–75) Recipients with asymptomatic TASC II A/B requiring angioplasty Donor: 100% LD, age: 51.4 \pm 9.1	$N = 24$ implantations on prosthetic graft ($N = 22$ graft before KTx, $N = 2$ simultaneously, $N = 3$ after KTx)	1,2,5,6
Han Korea	2014	R OBS	$N = 748$	Recipients transplanted with usage of a vascular conduit because of severe VC Donor: 20% LD, age: 61 \pm 9 Recipients who gave consent to provide an iliac artery specimen during KTx Donor: 78% LD, age: 42.5 \pm 10.3	$N = 8$ donor iliac artery graft $N = 2$ saphenous vein graft	1,2,3,4,6
Coleman USA	2014	R OBS	$N = 10$		NA	1,2,4,5,6
Hwang Korea	2015	P OBS	$N = 90$ ($N = 48$ positive intimal microcalcification, $N = 42$ negative intimal microcalcification)			3,4
Sagban Germany	2016	R OBS	$N = 208$ $N = 121$ from literature	Recipients who received a kidney transplant on a prosthetic graft Donor: 0% LD, age: 56 Recipients predicted to have complications with the arterial anastomosis because of VC Donor: 92.3% LD, age: 60.2 \pm 10.4, 0% pre-emptive	$N = 4$ anastomosis on a prosthetic graft $N = 121$ anastomosis on a prosthetic graft from the literature $N = 13$ EAT before anastomosis ($N = 10$ anastomosis on EIA, $N = 1$ on CIA, $N = 1$ on IIA, $N = 1$ on prosthetic graft)	1,2,5,6
Nanmoku Japan	2017	R OBS	$N = 13$			1,2,3,6
Chavent France	2017	R OBS	$N = 100$ (divided in quartiles, based on calcium score)	Recipients with non-contrast enhanced abdominal CT-scan. Donor: 5% LD, age: 60.3 \pm 12.8, 5% pre-emptive	NA	1,3,5
Franquet France	2018	R OBS	$N = 11$	Recipients who received a kidney transplant on a vascular bypass Age: 61 (57–62.5)	$N = 9$ aorto-femoral bypass $N = 1$ aorto-bifemoral bypass $N = 1$ aorto-iliac bypass	1,2,6

Table 2. Continued.

Study	Year	Design	Study size	Study population	Treatment	Outcome
Rijkse Netherlands	2019	R OBS	$N = 374$ ($n = 88$ with TASC II lesions, $n = 286$ without TASC II lesions)	Recipients with aorto-iliac stenosis compared to recipients without stenosis Donor: 61% LD, age: 59.6 ± 12.7 , 17% pre-emptive	$N = 3$ aortic bifurcation prosthesis, $N = 1$ iliac femoral bypass, $N = 2$ iliac endarterectomy with patch angioplasty, $N = 2$ Gore-Tex angioplasty	1,2,3,6

CIA, common iliac artery; CT, computed tomography; EAT, endarterectomy; EIA, external iliac artery; IIA, internal iliac artery; KTx, kidney transplantation; LD, living donor; NA, not applicable; P OBS, prospective observational; PTFE, polytetrafluoroethylene; R OBS, retrospective observational; TASC, Trans-Atlantic inter-Society Consensus.

Outcomes: 1, patient survival; 2, uncensored graft survival; 3, death-censored graft survival; 4, delayed graft function; 5, kidney function; 6, postoperative complications.

mean follow-up of 4.18 ± 1.64 years ($P = 0.7$) [28]. Hwang et al. [31] found a significant difference in overall death-censored graft survival between patients with positive intimal calcification in comparison with patients with negative intimal micro-calcification (log-rank test $P = 0.017$). Rijkse et al. [29] found no significant association between the presence of aorto-iliac stenosis as classified with the TASC II classification and death-censored graft loss [TASC II A/B: HR 0.78 (0.41–1.50), TASC II C/D: HR 1.85 (0.74–4.65)].

Delayed graft function

Out of five studies reporting the incidence of DGF, four provided data for meta-analysis. The pooled RR, as shown in the forest plot depicted in Fig. 5, showed no statistical significant difference for risk of DGF ($n = 1391$; RR: 1.24; 95% CI 0.98–1.58; $P = 0.08$) [22,25,26,30]. Hwang et al. [31] found no significant difference in the incidence of DGF between patients with and without intimal microcalcification (22.9% vs. 21.4%, $P = 0.204$).

