



## Original article

# Vascular access type and mortality in elderly incident hemodialysis patients

Marisa Roldão\*, Cátia Figueiredo, Rachele Escoli, Hernâni Gonçalves, Flora Sofia, Karina Lopes

Department of Nephrology, Centro Hospitalar do Médio Tejo, Torres Novas, Portugal

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### ABSTRACT

**Introduction:** The ideal vascular access type for elderly hemodialysis (HD) patients remains debatable. The aim of this study was to analyze the association between patterns of vascular access use within the first year of HD and mortality in elderly patients.

**Methods:** Single-center retrospective study of 99 incident HD patients aged  $\geq 80$  years from January 2010 to May 2021. Patients were categorized according to their patterns of vascular access use within the first year of HD: central venous catheter (CVC) only, CVC to arteriovenous fistula (AVF), AVF to CVC, and AVF only. Baseline clinical data were compared among groups. Survival outcomes were analyzed using Kaplan–Meier survival curves and Cox's proportional hazards model.

**Results:** When compared with CVC to AVF, mortality risk was significantly higher among CVC only patients and similar to AVF only group [HR 0.93 (95% CI 0.32–2.51)]. Ischemic heart disease [HR 1.74 (95% CI 1.02–2.96)], lower levels of albumin [HR 2.16 (95% CI 1.28–3.64)] and hemoglobin [HR 4.10 (95% CI 1.69–9.92)], and higher levels of c-reactive protein [HR 1.87 (95% CI 1.11–3.14)] were also associated with increased mortality risk in our cohort,  $p < 0.05$ .

**Conclusion:** Our findings suggested that placement of an AVF during the early stages of dialysis was associated with lower mortality compared to persistent CVC use among elderly patients. AVF placement appears to have a positive impact on survival outcomes, even in those who started dialysis with a CVC.

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\* Corresponding author.

E-mail address: [marisa.roldao@chmt.min-saude.pt](mailto:marisa.roldao@chmt.min-saude.pt) (M. Roldão).

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## Tipo de acceso vascular y mortalidad en pacientes ancianos en programa de hemodiálisis

### RESUMEN

#### Palabras clave:

Anciano  
Hemodiálisis  
Acceso vascular  
Mortalidad

**Introducción:** El tipo de acceso vascular ideal para pacientes ancianos en hemodiálisis (HD) sigue siendo discutible. El objetivo de este estudio fue analizar la asociación entre los patrones de uso del acceso vascular en el primer año de HD y la mortalidad en pacientes ancianos.

**Métodos:** Estudio retrospectivo unicéntrico de 99 pacientes incidentes en HD con edades  $\geq 80$  años desde enero de 2010 hasta mayo de 2021. Los pacientes fueron categorizados según sus patrones de uso del acceso vascular en el primer año de HD: catéter venoso central (CVC) solo, CVC a fístula arteriovenosa (FAV), FAV a CVC y FAV solamente. Los datos clínicos iniciales se compararon entre los grupos. Los resultados de supervivencia se analizaron mediante las curvas de supervivencia de Kaplan-Meier y el modelo de riesgo proporcional de Cox.

**Resultados:** En comparación con el CVC para la FAV, el riesgo de mortalidad fue significativamente mayor entre los pacientes que solo recibieron CVC y similar al grupo que solo utilizó FAV (HR: 0,93; IC 95%: 0,32-2,51). Cardiopatía isquémica (HR: 1,74; IC 95%: 1,02-2,96), niveles más bajos de albúmina (HR: 2,16; IC 95%: 1,28-3,64) y de hemoglobina (HR: 4,10; IC 95%: 1,69-9,92), y niveles más altos de proteína C reactiva (HR: 1,87; IC 95%: 1,11-3,14) también se asociaron con un mayor riesgo de mortalidad en nuestra cohorte ( $p < 0,05$ ).

**Conclusión:** Nuestros hallazgos sugirieron que la colocación de una FAV durante las primeras etapas de la diálisis se asoció con una menor mortalidad en comparación con el uso persistente de CVC en pacientes ancianos. La colocación de una FAV parece tener un impacto positivo en los resultados de supervivencia, incluso en aquellos que comenzaron la diálisis con un CVC.

