

Disparities in early mortality among chronic kidney disease patients who transition to peritoneal dialysis and hemodialysis with and without catheters

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Abstract

Purpose The early period after chronic kidney disease (CKD) patients transition to end-stage renal disease (ESRD) represents the highest mortality risk but is variable among different patient populations and clinical circumstances. We compared early mortality outcomes among a diverse CKD population that transitioned to ESRD.

Methods A retrospective cohort study (1/1/2002 through 12/31/2013) of CKD patients (age \geq 18 years) who transitioned to peritoneal dialysis (PD), hemodialysis (HD) with arteriovenous fistula/grafts, and HD with catheters was performed. Multivariable Cox regression modeling was used to estimate 6-month all-cause mortality hazard ratios (HR) among the three treatment groups after adjustment for patient and clinical characteristics.

Results Among 5373 ESRD patients (62.7 years, 41.3% females, 37.5% Hispanics, 13.3% PD, 34.9% HD with fistula/graft, 51.8% HD with catheter), 551 (10.3%) died at 6 months. Mortality rates were highest immediately after transition (299 deaths per 1000 person-years in first month). Compared to PD patients, the 6-month mortality HR (95% CI) was 1.87 (1.06–3.30) in HD with fistula/graft patients and 3.77 (2.17–6.57) in HD with catheter patients. Inpatient transition (HR 1.32), acute kidney injury (HR 2.06), and an eGFR \geq 15 vs 5–9 (HR 1.68) at transition were also associated with higher early mortality risk. **Conclusion** Among a diverse CKD population who transitioned to ESRD, we observed considerable differences in early mortality risk among PD, HD with fistula/graft, and HD with catheter patients. The identification of patient-specific and clinical environmental factors related to high early mortality may provide insights for managing advanced stages of CKD

and shared decision making.

Keywords Mortality · End-stage renal disease transition · Chronic kidney disease · Epidemiology · Comparative outcomes

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Introduction

While the end-stage renal disease (ESRD) population itself experiences high mortality, the early period after transition to ESRD appears to be most vulnerable. High mortality rates of up to 30% have been described within the first year of transition from chronic kidney disease (CKD) to ESRD [1–6]. While the evidence is not entirely consistent, studies have described a mortality spike within the first few weeks to months of ESRD transition while others have not [3, 7–9]. This alarmingly high rate of mortality has been the impetus to one of the Healthy People 2020 goals to reduce ESRD mortality to 329 deaths per 1000 patient years at risk within the first 3 months of ESRD transition [10].

A better understanding of the late CKD and early ESRD populations could provide insight into the reasons for this

high transition mortality. It could have important implications for prognostication and help guide management strategies and shared decision making. An important consideration is to identify the patients that pose risk for early mortality as they may represent different populations and circumstances altogether. The identification of these populations earlier in the course of CKD could help inform treatment plans and manage expectations through shared decision making.

Currently, there are limitations to available information on late-stage CKD and early ESRD outcomes. Large data registries such as the US Renal Data System (USRDS) historically have not comprehensively captured the early period after transition, due to a 3-month gap or an ascertainment bias [2, 11]. Medicare coverage does not go into effect until 90 days after start of ESRD for those without existing Medicare including those under age 65 years without disabilities [3]. This lag period can contribute to an under-ascertainment of ESRD patients in the USA [2]. In addition, late-stage CKD cohort studies often pool information from different data sources and clinical care environments, potentially limiting the interpretation of findings [12, 13]. Thus, the ideal management strategies prior to and at the initiation of renal replacement continue to evolve.

We hypothesize that early dialysis mortality is affected by the circumstances at transition from CKD to ESRD. These factors may include the timing, clinical setting and events preceding dialysis, and ultimately the dialysis modality itself which may be manifestation of a pre-ESRD care. In this USRDS special center study [1, 14], we identified a large diverse CKD population that transitioned to ESRD from a single integrated health system with comprehensive electronic health records (EHR). Among patients who started peritoneal dialysis (PD), hemodialysis (HD) with a fistula/ graft (AVF/AVG), and HD with a catheter, we sought to compare 6-month mortality risk and determine risk factors for early ESRD mortality.

Methods

Kaiser Permanente Southern California (KPSC) is an integrated health system geographically located in southern California and comprised of 15 medical centers and over 200 satellite medical office buildings. Members have similar coverage benefits including co-pays for visits and medications. This study was approved by the KPSC institutional review board and exempted from informed consent (IRB #10591).