Kidney function

Five studies investigated the impact of VC on 1-year creatinine. Due to the large heterogeneity between studies (I^2 of 81%), it was decided not to pool the results. Aalten et al. [22] found a mean serum creatinine of 130 ± 38 $\mu\text{mol/l}$ in KTR with VC and 131 ± 41 $\mu\text{mol/l}$ in KTR without VC, which was not statistically different. The study of Aitken et al. also found no statistical significant difference (VC: 148.5 ± 9.6 , nVC: 140.3 ± 8.9) [30]. Munguia et al. investigated serum creatinine after 1 month (nVC: 1.90 ± 0.13 mg/dl, VC: 1.79 ± 0.10 mg/dl), 3 months (nVC: 1.63 ± 0.06 mg/dl, VC: 1.69 ± 0.10 mg/dl), and 1 year (nVC: 1.57 ± 0.07 mg/dl, VC: 1.55 ± 0.11 mg/dl). No significant difference was found [26]. Hernandez et al. investigated the percentage of patients with a serum creatinine >2 mg/dl at discharge. He found that the proportions were not significantly different (VC 23.4%, nVC 27.3%, P -value 0.248) [1]. Chavent et al. found that creatinine levels were significantly higher at last-follow-up (mean 4.18 ± 1.64) in the fourth quartile with the most severe calcification ($P = 0.046$), but this result was not significant when the glomerular filtration rate was calculated with the MDRD formula ($P = 0.1$) [28].

