




ORIGINAL ARTICLE

Survival on four compared with three times per week haemodialysis in high ultrafiltration patients: an observational study

James Fotheringham ^{1,2}, Nicholas Latimer², Marc Froissart³, Florian Kronenberg⁴, Peter Stenvinkel⁵, Jürgen Floege⁶, Kai-Uwe Eckardt⁷ and David G. Wheeler^{8,9}

¹Sheffield Kidney Institute, Sheffield Teaching Hospitals, Sheffield, UK, ²School of Health and Related Research, University of Sheffield, Sheffield, UK, ³Clinical Trial Unit, Centre Hospitalier Universitaire Vaudois, Lausanne, Switzerland, ⁴Institute of Genetic Epidemiology, Medical University of Innsbruck, Innsbruck, Austria, ⁵Department of Renal Medicine, Karolinska University Hospital Huddinge, Stockholm, Sweden, ⁶Department of Nephrology, RWTH Aachen University Hospital, Aachen, Germany, ⁷Department of Nephrology and Medical Intensive Care, Charité–Universitätsmedizin-Berlin, Berlin, Germany, ⁸Department of Renal Medicine, University College London, UK and ⁹George Institute for Global Health, Sydney, New South Wales, Australia

Correspondence to: James Fotheringham; E-mail: j.fotheringham@sheffield.ac.uk

ABSTRACT

Background. The harm caused by the long interdialytic interval in three-times-per-week haemodialysis regimens (3×WHD) may relate to fluid accumulation and associated high ultrafiltration rate (UFR). Four-times-per-week haemodialysis (4×WHD) may offer a solution, but its impact on mortality, hospitalization and vascular access complications is unknown.

Methods. From the AROii cohort of incident in-centre haemodialysis patients, 3×WHD patients with a UFR >10 mL/kg/h were identified. The hazard for the outcomes of mortality, hospitalization and vascular access complications in those who switched to 4×WHD compared with staying on 3×WHD was estimated using a marginal structural Cox proportional hazards model. Adjustment included baseline patient and treatment characteristics with inverse probability weighting used to adjust for time-varying UFR and cardiovascular comorbidities.

Results. From 10 637 European 3×WHD patients, 3842 (36%) exceeded a UFR >10 mL/kg/h. Of these, 288 (7.5%) started 4×WHD and at baseline were more comorbid. Event rates while receiving 4×WHD compared with 3×WHD were 12.6 compared with 10.8 per 100 patient years for mortality, 0.96 compared with 0.65 per year for hospitalization and 14.7 compared with 8.0 per 100 patient years for vascular access complications. Compared with 3×WHD, the unadjusted hazard ratio (HR) for mortality on 4×WHD was 1.05 [95% confidence interval (CI) 0.78–1.42]. Following adjustment for baseline demographics, time-varying treatment probability and censoring risks, this HR was 0.73 (95% CI 0.50–1.05; P = 0.095). Despite these adjustments on 4×WHD, the HR for hospitalization remained elevated and vascular access complications were similar to 3×WHD.

Received: 13.7.2020; Editorial decision: 26.10.2020

© The Author(s) 2020. Published by Oxford University Press on behalf of ERA-EDTA.

This is an Open Access article distributed under the terms of the Creative Commons Attribution Non-Commercial License (<http://creativecommons.org/licenses/by-nc/4.0/>), which permits non-commercial re-use, distribution, and reproduction in any medium, provided the original work is properly cited. For commercial re-use, please contact journals.permissions@oup.com

Conclusions. This observational study was not able to demonstrate a mortality benefit in patients switched to 4×WHD. To demonstrate the true benefits of 4×WHD requires a large, well-designed clinical trial. Our data may help in the design of such a study.

Keywords: four-times-per-week haemodialysis, hospitalization, survival analysis, ultrafiltration, vascular access

INTRODUCTION

For the majority of in-centre haemodialysis (HD) patients, a three-times-per-week HD (3×WHD) schedule is unphysiological with interdialytic intervals of between 48 and 72 h. Harm associated with the accumulation of uraemic toxics, potassium and fluid may manifest in increases in arrhythmias [1], cardiac failure [2], hospitalization, mortality and symptom burden [3–5]. More intensive HD theoretically mitigates some of these harms. Observational data suggest that 3×WHD with a longer session length is associated with improved overall survival, but increases in mortality after the long interdialytic interval persist [6]. Obtaining outcome data on benefits of more frequent dialysis through clinical trials has proven challenging [7]. Six-times-per-week HD may be clinically beneficial and cost effective but is only practical in a subgroup of patients [8–11].

Recent analyses suggest that compared with the short interdialytic interval, the longer 2-day interdialytic interval is associated with increases in hospitalization and mortality of 80–100% [12]. Specifically targeting this period with an additional dialysis session may be attractive both to providers and to patients: one-fifth of patients who report being bothered by their fluid restriction state that they would accept an additional HD session if offered it [13]. Four-times-per-week HD (4×WHD) is the third most commonly prescribed HD frequency after three- and two-times-per-week schedules; however, previous evaluations have combined 4×WHD with other augmented regimes such as extended-hours dialysis, making the impact of the discrete removal of the long interdialytic interval challenging [14].