Study population

between January 2002 and March 2015 and had at least one estimated glomerular filtration (eGFR) measurement within 90 days before transition. CKD was defined as two separate eGFR < 45 ml/min/1.73 m² that were 90 days or more apart. eGFR < 45 ml/min/1.73 m² was chosen as the inclusion criteria for the CKD cohort because of the higher risk for ESRD in this population regardless of proteinuria [15]. The eGFR was calculated using the Chronic Kidney Disease Epidemiology Collaboration (CKD-EPI) equation [16]. ESRD was defined as treatment with HD (in center, home, or nocturnal) and PD. Patients who underwent preemptive renal transplantation were excluded.

All health information is collected as part of routine clinical care in a shared EHR. Study information on demographics, vitals, comorbidities, medications, and outcomes was extracted from the EHR and KPSC regional quality information data sources. The KPSC Renal Business Group captures and tracks all patients who initiated renal replacement therapy at an outpatient dialysis center or received a renal transplant. It also maintains a patient registry of all adult KPSC members who meet criteria for CKD [17]. Patient characteristics including vital signs, demographics, and laboratory results within 90 days prior to ESRD transition were extracted from the EHR. Comorbidity information was retrieved from the EHR using the International Classification of Diseases (ICD) codes. The cause of ESRD was extracted from the internal KPSC Renal Business Group records which was derived from clinician assessment and documentation by the physicians. Information on type of HD access used at the start of HD (AVF/AVG or vascular catheter], setting of first ESRD (inpatient vs outpatient start), and acute kidney injury (based on ICD codes) within 90 days before ESRD transition were also extracted. Patients were categorized based on ESRD treatment types (PD, HD with AVF/AVG, or HD with catheter), age in years (>80, 70–79, 60–69, < 60), and pre-ESRD transition eGFR. Patients were passively followed through the EHR for up to 6 months after ESRD transition.

Outcomes

The primary outcome evaluated was early all-cause mortality defined as within 6 months after transition to ESRD. Mortality information was obtained from the KPSC Mortality database which combines information from seven data sources, including California State Death Master Files, California State Multiple Cause of Death Master Files, Social Security Administration Death Master Files, KPSC Hospital and Emergency Room records, KPSC Membership System, Perinatal Data Mart, and Outside Claims Processing System. Mortality rates were calculated for each month following ESRD transition up to 6 months and reported as events per 1000 patient years. Patients were followed until a mortality outcome occurred, until 6 months of follow-up after ESRD, or until the end of the study observation period (March, 2015). Because mortality information was captured from KPSC and California state files, we did not censor any patients who lost membership or left the health plan. Despite the fact that ESRD patients may switch modalities or receive renal transplantation within the 6 months after ESRD transition, we analyzed our variables at the time of ESRD similar to an "intention to treat" approach.

Statistical analysis

Patient characteristics were compared across the three ESRD treatment types. Categorical variables were presented as counts (frequencies), and continuous variables were presented as means (standard deviation). χ^2 test was performed to compare independent categorical variables, and ANOVA was used to test the difference of continuous variables. Mortality rates were compared by treatment type (PD, HD with AVF/AVG, HD with catheter), prelude (pre-ESRD transition) eGFR, and age categories. The primary analysis was 6-month mortality risk after transition to ESRD using Cox proportional hazards modeling. Six-month HRs were estimated for ESRD treatment type. Multivariable hazard ratios (HR) were estimated adjusting for age, sex, race/ethnicity, comorbidities DM and heart failure, transition setting (inpatient or outpatient, acute kidney injury, and prelude laboratory values, including eGFR). We tested proportional assumption by introducing the interactions between exposure variables and time. No indication of proportionality violation was found. This was also confirmed by supremum test for proportional hazards. All hypothesis tests were twosided with a significance level of p < 0.05. Statistical analysis was conducted using SAS Enterprise Guide 5.1 (SAS Institute, Cary, NC).

Results

Among 71,127 individuals identified with CKD between 2002 and 2013, a total of 5819 (8.2%) transitioned to ESRD before March, 2015. The study cohort was comprised of 5373 CKD patients who transitioned to ESRD, had a documented eGFR within 90 days pre-transition, and had an ESRD modality identified (Fig. 1). The mean age was 62.7 years with 41.3% females, 37.5% Hispanics, 27.7% non-Hispanic whites, 22.7% blacks, and 9.8% Asians. The cause of ESRD was DM (69.1%), hypertension (12.9%), glomerulonephritis (5.8%), and other (12.2%). Overall, DM was identified in 79.1% and listed as the primary cause of ESRD for 69.1% of the study cohort. A total of 2266 (42.2%) patients had acute kidney injury within 90 days before ESRD transition (Table 1). The median hemoglobin level was

10.2 g/L, and 26.4% had a serum CO_2 measurement under 20 mEq/L (Table 1).