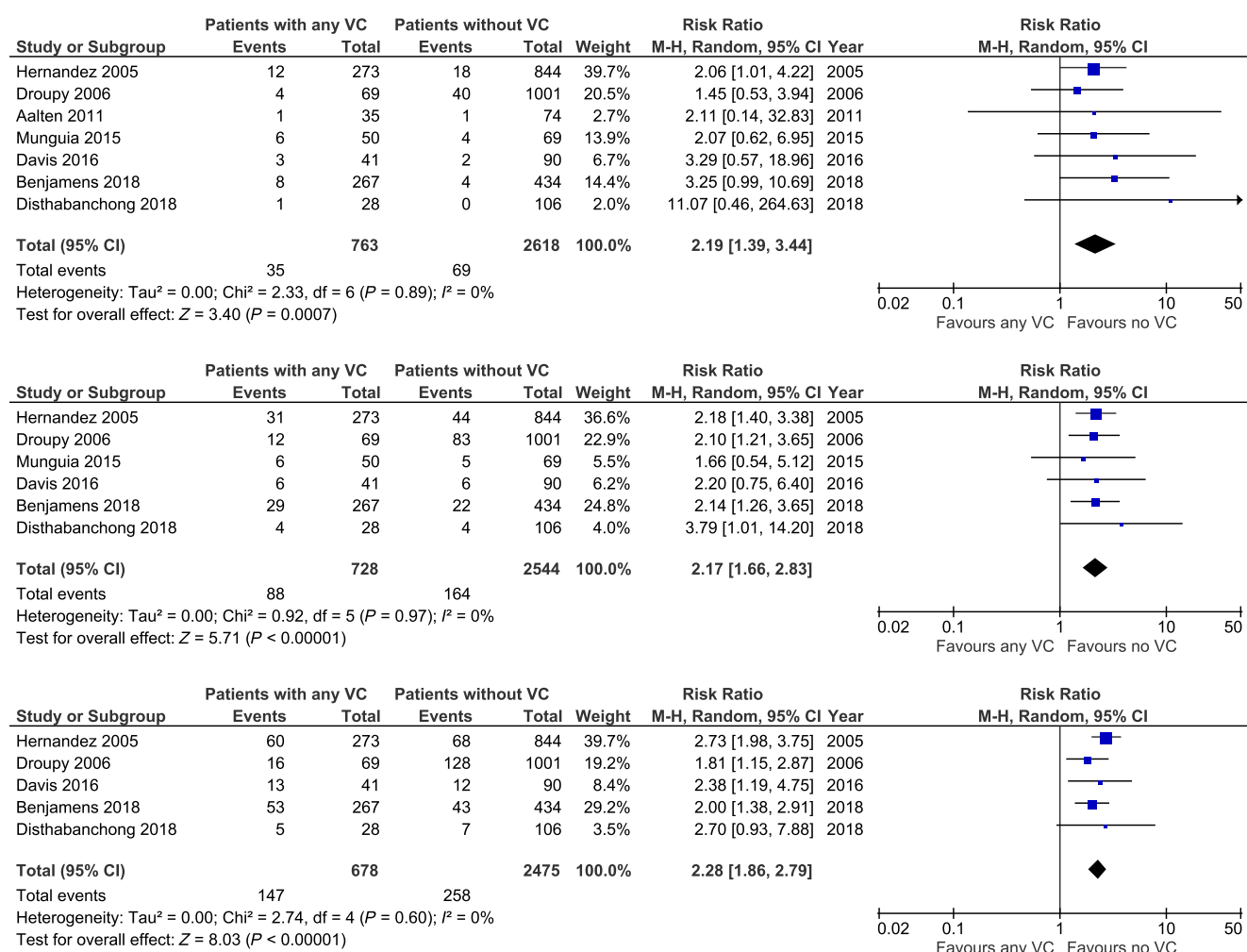
KTx on a prosthetic graft

Nine studies were published in which kidney transplants were connected to a prosthetic graft. In three

Table 3. Baseline characteristics of patients included in meta-analysis.

Characteristics	Studies	nVC	Total patients	Any VC	Total patients	P-value
Recipient age, mean (SD)	6 ^{1,22,23,25,26,27}	42.0 (12.4)	2529	54.3 (10.8)	721	<0.001*
Male sex, <i>n</i> (%)	5 ^{1,22,23,26,27}	977 (63.9)	1528	421 (64.6)	652	0.779
Smoking, <i>n</i> (%)	2 ^{22,23}	99 (19.5)	508	59 (19.5)	302	<0.986
Hypertension, <i>n</i> (%)	4 ^{1,22,23,27}	1106 (75.8)	1459	518 (86.0)	602	<0.001*
DM, <i>n</i> (%)	5 ^{1,22,23,26,27}	160 (10.5)	1528	211 (32.4)	652	<0.001*
Hypercholesterolemia, <i>n</i> (%)	3 ^{1,23,27}	387 (27.9)	1385	251 (44.3)	567	<0.001*
Pre-emptive KTx, <i>n</i> (%)	5 ^{1,22,23,25,26}	176 (7.3)	2422	113 (16.3)	694	<0.001*
Living donor, <i>n</i> (%)	3 ^{1,22,25}	41 (2.1)	1919	20 (5.3)	377	<0.001*
History MI, <i>n</i> (%)	2 ^{22,23}	34 (6.7)	508	43 (14.2)	302	<0.001*
History CVA/TIA, <i>n</i> (%)	2 ^{22,23}	28 (5.5)	508	27 (8.9)	302	0.061

CVA, cerebrovascular accident; DM, diabetes mellitus; KTx, kidney transplantation; MI, myocardial infarction; nVC, no vascular calcification; SD, standard deviation; TIA, transient ischemic attack; VC, vascular calcification. * indicates statistical significance.

**Figure 2** Risk of 1-, 3- and 5-year mortality in recipients with any degree of vascular calcification (VC) and without VC.

studies, numbers were too small for further analysis [32–34]. One study did not describe the results of those patients separately [29]. Table 4 presents the eligible studies. A total number of 57 cases were

described in which the kidney transplant was connected to a prosthetic graft [35–39]. The incidence of DGF was described in 53 patients from which 4 (7.5%) had a DGF [35–37,39]. The incidence of