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## Introduction

The population of elderly patients with end-stage renal disease is growing worldwide. The 2019 Annual Report of the ERA-EDTA registry reported that 45% of the total prevalent European renal replacement therapy patients were aged  $\geq 65$  years.<sup>1</sup> The high prevalence of comorbidities, limited life expectancy and complex quality of life issues associated with this population pose unique challenges for clinicians, including choosing the ideal vascular access type for hemodialysis (HD).<sup>2-4</sup> There is a wide debate about whether the “Fistula First” Initiative should be applied to elderly patients in HD, given lower maturation rates, longer maturation times and emerging data indicating the lack of a survival benefit of arteriovenous fistula (AVF) compared with central venous catheter (CVC) use in this group of patients.<sup>5,6</sup> A new approach introduced by the KDOQI Clinical Practice Guideline for Vascular Access (2019) proposed an individualized and comprehensive map for dialysis modalities for the lifetime of the patient called the “ESKD Life-Plan,” achieved by creating a plan for each patient that considers the Patient’s Life-Plan and corresponding Access Needs. This comprehensive patient-adjusted vascular access management plan throughout the period of end-stage renal disease, appears especially important in elderly patients.<sup>7</sup> In fact, most elderly patients start dialysis with a CVC as their primary vascular access. A recent study

suggested a lower mortality risk among elderly incident HD patients who had an AVF placed from a CVC in the early phases of dialysis, compared with those with persistent CVC use.<sup>8</sup> The aim of our study was to analyze whether specific patterns of vascular access use within the first year of dialysis were associated with differential survival outcomes among a cohort of incident HD patients aged  $\geq 80$  years.

## Material and methods

We conducted a single-center retrospective cohort study of incident HD patients at Centro Hospitalar do Médio Tejo, EPE HD unit between January 2010 and May 2021. All patients with  $\geq 80$  years were included. Patients who recovered renal function, lost follow-up, or switched to another renal replacement therapy were excluded. Patients were categorized according to their patterns of vascular access use within the first year of HD: CVC only, CVC to AVF, AVF only and AVF to CVC use.

The primary outcome was all-cause mortality, and secondary outcomes included laboratory data related to control of anemia, mineral bone disease and inflammatory markers, during the first year of treatment.

Baseline clinical and demographic characteristics, including age, gender, cause of end-stage renal disease, and previous nephrology follow-up were collected, as well as patient

**Table 1 – Baseline patient characteristics stratified by patterns of vascular access type during the first year of dialysis.**

	Overall cohort	CVC only	CVC to AVF	AVF only	AVF to CVC	p-Value
Patients, n (%)	99 (100%)	55 (56.6%)	19 (19.2%)	13 (13.1%)	11 (11.1%)	
Age (years) mean $\pm$ SD	85.1 $\pm$ 3.9	85.1 $\pm$ 4.1	84.9 $\pm$ 3.2	85.7 $\pm$ 5.1	84.8 $\pm$ 3.5	0.949
Male, n (%)	48 (48.5%)	27 (51.8%)	10 (47.4%)	4 (69.2%)	7 (63.6%)	0.427
ESRD etiology, n (%)						0.451
Diabetes mellitus	23 (23.2%)	14 (25%)	4 (21.1%)	4 (30.8%)	1 (9.1%)	
Hypertension	15 (15.2%)	4 (7.1%)	2 (10.5%)	1 (7.7%)	1 (12.5%)	
Chronic glomerulonephritis	9 (9.1%)	5 (8.9%)	2 (10.5%)	2 (15.4%)	1 (12.5%)	
Unknown	21 (21.2%)	9 (16.1%)	3 (15.8%)	4 (30.8%)	5 (45.5%)	
Other	31 (31.3%)	23 (41.8%)	8 (42.1%)	2 (15.4%)	3 (20.4%)	
Comorbidities, n (%)						
Diabetes mellitus	44 (44.5%)	25 (44.6%)	10 (52.6%)	5 (38.5%)	4 (36.4%)	0.802
Ischemic heart disease	60 (60.6%)	37 (66.1%)	10 (52.6%)	7 (53.8%)	6 (54.5%)	0.654
Cerebrovascular disease	16 (16.2%)	8 (14.3%)	5 (26.3%)	1 (7.7%)	2 (18.2%)	0.510
Peripheral artery disease	15 (15.2%)	11 (19.6%)	2 (10.5%)	1 (9.1%)	1 (15.2%)	0.558
CCI ( $\geq$ 8)	78 (78.8%)	49 (87.5%)	15 (78.9%)	9 (69.2%)	5 (45.5%)	<b>0.014</b>
CCI mean $\pm$ SD	8.41 $\pm$ 1.65	8.73 $\pm$ 1.51	8.63 $\pm$ 1.57	7.69 $\pm$ 1.18	7.27 $\pm$ 2.33	<b>0.015</b>
Previous Nephrology follow-up, n (%)	69 (69.7%)	35 (62.5%)	15 (78.9%)	11 (84.6%)	8 (72.7%)	0.313
Urgent HD start n (%)	79 (79.8%)	50 (89.3%)	18 (94.7%)	5 (38.5%)	6 (54.6%)	<b>0.001</b>
Inpatient n (%)	86 (86.9%)	52 (92.9%)	18 (94.7%)	8 (61.5%)	8 (72.7%)	<b>0.007</b>