We present a target trial observational data analysis where a large dataset was used to emulate the desired clinical trial with inclusion and exclusion criteria, follow-up time, adherence and endpoints reflecting benefits and harms [15, 16]. Identifying high-risk individuals using the ultrafiltration rate (UFR) representing the clinical driver to initiate 4×WHD [17], it employs marginal structural modelling [18] to address the relationship between UF and the risk of initiating of 4×WHD and mortality, which may lead to biased estimates of the benefits of treatment. This study design recognizes the clinical indications to offer a 3×WHD patient a regular additional session, appropriately factoring in the variables that are associated with the time-varying risk of events the clinician and patient are trying to avoid.

MATERIALS AND METHODS

Cohort and data

The Analyzing Data, Recognizing Excellence and Optimizing Outcomes (ARO) cohort was a prospective observational cohort study of electronic medical records capturing anonymized longitudinal individual-level data for incident HD patients enrolled at 1 of the 312 Fresenius Medical Care (FMC) facilities across 15 European countries between 2007 and 2009 and followed up until the end of 2014. All local ethical and regulatory obligations concerning patient data for each of the 15 participating countries were met. These approvals encompass subsequent

analyses including those described here. Informed consent was obtained from all patients by FMC (Europe).

Data on demographics, comorbidities, laboratory results, hospitalizations, mortality and individual HD sessions were captured [19]. The presence of 11 comorbid conditions was identified using International Classification of Diseases, Tenth revision (ICD-10) codes from administrative data using existing schema (ischaemic heart disease, congestive heart failure, cerebrovascular accident, peripheral vascular disease, other cardiac disease, chronic obstructive pulmonary disease, gastrointestinal bleeding, liver disease, dysrhythmia, cancer and diabetes [20]). Hospitalization was defined as an admission to hospital lasting at least 1 day. Within these hospitalizations, ICD-10 codes for vascular access complications were identified (see [Supplementary data, Table S1](#), for associated ICD-10 codes).

Eligibility, exposure, adherence and follow-up

The inclusion criteria of an UFR >10 mL/kg/h was based on the progressive mortality increase associated with this range of UF [17]. Patients were classed as eligible for 4×WHD and included in the analysis from the point they exceeded this UFR across three HD sessions while prescribed 3×WHD. The UFR was calculated for each session from the recorded difference in pre- and post-dialysis weights divided by treatment time in minutes. There were no clinical exclusion criteria. Exposure to 4×WHD was classed as receiving four HD sessions in week 1, four sessions per week 4 weeks later and a further four sessions per week 2 weeks after that. This was in order to address any incorrect identification of 4×WHD associated with rescheduling HD for elective admissions or *ad hoc* HD to address specific issues. Adherence was assessed while receiving 3×WHD and 4×WHD using the mean number of delivered sessions per week. Attendance for HD was defined as the presence of a recording for blood pressure, pre-dialysis weight and HD treatment time. Follow-up was up to 3 years from the date of first eligibility or until censoring for transplantation, moving to a non-FMC facility, changing dialysis modality, withdrawal of consent or death.

Statistical methods

During the course of the analysis, an individual's dialysis frequency (moving from 3×WHD to 4×WHD) and the clinical parameters that are associated with this change in frequency and also the outcomes (e.g. increasing UFR) vary. Marginal structural Cox proportional hazards models were used to estimate the time-varying association between dialysis frequency (4×WHD versus 3×WHD) and the endpoints of mortality, hospitalization and hospitalization for vascular access complications. Marginal structural models are designed to account for confounding introduced by time-varying clinical parameters and their response to treatment over time by weighting an individual patient's observations to create a pseudo-population where time-varying covariates are more evenly distributed between treatment arms. They still assume there are no unmeasured confounders. First, logistic regression was used to obtain

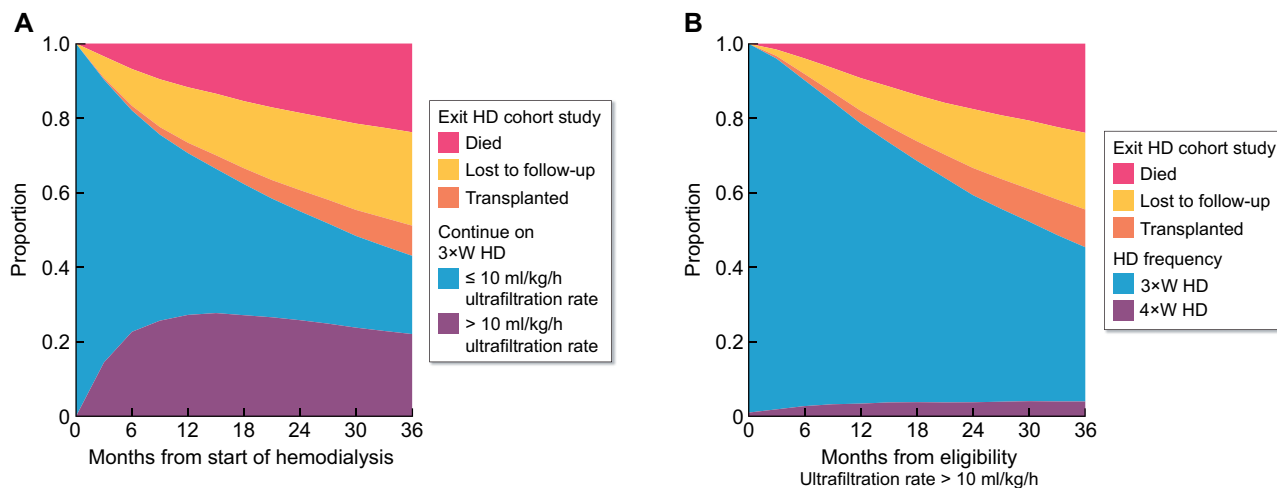


FIGURE 1: (A) The prevalence of patients meeting the inclusion criteria (10 mL/kg/h UFR) for the four-times-per-week target trial. (B) The proportion of patients meeting the inclusion criteria who go on to receive 4xWHD, and other competing events.