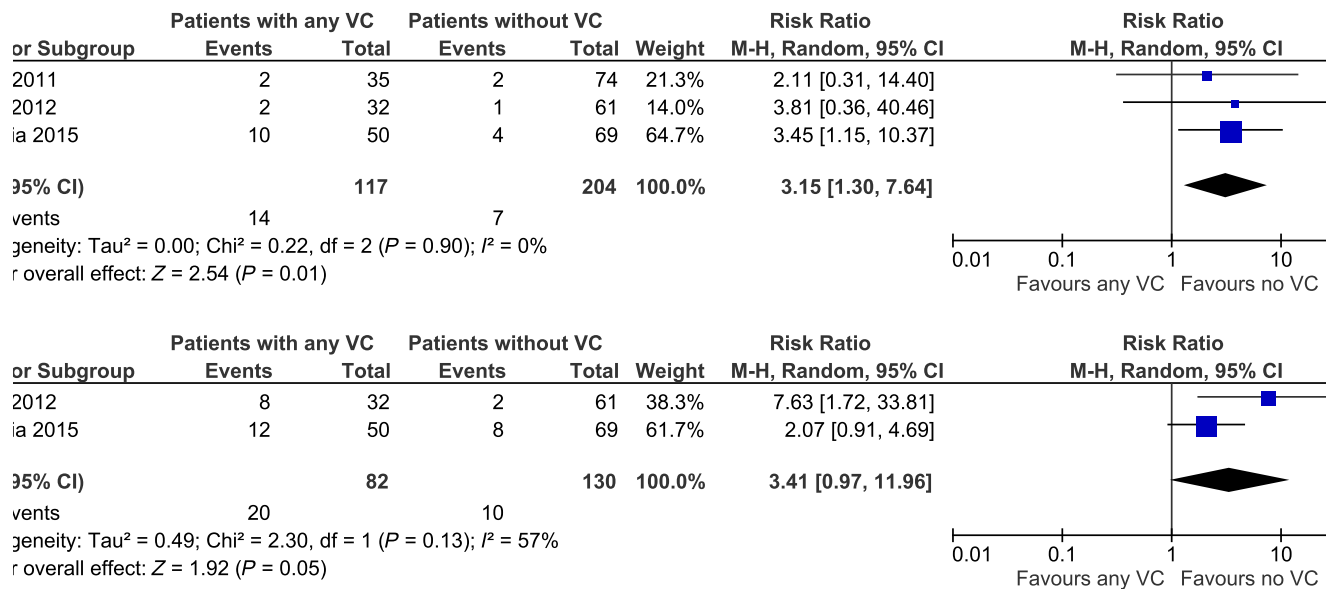


Figure 3 Risk of 1- and 3-year graft loss uncensored for death in recipients with any degree of vascular calcification (VC) and without VC.

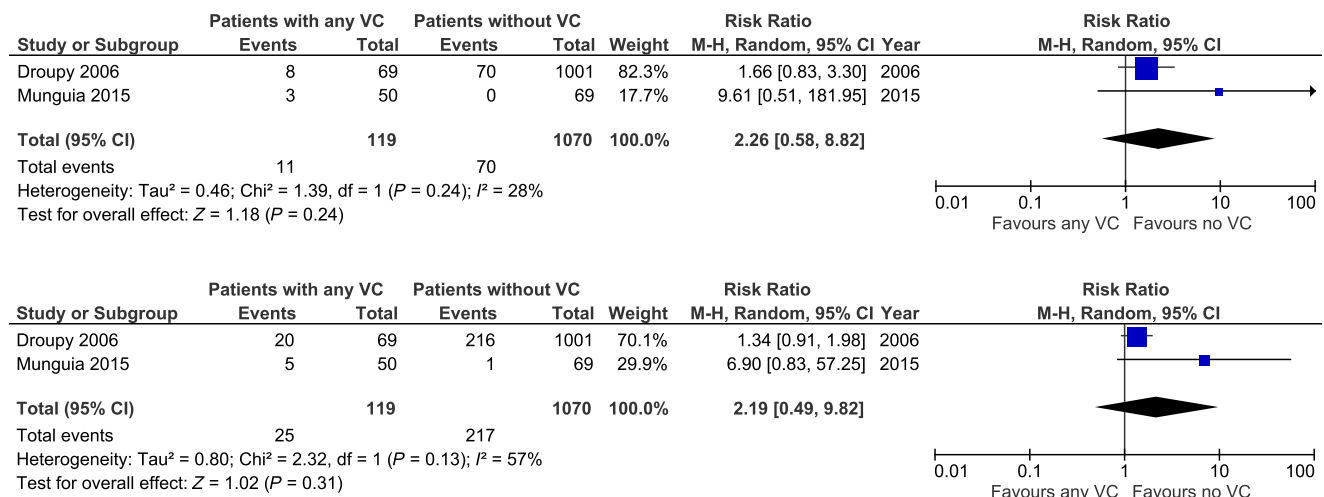


Figure 4 Risk of 1- and 3-year death-censored graft loss in recipients with any degree of vascular calcification (VC) and without VC.