Within the study cohort, 714 (13.3%) patients transitioned to PD, 1874 (34.9%) patients transitioned to HD with AVF/AVG, and 2785 (51.8%) patients transitioned to HD with a catheter. There were 1915 (35.6%) patients who transitioned to ESRD in an inpatient setting (40.7% of HD patients and 2.5% of PD patients). Arteriovenous grafts were present in 4% of the total study cohort. The majority of the cohort transitioned to ESRD with an eGFR in the range of 5–14 mL/min/1.73 m².

Mortality

At 6 months following transition to ESRD, 551 patients had died (14 PD, 97 HD with AVF/AVG, and 440 HD with a catheter). Mortality rates were highest immediately (within month 1) after transition and then declined (Fig. 2a). Mortality rates were 299 (deaths per 1000 person-years) at month 1 and then declined to 224 by month 3 (Fig. 2a). This trend was mostly observed among the HD population but not among PD patients (Fig. 2b). Patients with an HD catheter had higher early mortality compared to HD patients who transitioned with an AVF/AVG. Higher early mortality was also observed in those who transitioned with eGFR \geq 15 mL/min/1.73 m² and in older patients (Fig. 2c, d).

Regression analyses for early ESRD mortality

Compared to patients who transitioned to PD, the fully adjusted 6-month mortality HRs (95% CI) were 1.87 (1.06-3.30) and 3.77 (2.17-6.57) for HD with AVF/AVG and HD with a catheter, respectively (Table 2). There was no difference in mortality risk between males and females. Older age demonstrated an incremental pattern of higher mortality risk with age ≥ 80 years having a 6-month HR of 4.19 (3.14-5.59) compared to age < 60 years. Non-Hispanic whites had worse survival compared to Hispanics, Asians, and blacks. Pre-ESRD transition eGFR < 5, 10-14, and \geq 15 ml/min/1.73 m² had 6-month mortality HRs of 1.04 (0.69–1.58), 1.38 (1.11–1.73), and 1.68 (1.34–2.11), respectively, compared to eGFR 5-9 ml/min/1.73 m². Heart failure or $CO_2 < 20$ mEq/L was not associated with higher early mortality risk. Overall, having DM had a lower 6-month mortality HR [0.67 (0.55–0.81)] within our study population (Supplemental Table 1).

The circumstances around the transition appeared to affect early mortality. Patients who transitioned to ESRD in the hospital (inpatient) had an HR of 1.32 (1.11–1.58) compared to those who started dialysis on an outpatient basis. Patients diagnosed with acute kidney injury within 90 days prior to ESRD had a 6-month HR of 2.06 (1.66–2.54) (Supplemental Table 1).



Fig. 1 Among a total of 71,127 individuals (between 2002 and 2013) who were identified with CKD, 5819 (8.2%) transitioned to ESRD. The study cohort was comprised of 5373 CKD patients who transi-

tioned to ESRD, had a documented eGFR within 90 days pre-transition, and had an ESRD modality identified

Sensitivity analyses using cause of ESRD in the multivariable modeling demonstrated 6-month mortality HRs of 1.96 (1.11–3.46) and 4.62 (2.67–8.00) for HD with AVF/ AVG and HD with a catheter, respectively, compared to PD (Supplemental Table 1).

Discussion

We evaluated early ESRD mortality in 5373 CKD patients who transitioned to dialysis among three different treatment types (PD, HD with an AVF/AVG, and HD with a catheter) and found that PD patients had the lowest risk for early mortality. Compared to PD patients, HD patients had a twofold to fourfold higher 6-month mortality risk depending on the type of access used at ESRD transition. Early mortality was also associated with patients who started dialysis at a higher eGFR, who were older, and who endured more acute circumstances prior to ESRD transition. The highest mortality was manifested within the first month. Our findings from a real-world clinical care environment demonstrate that the period immediately after ESRD represents the most vulnerable stage for CKD patients transitioning to ESRD.

These insights into the high mortality in early ESRD could have important implications for both prognostication and management purposes. Catheter use, volume overload, infrequent CKD care, and perhaps rapid eGFR decline emphasizes the importance of pre-ESRD transition care [18–21]. Early initiation of dialysis, however, does not seem to be associated with ESRD survival [22, 23]. By contrast, earlier ESRD transition may even be associated with worse outcomes [23, 24]. Although confounding by clinical indication affects the assessment, we similarly observed that patients who transitioned with eGFR 15 mL/min/m² or greater experienced the highest early mortality.