p-Values represent an ANOVA test for continuous variables and Chi-square test for categorical variables.

ESRD, end-stage renal disease; Hgb, hemoglobin; SD, standard deviation; IQR, interquartile range. In bold values with statistical significance.

comorbidity score according to the Charlson Comorbidity Index (CCI, low  $<$  8 vs high  $\geq$  8).

Laboratory parameters were measured at dialysis initiation and monthly during the first year of HD, including hemoglobin (Hgb), phosphorus ( $P^+$ ), calcium ( $Ca^{2+}$ ), albumin, c-reactive protein (CRP) and potassium ( $K^+$ ). Serum intact parathyroid hormone (iPTH) and ferritin were measured every 3 months.

Statistical analysis was performed using SPSS version 23.0 for Mac OS X. Continuous variables were presented as mean and standard deviation or median and interquartile range (IQR) for variables with skewed distributions. Nominal variables were presented as number (frequency) and percentage. A  $p$  value  $<$  0.05 was considered statistically significant.

Baseline characteristics and laboratory data were compared among groups using the ANOVA test for normally distributed continuous variables ( $P^+$ ), Kruskal-Wallis test for skewed distributed continuous variables (Hgb,  $Ca^{2+}$ , albumin, CRP,  $K^+$ , iPTH and ferritin) and Chi-square test for categorical variables.

The association between patterns of vascular access use within the first year of treatment and all-cause mortality risk were examined using standard survival analysis methods. Survival time was calculated from the date of dialysis initiation until the date of death or end of follow-up (May 31, 2021). Kaplan-Meier curves were developed for each group and compared by log-rank tests. Univariate hazard ratios (HRs) and 95% confidence intervals (CIs) for all-cause mortality were estimated using the Cox proportional hazards risk model.

## Results

Ninety-nine patients aged  $\geq$  80 years were recruited into the study: 48 (48.5%) were male, 44 (44.4%) diabetic, 60 (60.6%) had ischemic heart disease and 15 (15.2%) peripheral artery

disease. Mean CCI was  $8.41 \pm 1.65$ . Mean age was  $85.1 \pm 3.9$  years and eleven patients (11.1%) were over 90 years old. Seventy-four patients (75.8%) started dialysis with a CVC. Over the first year of dialysis, we observed that 56.6% ( $n = 56$ ) of patients persisted in CVC use only, 19.2% ( $n = 19$ ) underwent placement of an AVF from a CVC, 13.1% ( $n = 13$ ) persisted in AVF use only and 11.1% ( $n = 11$ ) underwent placement of a CVC from an AVF. The duration of CVC use was  $219.5 \pm 100.6$  days in the CVC to AVF group and  $295.4 \pm 107.4$  days in the AVF to CVC group.

No statistically significant differences were found in age, gender, or cause of end-stage renal disease (ESRD) among groups. Compared with other patients, those who persisted on having a CVC only were more likely to initiate dialysis with urgent criteria, as inpatients and to have  $CCI \geq 8$  (Table 1).

Laboratory data related to ESRD complications (anemia and mineral bone disease), nutrition and inflammation parameters in the first year of HD were similar among groups, except for iPTH which was significantly lower in the CVC only group ( $p <$  0.05). Hemodialysis efficacy (spKT/V) was also similar (Table 2).

Among the patients enrolled in the study, there were 64 deaths over a follow-up period of 228 person-years. The mean follow-up was 2.3 years. The overall mortality rates by pattern of vascular access use within the first year of dialysis were 42.3, 17.4, 15.1 and 19.8 per 100 person-years for CVC only, CVC to AVF, AVF only and AVF to CVC, respectively. Fig. 1 shows non-adjusted Kaplan-Meier curves for all-cause death according to groups stratified by vascular access type. Patients in CVC only group showed a higher incidence rate of all-cause death compared with the other groups (log-rank test,  $p <$  0.05).