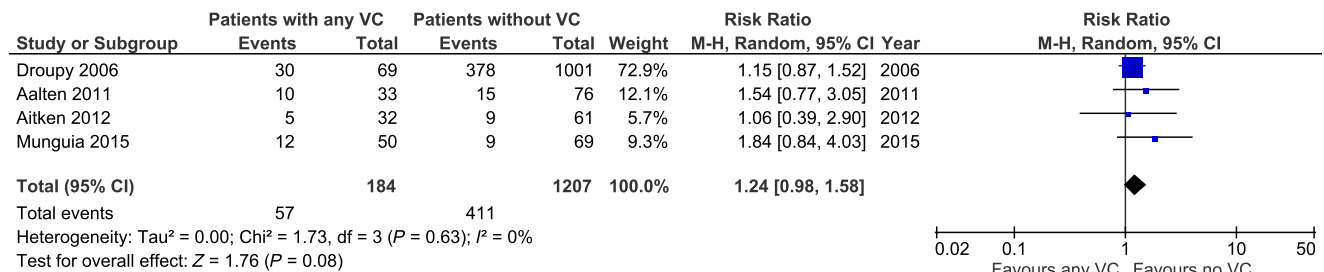


Figure 5 Risk of delayed graft function in recipients with any degree of vascular calcification (VC) and without VC.

postoperative complications was 19.3% and complications described were renal vein thrombosis ($n = 1$), bleeding ($n = 5$), rejection ($n = 1$), lower limb amputation ($n = 1$), thrombosis below the graft ($n = 1$), infection ($n = 2$), and surgical wound dehiscence ($n = 1$) [35–39]. A re-operation was needed in

Table 4. Outcomes of KTx on a vascular bypass.

Study	N	DGF (%)	Re-operation (%)	Patient survival (%)				Graft survival DC (%)				Uncensored graft survival (%)				
				EC (%)	30 days	1 year	3 years	5 years	30 days	1 year	3 years	5 years	30 days	1 year	3 years	5 years
Ozcelik Germany	2007	11	2/11 (18.2)	3/11 (27.3)	1/11 (9.1)	NA	NA	NA	8/11 (72.7)	NA	NA	NA	8/11 (72.7)	NA	NA	NA
Patrono Belgium	2013	27	2/27 (7.4)	5/27 (18.5)	1/27 (3.7)	NA	24/27 (88.9)	23/27 (85.2)	26/27 (96.3)	26/27 (96.3)	24/27 (88.9)	NA	24/27 (88.9)	22/27 (81.5)	20/27 (74.1)	
Tozzi, Italy	2013	4	0/4 (0)	0/4 (0)	0/4 (0)	4/4 (100)	4/4 (100)	NA	4/4 (100)	4/4 (100)	NA	4/4 (100)	4/4 (100)	NA	NA	
Sagban German	2016	4	NA	1/4 (25)	1/4 (25)	4/4 (100)	4/4 (100)	NA	4/4 (100)	4/4 (100)	4/4 (100)	4/4 (100)	4/4 (100)	NA	NA	
Franquet France	2018	11	0/11 (0)	2/11 (18.2)	1/11 (9.1)	11/11 (100)	11/11 (100)	9/11 (81.8)	9/11 (81.8)	9/11 (81.8)	NA	11/11 (100)	9/11 (81.8)	NA	NA	
Total		57	4/53 (7.5)	11/57 (19.3)	4/57 (7.0)	19/19 (100)	43/46 (93.5)	32/38 (84.2)	43/46 (93.5)	30/31 (96.8)	28/31 (90.3)	27/30 (90.0)	41/46 (89.1)	22/27 (81.5)	20/27 (74.1)	

DC, death-censored; DGF, delayed graft function; EC, early complications; NA, not applicable.