probabilities of treatment (switching from 3xWHD to 4xWHD) and censoring (transplantation or being lost to follow-up) for each month from inclusion in the study by meeting the eligibility until the end of follow-up [18, 21]. Baseline covariates (comorbidities at eligibility, achieved dialysis session duration, dialysis catheter use, equilibrated Kt/V, time on dialysis, age, serum phosphate level and post-dialysis weight) and time-varying covariates (UFR, systolic blood pressure and the comorbidities of congestive heart failure and ischaemic heart disease, which varied within individuals during their follow-up) were included. Continuous variables were split by quantiles into five equal groups. Probabilities from these logistic regression models were converted into weights by dividing the probabilities estimated from the baseline covariates (numerator) by the probabilities estimated using the baseline and time-varying covariates (denominator). Treatment and censoring weights were calculated separately and multiplied together, resulting in stabilized weights with a mean of 1.006 [standard deviation (SD) 0.174]. The final marginal structural models were adjusted for baseline covariates because they appeared in both the numerator and the denominator of the stabilized weights [18, 21]. The time-varying data include a variable reflecting if the patient is receiving 3xWHD or 4xWHD and weights were set to 1 following the initiation of 4xWHD [18]. The hazard for the endpoints associated with the time-varying exposure to 4xWHD is reported following sequential adjustment: (1) adjusted for baseline covariates, which leaves residual confounding, because clinical parameters and their response to treatment vary over time; (2) adjusted for time-varying covariates using inverse probability weighting for treatment; and (3) employing weighting to address the time-varying risk of being censored by transplantation or lost to follow-up, which may be associated with the outcomes of interest. Proportional hazards assumptions were assessed graphically using Schoenfeld residuals. When time-varying laboratory and HD data were missing we used the last-observation-carried forward approach (2.97% and 1.42% of patient follow-up time beyond 35 and 3 days, respectively); as in clinical practice, decisions to initiate 4xWHD may be made from historical observations. Patients with missing data at the time of death were excluded entirely, which only affected 14 patients who never received 4xWHD.

In the main analysis, patients who were exposed to 4xWHD were treated as receiving this until the end of follow-up, as adverse consequences of 4xWHD (which may manifest once a patient returns to 3xWHD) would then be captured. A sensitivity analysis was performed to explore any modification of the effect of 4xWHD accounting for patients who receive the treatment but return to 3xWHD, whereby patients who moved from 4xWHD back to 3xWHD were treated as on 3xWHD from 3 months after the observed treatment switch. Reverting to 3xWHD was defined as 3 consecutive non-hospitalized weeks receiving ≤ 3 sessions per week. This lagged per-protocol analysis assigned any events occurring within the 3 months of switching from 4xWD back to 3xWHD to the 4xWHD treatment. Statistical analyses were conducted in R version 3.4.4 (R Foundation for Statistical Computing, Vienna, Austria).

RESULTS

Patient characteristics

A total of 10 637 incident HD patients were screened for inclusion in the study, with 4009 (37.7%) patients meeting the inclusion criteria of an average UFR across three HD sessions > 10 mL/kg/h following a median of 6.1 months of HD. Figure 1A shows the proportion of patients on 3xWHD with a UFR < 10 mL/kg/h, 3xWHD with a UFR > 10 mL/kg/h (eligible for 4xWHD), transplanted, lost to follow-up or died. The prevalence of patients eligible for 4xWHD stabilizes at 20–25% following 12 months of HD.

Having exceeded a UFR of 10 mL/kg/h and classed as eligible for 4xWHD, 7.5% of patients subsequently went on to receive 4xWHD and were suitable for analysis. The prevalence of patients receiving 4xWHD stabilized at 4% after ~12 months, as new patients commenced 4xWHD and patients on 4xWHD left the study (Figure 1B). Patient flow through the screening process, inclusion criteria, subsequent treatment and inclusion in the analysis are shown in Figure 2. The demographics of patients who remained on 3xWHD or subsequently went onto 4xWHD having met the inclusion criteria are illustrated in Table 1, showing baseline differences in age (62.5 versus 60.8 years), diabetes (34.7% versus 48.3%), weight (64.5 and