The high rates of mortality immediately after ESRD were mostly accounted for by the HD with catheter population. Our findings were similar to prior observations that demonstrated that HD patients with catheters had higher mortality compared to HD with AVF/AVG [20, 25, 26]. The HD with AVF/AVG population also demonstrated a higher mortality rate immediately after ESRD transition, though

Table 1	Characteristics and	comparisons b	by treatment grou	ip among CKE) patients who	transitioned to I	ESRD (N = 5373)
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Characteristics	Total	PD	HD with AVF/AVG	HD with catheter	P value
	N=5373	N=714	N=1874	N=2785	
Age group, N (%)					< 0.0001
< 50	2049 (38.3)	403 (56.5)	655 (34.9)	991 (35.6)	
60–69	1532 (28.5)	178 (24.9)	554 (29.6)	800 (28.7)	
70–79	1332 (24.8%)	104 (14.6)	506 (27.0)	722 (25.9)	
80+	460 (8.6%)	29 (4.1)	159 (8.5)	272 (9.8)	
Female, $N(\%)$	2221 (41.3)	301 (42.2)	746 (39.8)	1174 (42.2)	0.3
Race/ethnicity, $N(\%)$					0.0002
White	1488 (27.7)	180 (25.2)	512 (27.3)	796 (28.6)	
Asian	525 (9.8)	91 (12.7)	207 (11.0)	227 (8.2)	
Black	1221 (22.7)	133 (18.6)	413 (22.0)	675 (24.2)	
Hispanic	2014 (37.5)	291 (40.8)	704 (37.6)	1019 (36.6)	
Other	125 (2.3)	19 (2.7)	38 (2.1)	68 (2.5)	
Diabetes, N (%)	4252 (79.1)	522 (73.1)	1514 (80.8)	2216 (79.6)	0.0001
Heart Failure, N (%)	3657 (68.1)	343 (48)	1213 (64.7)	2101 (75.4)	< 0.0001
HTN, N (%)	5330 (99.2)	711 (99.6)	1868 (99.7)	2751 (98.8)	0.002
Inpatient transition, N (%)	1915 (35.6)	18 (2.5)	476 (25.4)	1421 (51)	< 0.0001
AKI in 90 days prior to ESRD, $N(\%)$	2266 (42.2)	114 (16)	419 (22.4)	1733 (62.2)	
eGFR category, N (%)					< 0.0001
<5	345 (6.4%)	27 (3.8)	106 (5.7)	212 (7.6)	
5–9	2396 (44.6%)	348 (48.7)	966 (51.6)	1082 (38.9)	
10–14	1743 (32.4%)	289 (40.5)	665 (35.5)	789 (28.3)	
≥15	889 (16.6%)	50 (7.0)	137 (7.3)	702 (25.2)	
CO ₂ in blood, median mEq/L (IQR)	22 (19.5-25.0)	22.0 (20.0-25.0)	22.0 (20.0-24.5)	22.0 (19.0-25.0)	< 0.0001
<20	1420 (26.4%)	143 (20%)	458 (24.4%)	819 (29.4%)	
≥20	3884 (72.3%)	571 (80%)	1402 (74.8%)	1911 (68.6%)	
Unmeasured	69 (1.3%)	0 (0%)	14 (0.7%)	55 (2%)	
HGB, median g/dL (IQR)	10.2 (9.3–11.2)	10.9 (9.9–11.8)	10.4 (9.4–11.2)	10.0 (9.1–10.9)	< 0.0001
PTH, median pg/ml (IQR)	111.0, 287.0	194 (121.0–288.0)	190 (117.0-302.0)	178 (105.0-277.0)	0.02
Ferritin, median ng/ml (IQR)	241.9 (123. 3-425.5)	211 (106.7-384.0)	246 (124.0-420.0)	248.4 (130.7-449.1)	0.002
Albumin, median g/L(IQR)	3.1 (2.6–3.5)	3.5 (3.0–3.9)	3.2 (2.7–3.6)	2.9 (2.4–3.3)	< 0.0001

All laboratory data were defined by the value in 90 days prior to baseline

not as pronounced as the catheter population. Conversely, the PD population appeared to have stable monthly mortality rates for the first 6 months after ESRD transition. The improved short-term outcomes in PD vs HD appear consistent with prior observations that compared dialysis modalities [27–29].

Our findings underscore the need to recognize the differences in patient characteristics and clinical circumstances among patients that transition to ESRD. These patient-specific, clinical environment-related, and treatment-related influences may help identify more optimal management strategies prior to and at the initiation of renal replacement. Unfortunately, there remain questions about the ideal management strategies in terms of blood pressure, DM, CKDmineral bone disorders (CKD-MBD), and anemia management strategies. The real-world clinical care and outcomes among a large diverse ESRD transition population could help provide additional and important insights.