7.0% of the recipients [35–39]. Thirty-day patient, death-censored and uncensored graft survival was 100%, 90.0% and 90% respectively [35,36,38,39]. One-year patient survival, uncensored and death-censored graft survival was 93.5%, 93.5% and 89.1% respectively [35,37–39]. Five-year survival outcomes were only mentioned in one study. In this study, 5-year patient survival, death-censored and uncensored graft survival was 85.2%, 90.3% and 74.1% respectively [37]. Coleman et al. described the usage of a vascular conduit in 10 patients ($n = 8$ donor iliac artery graft, $n = 2$ saphenous vein graft) to facilitate KTx in case of limited anastomotic options due to iliac calcification [40]. No postoperative mortality or graft loss was observed. Two patients had a fascial dehiscence as a complication, and two patients had DGF [40].

KTx after endarterectomy

Six studies discussed clinical outcomes after KTx in patients who underwent simultaneous iliac artery endarterectomy. All of them mentioned 1-year patient survival, as shown in Table 5. One-year patient survival varied from 86.6% till 100% in patients who underwent endarterectomy (EAT) [25,32–34,39,41]. Three studies used a control group of patients without VC and they all showed no statistical significant difference for 1-year patient survival [25,32,41]. Three studies also investigated 5-year patient survival, from which one showed a significant difference in favor of patients without VC (nVC 87 ± 1 , VC + EAT 69 ± 8 , $P < 0.001$) [25]. Five studies noted 1-year uncensored graft survival, which varied from 80% to 100% [32–34,39,41]. Two studies compared the results with KTR without VC, which was not statistically different [32,41]. One-year death-censored graft survival was investigated in three studies and varied from 87–100%, from which two studies used a control group of recipients without VC [33,34,39]. Droupy et al. found no significant difference for 1-year death-censored survival, but 5-year death-censored graft survival was inferior in recipients who underwent EAT (70 ± 2 , 46 ± 7 , $P < 0.001$) [25]. In the study from Han et al., 1- and 5-year death-censored graft survival was equal between both groups [32]. The incidence of DGF was investigated in two studies. Droupy et al. found no difference in the incidence of DGF between KTR without VC and KTR with VC + EAT (37%, 42%, NS) [25]. Han et al. [32] confirms these results with an incidence of 2.0% in the KTR without VC group and 7.4% in the KTR with VC + EAT group ($P = 0.099$).

Discussion

Our meta-analysis demonstrated that the presence of any VC is associated with an increased mortality risk and risk of 1-year graft loss. This is in line with our expectations, because there is a strong association between large-vessel peripheral arterial disease and cardiovascular mortality [42]. Indeed, the incidence of cardiovascular death was more frequent in deceased KTR with any degree of VC. As expected, risk factors for vascular disease were more prevalent in patients with any degree of VC. Our meta-analysis did not demonstrate a statistical significant difference for risk of death-censored graft loss or DGF. However, the pooled RR's for these outcomes and the wide confidence intervals suggest that this may be attributable to the small sample size. Studies describing results from KTx on a prosthetic graft or after endarterectomy were scarce and results varied largely. Patients who died after bypass surgery and did therefore not receive a kidney transplant are

Table 5. Patient and graft survival in kidney transplant recipients who underwent endarterectomy.

Study	Year	Patient survival (%)				Uncensored graft survival (%)				Death-censored graft survival (%)			
		1-year		5-year		1-year		5-year		1-year		5-year	
		nVC	EAT	P-value	nVC	EAT	P-value	nVC	EAT	nVC	EAT	nVC	P-value
Galazka Poland	2002	94	98	>0.05	92	96	>0.05	87	89	>0.05	66	68	>0.05
Droupy France	2006	96	97.4	>0.05	87 ± 1	69 ± 8	<0.001	NA	NA	NA	NA	NA	>0.05
Tsivian Italy	2009	NA	86.6	NA	NA	NA	NA	NA	80	NA	NA	NA	NA
Tozzi Italy	2013	NA	100	NA	NA	NA	NA	NA	100	NA	NA	100	NA
Han Korea	2014	97.7	96.3	>0.05	96.3	96.3	>0.05	96.9	96.3	0.463	94.5	86.7	0.463
Nannmoku Japan	2017	NA	100	NA	NA	NA	NA	NA	100	NA	NA	100	NA

EAT, endarterectomy; NA, not applicable; nVC, without vascular calcification; VC, vascular calcification.

not taken into account, creating guarantee-time bias. As a result, survival outcomes of kidney transplantation on a prosthetic graft might be too optimistic.