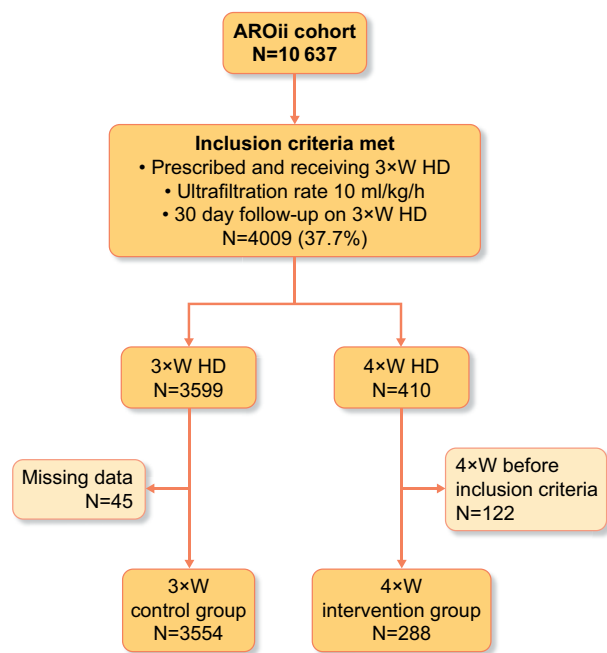


FIGURE 2: Incident patients screened throughout their follow-up and their flow through the analysis.

68.2 kg) and heart failure (16.4% versus 21.5%). The numbers of patients from each country are available in the [Supplementary Materials](#).

Both at baseline and during follow-up, there were differences in the prevalence of cardiovascular comorbidity and the proportion of patients who were transplanted or lost to follow-up between the time patients received 3xWHD (Figure 3A) and 4xWHD (Figure 3B). These time-varying associations support the use of inverse probability weighting methods for treatment and censoring. The demographics of patients according to their reasons for exiting the study are listed in [Supplementary data, Table S2](#).

Treatment initiation and adherence

In those who received it, 4xWHD was initiated a median of 12.5 months from the time patients first became eligible. From 6 weeks following the initiation of 4xWHD treatment, the median time from the initiation of 4xWHD to the end of follow-up or 3 consecutive weeks receiving less than four sessions per week was 6.1 months. For the duration of follow-up, the average number of sessions delivered per week, excluding hospitalized time, was 3.46 sessions per week in the 4xWHD arm and 2.96 sessions per week in the 3xWHD arm. The mean session duration was 229 min while receiving 4xWHD and 235 min while receiving 3xWHD.

Associations with mortality, hospitalization and vascular access complications

The crude mortality rate during follow-up was 10.8 per 100 patient years [95% confidence interval (CI) 10.1–11.6] while receiving 3xWHD and 12.6 per 100 patient years (95% CI 9.3–16.7) following the start of 4xWHD. The univariate hazard for survival while receiving 4xWHD compared with 3xWHD was 1.05 (95% CI 0.78–1.42; $P=0.735$). The hazard ratio (HR) was 0.90 (95% CI 0.65–1.24; $P=0.518$) after adjustment for baseline covariates

and the final multivariable adjusted HR was 0.81 (95% CI 0.58–1.14; $P=0.229$) after adjusting for time-varying covariates using inverse probability weighting for treatment. After weighting for censoring risk, this multivariable HR was 0.73 (95% CI 0.50–1.06; $P=0.096$). These sequential adjustments are shown in [Figure 4A](#).

The crude hospitalization rate during follow-up was 0.65 (95% CI 0.64–0.67) per patient year while receiving 3xWHD and was 0.96 (95% CI 0.86–1.06) per patient year while receiving 4xWHD [univariate HR 1.51 (95% CI 1.28–1.77), $P=0.008$]. Following adjustment for baseline and time-varying factors influencing treatment and censoring, 4xWHD had a multivariable HR of 1.28 (95% CI 1.06–1.53; $P=0.008$).

The crude vascular access complication rate was 8.0 (95% CI 7.3–8.7) per 100 patient years while receiving 3xWHD and was 14.7 (95% CI 11.1–19.1) per 100 patient years while receiving 4xWHD [univariate HR 1.91 (95% CI 1.50–2.61), $P<0.001$]. Sequential adjustments resulted in a final multivariable HR associated with 4xWHD of 1.15 (95% CI 0.78–1.72; $P=0.478$) and are shown in [Figure 4B and C](#).

The sensitivity analysis exploring the medium-term impact of switching from 4xWHD back to 3xWHD did not significantly alter the effect sizes for the HR associated with the 4xWHD treatment strategy ([Figure 5](#)).

DISCUSSION

This study is the first to explore the association between 4xWHD and endpoints important to clinicians and patients [15, 18, 22]. Thirty-six percent of 3xWHD patients experienced the high UFRs associated with myocardial stunning and increased mortality [17, 23] and are the subjects of performance indicators for dialysis clinics [24]. Despite this, only 7.5% went on to receive 4xWHD for at least 6 weeks. Patient characteristics predicting mortality and transplantation were less favourable in those who received 4xWHD, and accounting for this, 4xWHD was not associated with an improvement in survival. Vascular access complications were comparable following adjustment and hospitalization remained elevated when compared with patients with high UF on 3xWHD.