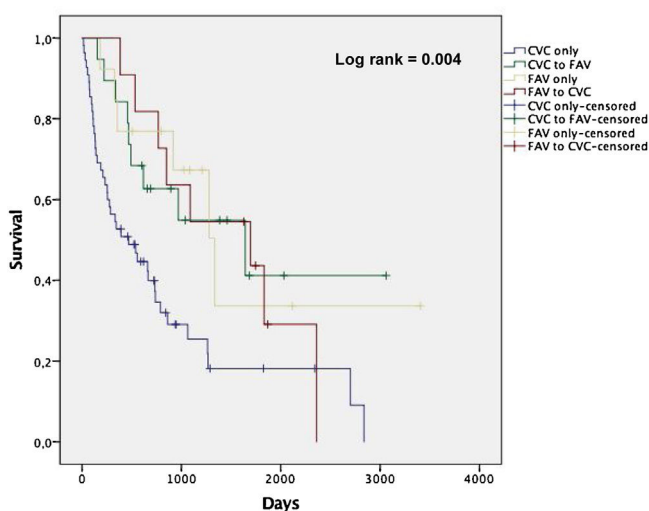
The mean and median survival times (in days) revealed a statistically significant disadvantage of CVC only group ( $p <$  0.05), demonstrating that the observed difference in survival curves was not due to chance. CVC to AVF patients

**Table 2 – Clinical and analytical outcomes stratified by patterns of vascular access type during the first year of dialysis.**

	CVC only	CVC to AVF	AVF only	AVF to CVC	p-Value
Hgb (g/dL) median (IQR)	10.68 (2)	11.05 (1.3)	11.18 (1)	10.18 (0.7)	0.260
Albumin (g/dL) median (IQR)	2.8 (1.2)	3.13 (0.4)	3.11 (0.8)	3.03 (0.7)	0.271
iPTH (ng/mL) median (IQR)	123.7 (198.3)	255.1 (201)	229.25 (391)	247 (352.2)	<b>0.009</b>
Calcium (mg/dL) median (IQR)	8.63 (0.3)	8.48 (0.8)	8.96 (0.8)	8.73 (0.6)	0.062
Phosphorus (mg/dL) mean ± SD	3.49 ± 1.49	3.9 ± 1.26	3.87 ± 0.64	4.18 (1.01)	0.325
Potassium (mmol/L) median (IQR)	4.65 (1.1)	4.58 (0.8)	4.97 (0.8)	5.03 (0.7)	0.166
CRP (mg/dl) median (IQR)	1.31 (3.7)	0.98 (1.3)	0.58 (0.8)	1.35 (3.5)	0.095
Ferritin (mg/dL) median (IRQ)	348.98 (368)	261.05 (168)	410.25 (351)	191.85 (359)	0.064
BMI (kg/m <sup>2</sup> ) mean ± SD	23.92 ± 5.36	25.10 ± 5.23	24.67 ± 4.95	24.39 ± 5.11	0.386
Kt/V median (IRQ)	1.73 (0.54)	1.85 (0.43)	1.89 (0.32)	1.79 (0.54)	0.255

p-Values represent an ANOVA test for normally distributed continuous variables (phosphorus) and a Kruskal–Wallis test for skewed distributed continuous variables.

Hgb, hemoglobin; SD, standard deviation; IQR, interquartile range; iPTH, intact parathormone; CRP, c-reactive protein. In bold values with statistical significance.

**Fig. 1 – Kaplan–Meier curves for all-cause mortality in the 4 groups.**

showed a similar mean and median survival times as AVF only (Table 3). Among the CVC only patients, the rate of CVC-related bloodstream infection was 0.6 per 1000 days of CVC and infection-related mortality was 10.3 per 100 person-years (24.4% of all deaths in this group).