Our meta-analysis has some important limitations. First, all studies were observational, which means that there is much confounding we could not correct for. We compared baseline characteristics to give an indication of the existing confounders. Confounders known to increase the mortality risk were more prevalent in KTR with any degree of VC. Factors associated with better graft survival, such as a living donor transplant and preemptive transplantation, were also more frequent in KTR with any degree of VC. This may be due to possible selection bias of the transplant surgeon to improve outcomes and to guarantee daytime surgery. Another limitation is instrument variability because of the different methods used to diagnose VC. In two studies, VC was diagnosed during surgery, which may lead to underdetection. Other studies used an X-ray or CT-scan to diagnose VC. According to Aitken et al. [30], sensitivity and specificity for pelvic X-ray to diagnose VC in comparison to CTA were 98.5% and 92.6% respectively. Both pelvic X-ray (sensitivity 95.5%, specificity 83.1%) and CTA (sensitivity 100%, specificity 92%) correlated well with intraoperative assessment of the vessels [30]. Therefore, we think that the usage of different imaging modalities did not largely affect our results. Because of the different scoring systems used and the different methods to diagnose VC, we could not investigate a dose-response relationship. Therefore, we decided to use the dichotomous outcome of no VC/ any degree of VC to decrease misclassification. The studies in our meta-analysis did not all describe if patients with any VC who received pre-transplant PTA were excluded. If patients after PTA were included in the study, graft survival estimates could have been diluted towards the null when compared to untreated VC. Also, no distinction could be made between VC and hemodynamically significant stenosis. It may be possible that VC only impacts death-censored graft survival outcomes in case of a hemodynamically significant stenosis. Rijkse et al. investigated this and did not find a statistical significant difference [TASC II A/B: HR 0.78 (0.41–1.50), TASC II C/D: HR 1.85 (0.74–4.65)] [29]. However, in this study, the sample size of KTR with significant stenosis was small. Larger studies are needed to provide definite answers to this question.

This meta-analysis is the first one to describe the overall prognosis of KTR with any degree of VC.

Besides the observational study designs, low between study heterogeneity was observed. We were able to show important factors associated with the less optimistic prognosis of KTR with any degree of VC. For this meta-analysis, we were only able to look at the dichotomous outcome of any VC/ no VC. Future studies should focus on finding a dose-response effect of the amount of VC. Also, a standardized classification should be used to reduce heterogeneity between studies and to allow risk stratification. Even though the study designs were not optimal for meta-analysis, we carefully selected studies to include in our meta-analysis and therefore, we believe this is the best available evidence on this subject. Because of all confounding factors, we could not investigate whether VC is an independent risk factor for mortality and graft loss. The value of VC as an independent risk factor should be investigated for usage in post-transplant risk adjustment models, such as the Scientific Registry of Transplant Recipients (SRTR) post-transplant risk adjustment models [43].

Conclusion

The presence of VC in KTR is associated with an increased mortality risk and increased risk of graft loss. No statistical significant association between VC and DGF or risk of death-censored graft loss could be demonstrated. However, for the interpretation of the outcomes, the quality, risk of bias and sample size of the available evidence should be taken into consideration.

Authorship

ER, RCM and HJANK: participated in the research design. ER and JLD: screened all studies, selected the studies to be included in this systematic review and analyzed the data. ER, JLD and RCM: wrote the article. JLD, JNMII, RCM, JIR and HJANK: revised the manuscript critically.

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Conflict of interest

The authors declare that they do not have any conflicts of interest to disclose.

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SUPPORTING INFORMATION

Additional supporting information may be found online in the Supporting Information section at the end of the article.

Table S1. Search terms used for this systematic review and meta-analysis.

Table S2. Summary of findings table for studies included in meta-analysis.

Figure S1. Funnel plot for outcome: 1-year mortality.

Figure S2. Funnel plot for outcome: 1-year uncensored graft loss.

Figure S3. Funnel plot for outcome: 1-year death-censored graft loss.

Figure S4. Funnel plot for outcome: delayed graft function.

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