In addition, the recognition of patients who are at highest risk for early mortality has potential benefit [9, 30]. The subgroup that experiences early mortality appears to represent a different population (or populations) altogether. More aggressive and timely management strategies such as earlier and more frequent nephrology care may be beneficial [21]. Conversely, the elderly and patients with high comorbidity burdens may have shortened life expectancies regardless of whether they receive dialysis [31, 32]. These represent populations/situations in which renal replacement may not improve or extend lifespan. The information may improve shared decision-making discussions which might more easily include conservative care of ESRD without dialysis as a viable option. With hundreds



Fig. 2 Monthly mortality rates up to 24 months of follow-up are reported in deaths per 100 person-years. **a** Mortality rates for the entire ESRD transition population (N=5373). **b** Mortality rates for hemodialysis with catheter (HD catheter), hemodialysis with AVF/AVG (HD-AVF/AVG) and peritoneal dialysis (PD) populations

separately not including transplant. **c** Mortality rates of the entire study cohort categorized by eGFR at transition (<5, 5–9, 10–14, and \geq 15 ml/min/1.73 m²). **d** Mortality rates of the entire study cohort categorized by age (<60, 60–69, 70–79, and \geq 80)

Table 2Hazard ratios for6-month mortality amongincident ESRD patients bytreatment group

	HR (95% CI) ^a	HR (95% CI) ^b	HR (95% CI) ^c
Modality			
PD	Reference		
HD with AVF/AVG	2.65 (1.51-4.64)	2.10 (1.20-3.68)	1.87 (1.06-3.30)
HD with Catheter	8.55 (5.02–14.55)	6.65 (3.90–11.34)	3.77 (2.17-6.57)

^aUnadjusted

^bAdjusted for age, sex, and race/ethnicity

^cAdjusted for age, sex, race/ethnicity, eGFR prior to transition, diabetes, heart failure, acute kidney injury, inpatient transition, bicarbonate level, and albumin

of thousands of people transitioning to ESRD throughout the world on an annual basis [33], the instances in which ESRD treatment and expenditures are futile and dialysis may not affect or prolong survival are of great concern. One potentially modifiable risk factor we observed was that patients with acidemia ($CO_2 < 20$) at ESRD transition had higher early mortality risk. Perhaps earlier and

improved management of acidemia may lead to better survival during the early period after starting dialysis.

The information from our integrated health system captures early mortality events more comprehensively and accurately than traditional USRDS data sources [2, 11]. Medicare coverage does not go into effect until 90 days after the start of ESRD for those without existing Medicare coverage including those under age 65 years without disabilities who experience early mortality. This accounts for up to 60% of patients who transition to ESRD [3]. By definition, our study from a single integrated health system with comprehensive clinical documentation and follow-up allowed a detailed assessment of pre-and post-ESRD transition outcomes that are likely to be captured without ascertainment bias or information gaps.

Potential limitations

Our study population may not reflect the experience of all individuals who start ESRD treatment given the fact that our cohort was selected based on CKD patients who were relatively stable, with two eGFR measurements 90 or more days apart. Thus, patients with normal baseline renal function who abruptly transitioned to ESRD were not included in our study. Catheter use and inpatient ESRD transitions were lower in our cohort than what has been described for the US ESRD population overall [1]. This may suggest that they may have been managed differently and perhaps were more "prepared" for ESRD transition. While we evaluated race/ethnicity, sex, and age on our outcome, we did not explore more granular patient information including socioeconomic and sociodemographic backgrounds. We also did not have reliable information on lifestyle including smoking and substance abuse. In addition, information on dialysis adherence or missed dialysis during the observation window was not available. Another consideration is that within the Kaiser Permanente integrated health system, there is more standardizations in care among the CKD population. CKD-specific interventions such as blood pressure control, anemia management, CKD-MBD management, and CKDrelated patient education are accessible to all CKD patients. Moreover, KPSC has an internal CKD registry that identifies patients and gives feedback to providers on management. Similar mechanisms are in place for other chronic conditions such as hypertension and patient safety nets [34, 35]. Thus, our findings may not reflect the heterogeneity in practice patterns and treatment environments found in pooled cohorts.

Another potential limitation of our study is the lack of information on cause of death. Cardiovascular events are high in the early ESRD period and are a major contributor to ESRD mortality [36]. Given the potential for discrepancies and reliability concerns, we were cautious about solely relying on the Center for Medicare and Medicaid Services (CMS) ESRD death notification form 2746 [37]. We also did not censor patients who switched dialysis modalities or received a renal transplant within the first 6 months. In addition, we did not censor patients that may have left the health plan because the state files would have captured any mortality although we may have missed deaths among patients who left the health plan and state and then died within 6 months.

Conclusion

Among a CKD population who transitioned to ESRD, we observed high mortality immediately after the start of dialysis especially among HD patients who transitioned with a catheter. Patients who transitioned in an inpatient setting, who had higher eGFR at transition, and who were recently diagnosed with AKI also had higher early mortality risk. This information may lead to more effective management strategies and inform shared decision making.