The favourable survival HR of 0.73 (95% CI 0.50–1.05; $P=0.095$) observed with 4xWHD should be cautiously interpreted alongside HRs of 0.54, 3.88 and 0.91 for the 12-month interventions in the Frequent Hemodialysis Network (FHN) Frequent, Nocturnal and ACTIVE (A Clinical Trial of Intensive Dialysis) studies, respectively [8, 25, 26]. Statistical bodies and prominent journals recommend that estimates of effects and their margins of error should be interpreted together to inform clinicians and regulatory agencies regarding an intervention, with less reliance on the absolute P -value [27–29]. One of the primary reasons to initiate 4xWHD is to modify the increase in mortality after the 2-day break in 3xWHD [3, 4]. If this short-term increase is reduced to that of the rest of the week compared with a dialysis week with this short-term increase present, the HR is only 0.88 ([Appendix 1 of Supplementary Materials](#)).

The potential mechanisms through which the 4xWHD schedule improves outcomes could be through reductions in the UFR and hyperkalaemia, lower time-averaged volume overload and myocardial stunning and reduced arrhythmias in the build up to and during the first HD session of the dialysis week [1, 17, 23, 30, 31]. A previous 12-month randomized trial of alternate-day dialysis with no long interdialytic interval showed improvements in left ventricular mass and systolic blood

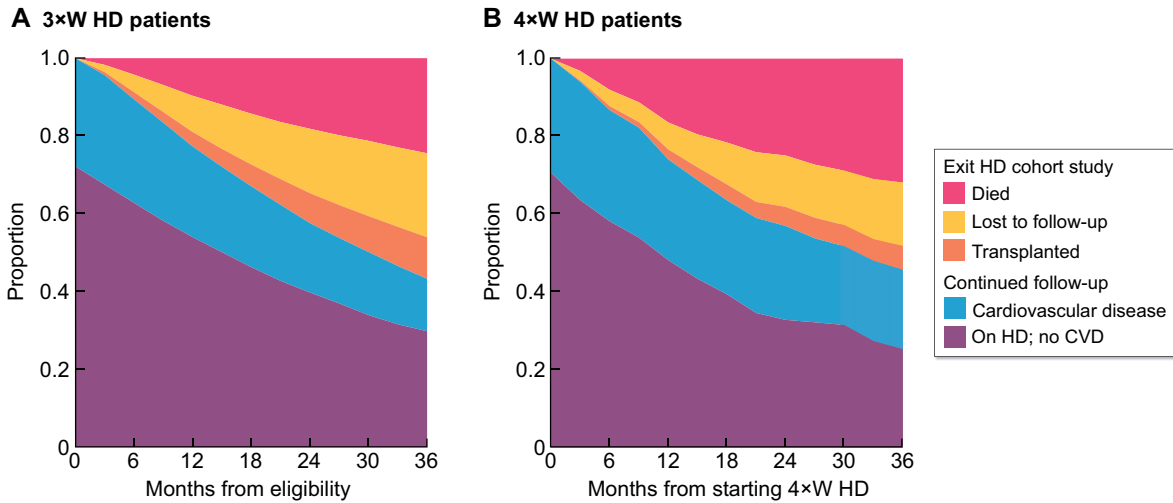


FIGURE 3: The prevalence of comorbid conditions and censoring events according to dialysis frequency: (A) 3xWHD and (B) 4xWHD.

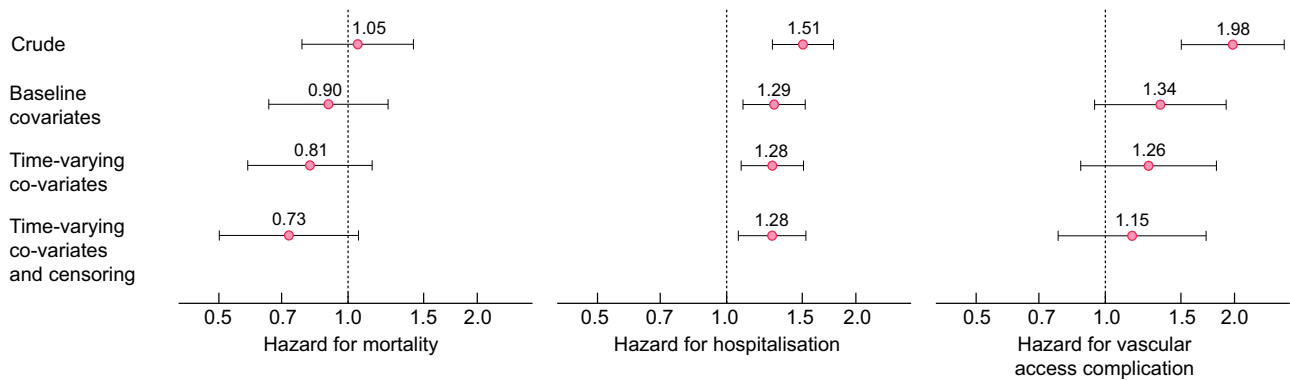


FIGURE 4: HR of 4xWHD compared with 3xWHD for the endpoints of mortality, hospitalization and vascular access complication. Adjustment for baseline covariates (comorbidities at eligibility, dialysis session duration, dialysis access type, equilibrated Kt/V, time on dialysis, age, serum phosphate level and post-dialysis weight) and time-varying covariates (UFR, systolic blood pressure and the comorbidities of congestive heart failure and ischaemic heart disease) are sequentially reported.

pressure compared with 3xWHD [32]. The extended-hours intervention (predominantly extended sessions 3xWHD) in the ACTIVE trial showed reductions in left ventricular mass in those individuals who had a reduction in UFR, although UFR and patient survival were not improved in those randomized to extended hours [26, 33]. However, augmented HD may be associated with potential harms: the FHN short daily trial was associated with improved survival in contrast to the nocturnal study's inferior survival [8, 25], and both tended towards a higher incidence of vascular access complications in the intervention arms [9, 34]. In our analysis, the increase in vascular access events in patients receiving 4xWHD was largely mitigated following adjustment, suggesting the patient characteristics that predict the vascular access complications and the clinical need for 4xWHD are similar.