Using the univariate Cox model, we compared patients with persistent CVC use to those with placement of an AVF

from a CVC [HR 0.42 (95% CI 0.21–0.87)] and AVF only [HR 0.38 (95% CI 0.16–0.89)] with both groups showing a lower mortality risk ( $p < 0.05$ ). Those with placement of a CVC from an AVF also showed a trend towards lower mortality risk [HR 0.49 (95% CI 0.23–1.07)], although it was not statistically significant ( $p > 0.05$ ) (Table 4). When the reference group was changed to patients with placement of an AVF from a CVC, those with AVF use only showed a similar mortality risk [HR 0.93 (95% CI 0.32–2.51)]. By using univariate Cox proportional hazards regression, we found that other significant risk factors for all-cause mortality among these patients were ischemic heart disease [HR 1.74 (95% CI 1.02–2.96)], lower albumin ( $< 3$  g/dL) [HR 2.16 (95% CI 1.28–3.64)], lower Hgb ( $< 8.5$  g/dL) [HR 4.10 (95% CI 1.69–9.92)] and higher CRP ( $> 2.5$  mg/dL) [HR 1.87 (95% CI 1.11–3.14)]. In the multivariable Cox regression model, after adjustment for these risk factors (albumin  $< 3$  g/dL, Hgb  $< 8.5$  g/dL and CRP  $< 2.5$  mg/dL), mortality risk remained higher in the CVC group when compared with those with AVF only [HR 0.40 (95% CI 0.19–0.83)] ( $p < 0.05$ ). The same trend was observed among those with placement of an AVF from a CVC [HR 0.51 (95% CI 0.26–1.03)] and CVC from an AVF [HR 0.71 (95% CI 0.41–1.24)] but did not reach statistical significance ( $p > 0.05$ ) (Table 4).

## Discussion

The use of CVCs among incident HD patients remains high, particularly in older patients ( $> 80$  years of age). In addition,

**Table 3 – Confidence intervals mean and median for survival time (in days) according to patterns of vascular access type during the first year of dialysis.**

Groups	Mean				Median			
	Estimate	Std. error	95% confidence interval		Estimate	Std. error	95% confidence interval	
			Lower bound	Upper bound			Lower bound	Upper bound
CVC only	939.944	162.708	621.037	1258.851	469.000	182.476	111.347	826.653
CVC to AVF	1708.427	316.036	1088.996	2327.859	1642.000	713.790	242.971	3042.029
AVF only	1740.663	436.323	885.471	2595.856	1692.000	654.570	413.043	2978.957
AVF to CVC	1467.109	240.123	996.467	1937.751	1335.000	225.462	893.094	1776.906
Overall	1268.743	137.320	999.596	1537.890	789.000	136.923	520.630	1057.370

CVC, central venous catheter; AVF, arteriovenous fistula.

**Table 4 – Association between patterns of vascular access type during the first year of dialysis and the risk of all-cause mortality examined by Cox proportional hazard models.**

Groups	Univariate analysis			Multivariate analysis <sup>a</sup>		
	HR	95% confidence interval	p-Value	HR	95% confidence interval	p-Value
CVC only		Reference			Reference	
CVC to AVF	0.42	0.21–0.87	<b>0.020</b>	0.51	0.26–1.03	0.063
AVF only	0.38	0.16–0.89	<b>0.030</b>	0.40	0.19–0.83	<b>0.013</b>
AVF to CVC	0.49	0.23–1.07	0.080	0.71	0.41–1.24	0.230
Ischemic heart disease	1.74	1.02–2.96	<b>0.036</b>			
Albumin (<3 g/dL)	2.16	1.28–3.64	<b>0.005</b>			
CRP (>2.5 mg/dL)	1.87	1.11–3.14	<b>0.023</b>			
Hgb (<8.5 g/dL)	4.10	1.69–9.92	<b>0.007</b>			

CVC, central venous catheter; AVF, arteriovenous fistula; CRP, c-reactive protein; Hgb, hemoglobin.

<sup>a</sup> Adjusted for albumin (<3 g/dL), CRP (>2.5 mg/dL) and Hgb (<8.5 g/dL). In bold values with statistical significance.