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Author contributions JJS and SJJ contributed to research area and study design; HZ and JS were involved in data acquisition; JJS, SJJ, HZ, JS, KKZ, and CPK participated in analysis or interpretation of data; JJS and SJJ were involved in study supervision; HZ was involved in statistical analysis; JJS drafted the manuscript; SJJ, HZ, SFS, KKZ, CPK, SH, and JS were involved in critical revision of the manuscript for important intellectual content; SFS and SJJ participated in administrative, technical, or material support. Each author contributed important intellectual content during the manuscript drafting or revision and accepts accountability of the overall work by ensuring that questions pertaining to the accuracy or integrity of any portion of the work are appropriately investigated and resolved. JJS takes responsibility that this study has been reported honestly, accurately, and transparently; that no important aspects of the study have been omitted, and that any discrepancies from the study as planned have been explained.

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Compliance with ethical standards

Conflict of interest All author declares that they have no conflict of interest

References

1. Saran R, Robinson B, Abbott KC, Agodoa LY, Ayanian J, Bragg-Gresham J, Balkrishnan R, Chen JL, Cope E, Eggers PW, Gillen D, Gipson D, Hailpern SM, Hall YN, Han Y, He K, Herman W, Heung M, Hutton D, Jacobsen SJ, Kalantar-Zadeh K, Kovesdy CP, Li Y, Lu Y, Molnar MZ, Morgenstern H, Nallamothu B, Nguyen DV, O'Hare AM, Obi Y, Plattner B, Pisoni R, Port FK, Rao P, Ravel V, Rhee CM, Sakhuja A, Schaubel DE, Selewski DT, Sim JJ, Song P, Streja E, Kurella Tamura M, Tentori F, White S, Woodside K, Hirth RA, Shahinian V (2017) US renal data system 2016 Annual data report: epidemiology of kidney disease in the United States. American Journal of Kidney Diseases: The Official Journal of the National Kidney Foundation 69 (3S1):A7–A8. https://doi.org/10.1053/j.ajkd.2016.12.004

- Foley RN, Chen SC, Solid CA, Gilbertson DT, Collins AJ (2014) Early mortality in patients starting dialysis appears to go unregistered. Kidney Int 86(2):392–398. https://doi.org/10.1038/ ki.2014.15
- Chan KE, Maddux FW, Tolkoff-Rubin N, Karumanchi SA, Thadhani R, Hakim RM (2011) Early outcomes among those initiating chronic dialysis in the United States. Clin J Am Soc Nephrol 6(11):2642–2649. https://doi.org/10.2215/CJN.03680411
- Robinson BM, Zhang J, Morgenstern H, Bradbury BD, Ng LJ, McCullough KP, Gillespie BW, Hakim R, Rayner H, Fort J, Akizawa T, Tentori F, Pisoni RL (2014) Worldwide, mortality risk is high soon after initiation of hemodialysis. Kidney Int 85(1):158– 165. https://doi.org/10.1038/ki.2013.252
- Sladoje-Martinovic B, Mikolasevic I, Bubic I, Racki S, Orlic L (2014) Survival of chronic hemodialysis patients over 80 years of age. Clin Interv Aging 9:689–696. https://doi.org/10.2147/CIA. S58976
- Bradbury BD, Fissell RB, Albert JM, Anthony MS, Critchlow CW, Pisoni RL, Port FK, Gillespie BW (2007) Predictors of early mortality among incident US hemodialysis patients in the dialysis outcomes and practice patterns study (DOPPS). Clin J Am Soc Nephrol 2(1):89–99. https://doi.org/10.2215/cjn.01170905
- Hwang SJ, Yang WC, Lin MY, Mau LW, Chen HC, Taiwan Society of N (2010) Impact of the clinical conditions at dialysis initiation on mortality in incident haemodialysis patients: a national cohort study in Taiwan. In: Nephrology, dialysis, transplantation : official publication of the European Dialysis and Transplant Association–European Renal Association 25 (8):2616–2624. https://doi.org/10.1093/ndt/gfq308
- Lukowsky LR, Kheifets L, Arah OA, Nissenson AR, Kalantar-Zadeh K (2012) Patterns and predictors of early mortality in incident hemodialysis patients: new insights. Am J Nephrol 35 (6):548–558. https://doi.org/10.1159/000338673
- Wick JP, Turin TC, Faris PD, MacRae JM, Weaver RG, Tonelli M, Manns BJ, Hemmelgarn BR (2016) A clinical risk prediction tool for 6-month mortality after dialysis initiation among older adults. Am J Kidney Dis. https://doi.org/10.1053/j.ajkd.2016.08.035
- 2020 HP https://www.healthypeople.gov/2020/topics-objectives/ topic/chronic-kidney-disease/objectives. Accessed 30 Jan 2017
- USRDS https://www.usrds.org/2016/view/v1_08.aspx. Accessed 30 Jan 2017
- Lash JP, Go AS, Appel LJ, He J, Ojo A, Rahman M, Townsend RR, Xie D, Cifelli D, Cohan J, Fink JC, Fischer MJ, Gadegbeku C, Hamm LL, Kusek JW, Landis JR, Narva A, Robinson N, Teal V, Feldman HI (2009) Chronic Renal Insufficiency Cohort (CRIC) study: baseline characteristics and associations with kidney function. Clin J Am Soc Nephrol 4(8):1302–1311. https://doi. org/10.2215/CJN.00070109
- Mariani L, Stengel B, Combe C, Massy ZA, Reichel H, Fliser D, Pecoits-Filho R, Lopes AA, Yamagata K, Wada T, Wong MM, Speyer E, Port FK, Pisoni RL, Robinson BM (2016) The CKD outcomes and practice patterns study (CKDopps): rationale and methods. Am J Kidney Dis 68(3):402–413. https://doi. org/10.1053/j.ajkd.2016.03.414