The strengths of this analysis include the incident nature of the cohort and the use of highly granular data to define inclusion criteria, exposure to the intervention, adherence and outcomes, further capitalized on by the target trial methodology and marginal structural modelling approach. Although less prone to bias, the limitation of these methods is that they deal with observed confounders and residual unobserved differences between patients, which could introduce bias. Other

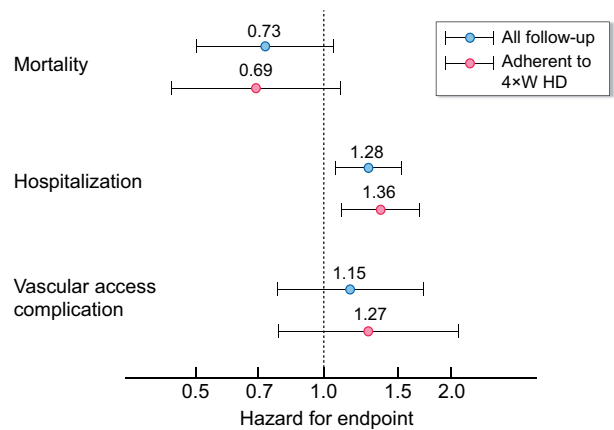


FIGURE 5: Sensitivity analysis comparing the HR for 4xWHD compared with thr3xWHD when patients who return to 3xWHD are treated as receiving this schedule from 3 months after the switch.

weaknesses include the absence of information on residual kidney function and quality of life. Our definition of 4xWHD means

Table 1. Demographics of patients at inclusion (10 mL/kg/hr UFR) and at exposure to 4×WHD

Characteristics	3×WHD at eligibility	4×WHD at eligibility	4×WHD at initiation
Patients, n	3554	288	288
Age (years), mean (SD)	62.5 (14.9)	60.8 (15.3)	62.40 (15.37)
Male, n (%)	2148 (60.4)	159 (55.2)	159 (55.2)
Days on dialysis, mean (SD)	355.2 (417.0)	221.7 (221.6)	803.9 (596.6)
Ischaemic heart disease, n (%)	563 (15.8)	40 (13.9)	52 (18.1)
Cancer, n (%)	260 (7.3)	22 (7.6)	26 (9.0)
Heart failure, n (%)	582 (16.4)	62 (21.5)	71 (24.7)
Chronic obstructive pulmonary disease, n (%)	234 (6.6)	20 (6.9)	23 (8.0)
Cerebrovascular disease, n(%)	350 (9.8)	34 (11.8)	40 (13.9)
Depression, n (%)	69 (1.9)	8 (2.8)	8 (2.8)
Diabetes, n (%)	1232 (34.7)	139 (48.3)	142 (49.3)
Arrhythmia, n (%)	335 (9.4)	37 (12.8)	50 (17.4)
Gastrointestinal disease, n (%)	76 (2.1)	5 (1.7)	6 (2.1)
Liver disease, n (%)	132 (3.7)	13 (4.5)	15 (5.2)
Other cardiac disease, n (%)	39 (1.1)	5 (1.7)	7 (2.4)
Peripheral vascular disease, n (%)	542 (15.3)	43 (14.9)	68 (23.6)
Ultrafiltration volume (L), mean (SD)	2.72 (1.00)	2.92 (0.80)	2.48 (1.01)
UFR (mL/kg/h), mean (SD)	11.00 (3.59)	11.37 (2.83)	9.61 (3.72)
Equilibrated Kt/V, mean (SD)	1.42 (0.30)	1.36 (0.30)	1.44 (0.29)
Phosphate, mean (SD)	1.55 (0.47)	1.61 (0.48)	1.54 (0.52)
HD session duration (min), mean (SD)	231.91 (16.98)	228.95 (17.46)	230.59 (20.50)
Loop diuretic use, n(%)	329 (9.3)	34 (11.8)	37 (12.8)
Weight (kg), mean (SD)	64.48 (12.83)	68.22 (14.13)	68.09 (14.24)
Systolic blood pressure (mmHg), mean (SD)	137.16 (22.90)	141.10 (23.96)	138.81 (25.58)
Diastolic blood pressure (mmHg), mean (SD)	71.17 (13.72)	71.64 (13.96)	69.42 (14.71)

that those who did not adhere within the first 6 weeks were not included. The median duration of 4×WHD was 6 months before the end of follow-up and we are unable to say with confidence what outcomes might be associated with longer treatment or sustained adherence. Our per-protocol analysis excluding follow-up time after 4×WHD patients returned to 3×WHD yielded similar effect sizes and statistical significance as the main analysis and suggests a legacy effect that was observed in the FHN daily trial but not the ACTIVE trial [8, 26].