a large proportion of patients who initiate treatment with a CVC do not switch to a permanent vascular access.<sup>8</sup> Among those who make the transition, only a minority develop a functional AVF.<sup>8</sup> Successful creation of an AVF requires suitable vasculature. Vein distensibility may be affected by a greater prevalence of peripheral vascular disease among elderly, which is supported by the greater risk of AVF failure found in this population.<sup>9,10</sup> Moreover, several aspects of CVCs, including immediate readiness for use and the absence of pain with cannulation may make this an appealing vascular access option among elderly HD patients with limited life expectancy.<sup>11</sup> In our study, most patients (75.8%) started dialysis with a CVC. Compared with other patients, those who started with a CVC were more likely to initiate dialysis as inpatients by urgent criteria, despite similar predialysis follow-up. Elderly patients are particularly susceptible to the development of acute kidney injury due to structural and functional deterioration of the kidneys, decreased renal reserve and the presence of comorbidities, reducing the ability to recover.<sup>12</sup> Thus, elderly patients may present with urgent indications for dialysis prior to placement and maturation of an AVF, contributing to the high incidence of CVC use at dialysis initiation in this population.<sup>13–15</sup> On the other hand, older patients lose kidney function at lower rates than younger ones, have a shorter survival due to competing risks of mortality and may be more likely to die before benefiting from an AVF.<sup>16–18</sup> Furthermore, they may be more likely to experience primary AVF failure which also increases the incidence of CVCs, morbidity, and mortality in this group.<sup>19</sup> Therefore, we aimed to determine whether placement of an AVF from a CVC during the first year of dialysis was associated with better survival outcomes when compared to persistent CVC use in patients aged  $\geq 80$  years. Among our cohort, compared with patients with persistent CVC use, those with AVF use only and placement of an AVF from a CVC had lower all-cause mortality risk. When the reference group was changed to patients with placement of an AVF from a CVC, those with AVF use only showed similar mortality risk. Our findings corroborate the results of Ko et al., suggesting a positive impact on survival outcomes associated with AVF placement in elderly, even in those who started dialysis with a CVC.<sup>8</sup> In this population the presence of ischemic heart disease, lower levels of albumin (<3 g/dL) and Hgb (<8.5 g/dL) and higher levels of CRP (>2.5 mg/dL) were also

associated with increased mortality risk. Yeh et al. described similar results, identifying hypoalbuminemia and high CRP as mortality risk factors in elderly HD patients,<sup>20</sup> suggesting an important role of inflammatory status in survival outcomes in this population.<sup>21,22</sup> Indeed, recent observational studies have suggested that elderly patient characteristics account for a large fraction of the excess mortality associated with CVC use in this population.<sup>23,24</sup> Although we cannot totally exclude this assumption, in our study, persistent CVC use was an independent mortality risk factor, even in models adjusted to patients' characteristics, such as hypoalbuminemia, anemia and high inflammatory markers. Our findings suggest that placement of an AVF from a CVC during the early phases of dialysis is associated with lower mortality compared to persistent CVC use and similar mortality compared to AVF use only, among incident HD patients aged  $\geq 80$  years. Despite our results, a pragmatic patient-centered approach is mandatory, considering the possibility that the 'AVF first' approach should not be an absolute for all patients. There were some limitations in this study. First, this was a retrospective study with a small sample size. Second, this is not a randomized study and biases may exist as patients with worst general condition tend to receive HD via CVC. Third, the patient number were different among groups. Fourth, the study reflects the elderly population of a hospital HD unit.

### Key concepts

- The high prevalence of comorbidities, limited life expectancy and complex quality of life issues pose unique challenges, including choosing the ideal vascular access type for HD in elderly patients.
- Our findings suggest that placement of an AVF from a CVC during the early phases of dialysis is associated with lower mortality compared to persistent CVC use and similar mortality compared to AVF use only, among incident HD patients aged  $\geq 80$  years.
- A pragmatic patient-centered approach is mandatory, considering the possibility that the 'AVF first' approach should not be an absolute for all patients.

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## Conflict of interest

No conflict of interest.