- 14. Kalantar-Zadeh K, Kovesdy CP, Streja E, Rhee CM, Soohoo M, Chen JL, Molnar MZ, Obi Y, Gillen D, Nguyen DV, Norris KC, Sim JJ, Jacobsen SS (2017) Transition of care from pre-dialysis prelude to renal replacement therapy: the blueprints of emerging research in advanced chronic kidney disease. Nephrol Dial Trans Off Pub Eur Dial Trans Assoc Eur Renal Assoc. https://doi. org/10.1093/ndt/gfw357
- Levey AS, de Jong PE, Coresh J, El Nahas M, Astor BC, Matsushita K, Gansevoort RT, Kasiske BL, Eckardt KU (2011) The definition, classification, and prognosis of chronic kidney disease: a KDIGO controversies conference report. Kidney Int 80(1):17–28. https://doi.org/10.1038/ki.2010.483
- Levey AS, Stevens LA, Schmid CH, Zhang YL, Castro AF 3rd, Feldman HI, Kusek JW, Eggers P, Van Lente F, Greene T, Coresh J, Ckd EPI (2009) A new equation to estimate glomerular filtration rate. Ann Int Med 150(9):604–612
- Rutkowski M, Mann W, Derose S, Selevan D, Pascual N, Diesto J, Crooks P (2009) Implementing KDOQI CKD definition and staging guidelines in Southern California Kaiser Permanente. Am J Kidney Dis Off J Natl Kidney Found 53(3 Suppl 3):S86–99. https ://doi.org/10.1053/j.ajkd.2008.07.052
- Rivara MB, Chen CH, Nair A, Cobb D, Himmelfarb J, Mehrotra R (2016) Indication for dialysis initiation and mortality in patients with chronic kidney failure: a retrospective cohort study. Am J Kidney Dis Off J Natl Kidney Found. https://doi.org/10.1053/j. ajkd.2016.06.024
- Naimark DM, Grams ME, Matsushita K, Black C, Drion I, Fox CS, Inker LA, Ishani A, Jee SH, Kitamura A, Lee JP, Nally J, Peralta CA, Rothenbacher D, Ryu S, Tonelli M, Coresh J, Gansevoort RT, Yatsuya H, Warnock DG, Woodward M, de Jong PE, Consortium CKDP (2016) Past decline versus current eGFR and subsequent mortality risk. J Am Soc Nephrol JASN 27(8):2456– 2466. https://doi.org/10.1681/ASN.2015060688
- Xue JL, Dahl D, Ebben JP, Collins AJ (2003) The association of initial hemodialysis access type with mortality outcomes in elderly Medicare ESRD patients. Am J Kidney Dis Off J Natl Kidney Found 42(5):1013–1019
- Fischer MJ, Stroupe KT, Kaufman JS, O'Hare AM, Browning MM, Sohn MW, Huo Z, Hynes DM (2016) Predialysis nephrology care and dialysis-related health outcomes among older adults initiating dialysis. BMC Nephrol 17(1):103. https://doi.org/10.1186/ s12882-016-0324-5
- Cooper BA, Branley P, Bulfone L, Collins JF, Craig JC, Fraenkel MB, Harris A, Johnson DW, Kesselhut J, Li JJ, Luxton G, Pilmore A, Tiller DJ, Harris DC, Pollock CA (2010) A randomized, controlled trial of early versus late initiation of dialysis. N England J Med 363(7):609–619. https://doi.org/10.1056/NEJMoa1000552
- 23. Yamagata K, Nakai S, Iseki K, Tsubakihara Y (2012) Late dialysis start did not affect long-term outcome in Japanese dialysis patients: long-term prognosis from Japanese Society for [corrected] dialysis therapy registry. Therapeutic Apheresis Dial 16(2):111–120. https://doi.org/10.1111/j.1744-9987.2011.01052.x
- Wright S, Klausner D, Baird B, Williams ME, Steinman T, Tang H, Ragasa R, Goldfarb-Rumyantzev AS (2010) Timing of dialysis initiation and survival in ESRD. Clin J Am Soc Nephrol CJASN 5(10):1828–1835. https://doi.org/10.2215/CJN.06230909
- Dhingra RK, Young EW, Hulbert-Shearon TE, Leavey SF, Port FK (2001) Type of vascular access and mortality in U.S. hemodialysis patients. Kidney Int 60(4):1443–1451. https://doi.org/10.1 046/j.1523-1755.2001.00947.x
- Moist LM, Trpeski L, Na Y, Lok CE (2008) Increased hemodialysis catheter use in Canada and associated mortality risk: data from the Canadian organ replacement registry 2001–2004. Clin J Am Soc Nephrol 3(6):1726–1732. https://doi.org/10.2215/cjn.01240 308