Building on the existing observational data on UFR, this study could be used to advise patients on potential interventions once a UF threshold of 10 mL/kg/h is reached. However, in our study it took up to a year from patients meeting this threshold to the initiation of 4×WHD, suggesting other factors may inform the decision-making process, such as struggling with fluid restriction and subsequent hospitalization for fluid overload [6]. The mean duration of 6 months for 4×WHD suggests that some clinicians are using this treatment in response to subacute issues that then resolve. The more widespread sustained adoption of 4×WHD would have staffing and capacity implications, which may be offset by the increasing adoption of incremental HD start, with schedules of less than three sessions per week [35]. Other capacity-generating initiatives such as shared- and self-care HD programmes and the more widespread use of home HD may offer solutions [36]. Taken together with the finding of other augmented HD clinical trials [7, 10, 37], our results could help in the design of a prospective trial evaluation of 4×WHD: to demonstrate this HR with an 80% power, $\alpha = 0.05$ and a 10% transplantation rate at 3 years would require 833 patients per arm, improving to 479 per arm by relaxing α (0.1) and significance to one-sided [38]. Sample sizes could be further reduced by adjustment for baseline variables such as cardiac failure [39], and stratification by suitability for transplantation should be considered.

With the mounting evidence of the range of harms associated with a long interdialytic interval, this study contributes to the supporting evidence for potential solutions; however, appropriately designed studies are required to ensure they are both clinically and cost effective, sustainable and acceptable to the patient.

DATA AVAILABILITY STATEMENT

The data underlying this article will be shared upon reasonable request to the corresponding author and approval of the ARO steering group.

SUPPLEMENTARY DATA

Supplementary data are available at [ndt online](https://ndt.oxfordjournals.org/).

ACKNOWLEDGEMENTS

The authors would like to thank Bruno Fouqueray and Karly Louie from Amgen for their assistance with this analysis.

FUNDING

The ARO CKD Research Initiative is a joint observational research commitment from Amgen and FMC (Europe), fully funded by Amgen (Europe), Rotkreuz, Switzerland. This article presents independent research funded by the National Institute for Health Research (NIHR). The views expressed are those of the authors and not necessarily those of the National Health Service, the NIHR or the Department of Health and Social Care. J.F. has received speaker honoraria from FMC and conducts research funded by Vifor Pharma

and Novartis. This specific research is funded by a National Institute for Health Research Clinician Scientist Fellowship awarded to J.F. (CS-2015-15-008). J.F. has received consultancy honoraria from Amgen, Fresenius and Vifor. D.C.W. has received consultancy fees from Amgen, AstraZeneca, Bayer, Boehringer Ingelheim, Janssen, GlaxoSmithKline, Mundipharma, Mitsubishi, Napp and Vifor Fresenius. Results presented in this article have not been published previously in whole or part, except in abstract format.

CONFLICT OF INTEREST STATEMENT

None declared.