## REFERENCES

- Kramer A, Pippias M, Noordzij M, Stel VS, Afentakis N, Ambuhl PM, et al. The European Renal Association – European Dialysis and Transplant Association (ERA-EDTA) Registry Annual Report 2015: a summary. *Clin Kidney J.* 2018;11:108–22.
- Roca-Tey R. El acceso vascular para hemodiálisis: la asignatura pendiente. *Nefrologia.* 2010;30:280–7.
- Roca-Tey R. El acceso vascular del paciente anciano en programa de hemodiálisis. *Nefrol Sup Ext.* 2012;3:13–20.
- Vachharajani TJ, Moist LM, Glickman MH, Vazquez MA, Polkinghorne KR, Lok CE, et al. Elderly patients with CKD-dilemmas in dialysis therapy and vascular access. *Nat Rev Nephrol.* 2014;10:116–22.
- Lee T, Thamer M, Zhang Q, Zhang Y, Allon M. Vascular access type and clinical outcomes among elderly patients on hemodialysis. *Clin J Am Soc Nephrol.* 2017;12:1823–30.
- Nadeau-Fredette AC, Goupil R, Montreuil B, Carignan A, Leblanc M. Arteriovenous fistula for the 80 years and older patients on hemodialysis: is it worth it? *Hemodial Int.* 2013;17:594–601.
- Lok CE, Huber TS, Lee T, Shenoy S, Yevzlin AS, Abreo K. KDOQI clinical practice guideline for vascular access: 2019 update. *Am J Kidney Dis.* 2020;75 Suppl. 2:S1–164.
- Ko JG, Rhee MC, Obi Y, Chang TI, Soohoo M, Kim TW, et al. Vascular access placement and mortality in elderly incident hemodialysis patients. *Nephrol Dial Transplant.* 2020;35:503–11.
- Wasse H, Speckman RA, Frankenfield DL, Rocco MV, McClellan WM. Predictors of delayed transition from central venous catheter use to permanent vascular access among ESRD patients. *Am J Kidney Dis.* 2007;49:276–83.
- Richardson AI, Leake A, Schmieder GC, Biuckians A, Stokes GK, Panneton JM, et al. Should fistulas really be first in the elderly patient? *J Vasc Access.* 2009;10:199–202.
- Hod T, Patibandla B, Vin T, Brown RS, Goldfarb-Rumyantzev AS. Arteriovenous fistula placement in the elderly: when is the optimal time? *J Am Soc Nephrol.* 2015;26:448–56.
- Rostami Z. When to start dialysis in elderly patients. *Nephro-Urol Month.* 2013;5:855–7.
- Chao CT, Tsai HB, Lin YF, Ko WJ. Acute kidney injury in the elderly: only the tip of the iceberg. *J Clin Gerontol Geriatr.* 2014;7–12.
- Moist L, Lok CE, Vachharajani TJ, Xi W, AlJaishi A, Polkinghorne KR, et al. Optimal vascular access in the elderly patient. *Semin Dial.* 2012;25:640–8.
- Lomonte C, Basile C, Mitra S, Combe C, Covic A, Davenport A, et al. Should a fistula first policy be revisited in elderly hemodialysis patients? *Nephrol Dial Transplant.* 2019;34:1636–43.
- Lee T, Thamer M, Zhang Y, Zhang Q, Allon M. Outcomes of elderly patients after predialysis vascular access creation. *J Am Soc Nephrol.* 2015;26:3133–40.
- Vachharajani TJ, Moossavi S, Jordan JR, Vachharajani V, Freedman BI, Burkart JM. Re-evaluating the fistula first initiative in octogenarians on hemodialysis. *Clin J Am Soc Nephrol.* 2011;6:1663–7.
- Escoli R, Luz I, Santos P, Vila Lobos A. Predialysis vascular access creation: to whom and when. *Port J Nephrol Hypert.* 2017;31:162–6.
- Xue JL, Dahl D, Ebben JP, Collins AJ. The association of initial hemodialysis access type with mortality outcomes in elderly Medicare ESRD patients. *Am J Kidney Dis.* 2003;42:1013–9.
- Yeh LM, Chiu SYH, Lai PC. The impact of vascular access types on hemodialysis patient long-term survival. *Sci Rep Vol.* 2019;9:10708.
- Kawaguchi T, Tong L, Robinson BM, Sen A, Fukuhara S, Kurokawa K. C-reactive protein and mortality in hemodialysis patients: The Dialysis Outcomes and Practice Patterns Study (DOPPS). *Nephron Clin Pract.* 2011;117:167–78.
- Mutsert R, Grootendorst DC, Indemans F, Boeschoten EW, Krediet RT, Dekker FW. Association between serum albumin and mortality in dialysis patients is partly explained by inflammation, and not by malnutrition. *J Ren Nut.* 2009;19:127–35.
- Ravani P, Quinn R, Oliver M. Examining the association between hemodialysis access type and mortality: the role of access complications. *Clin J Am Soc Nephrol.* 2017;12:955–64.
- Brown RS, Patibandla BK, Goldfarb-Rumyantzev AS. The survival benefit of “fistula first, catheter last” in hemodialysis is primarily due to patient factors. *J Am Soc Nephrol.* 2017;28:645–52.