- 27. Jaar BG, Coresh J, Plantinga LC, Fink NE, Klag MJ, Levey AS, Levin NW, Sadler JH, Kliger A, Powe NR (2005) Comparing the risk for death with peritoneal dialysis and hemodialysis in a national cohort of patients with chronic kidney disease. Ann Int Med 143(3):174–183
- Ganesh SK, Hulbert-Shearon T, Port FK, Eagle K, Stack AG (2003) Mortality differences by dialysis modality among incident ESRD patients with and without coronary artery disease. J Am Soc Nephrol JASN 14(2):415–424
- Vonesh EF, Snyder JJ, Foley RN, Collins AJ (2004) The differential impact of risk factors on mortality in hemodialysis and peritoneal dialysis. Kidney Int 66(6):2389–2401. https://doi.org/10.1 111/j.1523-1755.2004.66028.x
- Thamer M, Kaufman JS, Zhang Y, Zhang Q, Cotter DJ, Bang H (2015) Predicting early death among elderly dialysis patients: development and validation of a risk score to assist shared decision making for dialysis initiation. Am J Kidney Dis Off J Natl Kidney Found 66 (6):1024–1032. https://doi.org/10.1053/j. ajkd.2015.05.014
- Verberne WR, Geers AB, Jellema WT, Vincent HH, van Delden JJ, Bos WJ (2016) Comparative survival among older adults with advanced kidney disease managed conservatively versus with dialysis. Clin J Am Soc Nephrol 11(4):633–640. https://doi. org/10.2215/CJN.07510715
- Foote C, Ninomiya T, Gallagher M, Perkovic V, Cass A, McDonald SP, Jardine M (2012) Survival of elderly dialysis patients is predicted by both patient and practice characteristics. Nephrol Dial Transp 27(9):3581–3587. https://doi.org/10.1093/ndt/gfs096

- 33. Jha V, Garcia-Garcia G, Iseki K, Li Z, Naicker S, Plattner B, Saran R, Wang AY, Yang CW (2013) Chronic kidney disease: global dimension and perspectives. Lancet 382(9888):260–272. https://doi.org/10.1016/S0140-6736(13)60687-X
- 34. Sim JJ, Rutkowski MP, Selevan DC, Batech M, Timmins R, Slezak JM, Jacobsen SJ, Kanter MH (2015) Kaiser permanente creatinine safety program: a mechanism to ensure widespread detection and care for chronic kidney disease. Am J Med 128 (11):1204–1211. https://doi.org/10.1016/j.amjmed.2015.05.037
- Sim JJ, Handler J, Jacobsen SJ, Kanter MH (2014) Systemic implementation strategies to improve hypertension: the Kaiser permanente southern California experience. Can J Cardiol 30 (5):544–552. https://doi.org/10.1016/j.cjca.2014.01.003
- 36. Eckardt KU, Gillespie IA, Kronenberg F, Richards S, Stenvinkel P, Anker SD, Wheeler DC, de Francisco AL, Marcelli D, Froissart M, Floege J, Committee AROS (2015) High cardiovascular event rates occur within the first weeks of starting hemodialysis. Kidney Int 88(5):1117–1125. https://doi.org/10.1038/ki.2015.117
- 37. Rocco MV, Yan G, Gassman J, Lewis JB, Ornt D, Weiss B, Levey AS, Hemodialysis Study G (2002) Comparison of causes of death using HEMO Study and HCFA end-stage renal disease death notification classification systems. In: The National Institutes of Health-funded Hemodialysis. Health Care Financing Administration. American journal of kidney diseases: the official journal of the National Kidney Foundation 39 (1):146–153