REFERENCES

- Roy-Chaudhury P, Tumlin JA, Koplan BA et al. Primary outcomes of the monitoring in dialysis study indicate that clinically significant arrhythmias are common in hemodialysis patients and related to dialytic cycle. *Kidney Int* 2018; 93: 941–951
- Obokata M, Negishi K, Marwick TH et al. Comparison of different interdialytic intervals among hemodialysis patients on their echocardiogram-based cardiovascular parameters. *Am Heart J* 2015; 169: 523–530.e2
- Foley RN, Gilbertson DT, Murray T et al. Long interdialytic interval and mortality among patients receiving hemodialysis. *N Engl J Med* 2011; 365: 1099–1107
- Fotheringham J, Fogarty DG, El Nahas M et al. The mortality and hospitalization rates associated with the long interdialytic gap in thrice-weekly hemodialysis patients. *Kidney Int* 2015; 88: 569–575
- Ju A, Unruh M, Davison S et al. Establishing a core outcome measure for fatigue in patients on hemodialysis: a Standardized Outcomes in Nephrology–Hemodialysis (SONG-HD) consensus workshop report. *Am J Kidney Dis* 2018; 72: 104–112
- Fotheringham J, Sajjad A, Stel VS et al. The association between longer haemodialysis treatment times and hospitalization and mortality after the two-day break in individuals receiving three times a week haemodialysis. *Nephrol Dial Transplant* 2019; 34: 1577–1584
- Dember LM, Lacson E, Brunelli SM et al. The TiME trial: a fully embedded, cluster-randomized, pragmatic trial of hemodialysis session duration. *J Am Soc Nephrol* 2019; 30: 890–903
- Chertow GM, Levin NW, Beck GJ et al. Long-term effects of frequent in-center hemodialysis. *J Am Soc Nephrol* 2016; 27: 1830–1836
- Chertow GM, Levin NW, Beck GJ et al. In-center hemodialysis six times per week versus three times per week. *N Engl J Med* 2010; 363: 2287–2300
- Sergeyeva O, Gorodetskaya I, Ramos R et al. Challenges to enrollment and randomization of the Frequent Hemodialysis Network (FHN) Daily Trial. *J Nephrol* 2012; 25: 302–309
- Liu FX, Treharne C, Arici M et al. High-dose hemodialysis versus conventional in-center hemodialysis: a cost-utility analysis from a UK payer perspective. *Value Health* 2015; 18: 17–24
- Fotheringham J, Smith MT, Froissart M et al. Hospitalization and mortality following non-attendance for hemodialysis according to dialysis day of the week: a European cohort study. *BMC Nephrol* 2020; 21: 218
- Flythe JE, Mangione TW, Brunelli SM et al. Patient-stated preferences regarding volume-related risk mitigation strategies for hemodialysis. *Clin J Am Soc Nephrol* 2014; 9: 1418–1425
- Marshall MR, Polkinghorne KR, Kerr PG et al. Intensive hemodialysis and mortality risk in Australian and New Zealand populations. *Am J Kidney Dis* 2016; 67: 617–628
- Hernan MA, Robins JM. Using big data to emulate a target trial when a randomized trial is not available. *Am J Epidemiol* 2016; 183: 758–764
- Hernán MA, Robins JM. Per-protocol analyses of pragmatic trials. *N Engl J Med* 2017; 377: 1391–1398
- Flythe JE, Kimmel SE, Brunelli SM. Rapid fluid removal during dialysis is associated with cardiovascular morbidity and mortality. *Kidney Int* 2011; 79: 250–257
- Fewell Z, Wolfe F, Choi H et al. Controlling for time-dependent confounding using marginal structural models. *Stata J* 2004; 4: 402–420
- Steil H, Amato C, Carioni C et al. EuCliD—a medical registry. *Methods Inf Med* 2004; 43: 83–88
- Liu J, Huang Z, Gilbertson DT et al. An improved comorbidity index for outcome analyses among dialysis patients. *Kidney Int* 2010; 77: 141–151
- Cole SR, Hernan MA. Constructing inverse probability weights for marginal structural models. *Am J Epidemiol* 2008; 168: 656–664
- Evangelidis N, Tong A, Manns B et al. Developing a set of core outcomes for trials in hemodialysis: an international Delphi survey. *Am J Kidney Dis* 2017; 70: 464–475
- Burton JO, Jefferies HJ, Selby NM et al. Hemodialysis-induced cardiac injury: determinants and associated outcomes. *Clin J Am Soc Nephrol* 2009; 4: 914–920
- Kramer H, Yee J, Weiner DE et al. Ultrafiltration rate thresholds in maintenance hemodialysis: an NKF-KDOQI controversies report. *Am J Kidney Dis* 2016; 68: 522–532
- Rocco MV, Daugirdas JT, Greene T et al. Long-term effects of frequent nocturnal hemodialysis on mortality: the Frequent Hemodialysis Network (FHN) Nocturnal Trial. *Am J Kidney Dis* 2015; 66: 459–468
- Smyth B, Zuo L, Gray NA et al. No evidence of a legacy effect on survival following randomization to extended hours dialysis in the ACTIVE Dialysis trial. *Nephrology* 2020; 25: 792–800.
- Wasserstein RL, Lazar NA. The ASA statement on p-values: context, process, and purpose. *Am Stat* 2016; 70: 129–133
- Wasserstein RL, Schirm AL, Lazar NA. Moving to a world beyond “ $p < 0.05$ ”. *Am Stat* 2019; 73(Suppl 1): 1–19
- Harrington D, D’Agostino RB, Gatsonis C et al. New guidelines for statistical reporting in the *Journal*. *N Engl J Med* 2019; 381: 285–286
- Moissl U, Arias-Guillén M, Wabel P et al. Bioimpedance-guided fluid management in hemodialysis patients. *Clin J Am Soc Nephrol* 2013; 8: 1575–1582
- Brunelli SM, Du Mond C, Oestreich N et al. Serum potassium and short-term clinical outcomes among hemodialysis patients: impact of the long interdialytic interval. *Am J Kidney Dis* 2017; 70: 21–29

32. Katopodis KP, Dounousi E, Challa A et al. Switch from conventional to every other day hemodialysis: a comparison pilot study. *ASAIO J* 2009; 55: 41–46
33. Smyth B, Chan CT, Grieve SM et al. Predictors of change in left-ventricular structure and function in a trial of extended hours hemodialysis. *J Card Fail* 2020; 26: 482–491
34. Rocco MV, Lockridge RS Jr, Beck GJ et al. The effects of frequent nocturnal home hemodialysis: the Frequent Hemodialysis Network Nocturnal Trial. *Kidney Int* 2011; 80: 1080–1091
35. Kalantar-Zadeh K, Unruh M, Zager PG et al. Twice-weekly and incremental hemodialysis treatment for initiation of kidney replacement therapy. *Am J Kidney Dis* 2014; 64: 181–186
36. Fotheringham J, Barnes T, Dunn L et al. Rationale and design for SHAREHD: a quality improvement collaborative to scale up Shared Haemodialysis Care for patients on centre based haemodialysis. *BMC Nephrol* 2017; 18: 335
37. Jardine MJ, Zuo L, Gray NA et al. A trial of extending hemodialysis hours and quality of life. *J Am Soc Nephrol* 2017; 28: 1898–1911
38. Parmar MKB, Sydes MR, Morris TP. How do you design randomised trials for smaller populations? A framework. *BMC Med* 2016; 14: 183
39. Kahan BC, Jairath V, Doré CJ, Morris TP. The risks and rewards of covariate adjustment in randomized trials: an assessment of 12 outcomes from 8 studies. *Trials* 2014; 15: 139