

RESEARCH ARTICLE

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Limited referral to nephrologists from a tertiary geriatric outpatient clinic despite a high prevalence of chronic kidney disease and anaemia

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Abstract

Background: Chronic kidney disease (CKD) is increasing in prevalence world-wide with the largest growth being in the elderly. The aim of this study was to examine the prevalence of CKD in a geriatric outpatient clinic within a tertiary hospital and its association with anaemia and mortality with a focus on the referral patterns towards nephrologists.

Methods: Retrospective study utilising administrative databases. The cohort was defined as all patients that attended the geriatric outpatient clinics of a single tertiary hospital within the first 3 months of 2006. Patients were followed for 18 months for mortality and referral to a nephrologist.

Results: The mean Glomerular filtration rate (eGFR) of the 439 patients was 67.4 ± 29.1 mL/min/1.73 m² (44% <60 mL/min/1.73 m²). 11.8% had a haemoglobin < 110 g/L, with anaemia being significantly associated with kidney function in those with a eGFR < 60 mL/min/1.73 m² ($p = 0.0092$). Kidney function and anaemia were significantly associated with mortality on multivariate analysis ($p = 0.019$ and $p = 0.0074$). After 18 months, 8.8% of patients with CKD were referred to a nephrologist.

Conclusion: Despite a high prevalence of CKD in patients attending a geriatric outpatient clinic and its association with anaemia and mortality, few of these patients were referred to a nephrologist. An examination of the reasons behind this bias is required.

Keywords: Anaemia, Chronic kidney failure, Geriatrics, Referral and consultation

Background

Chronic kidney disease (CKD) is a growing problem due to our ageing population, many of who have increased comorbidities [1]. Cross-sectional studies in the general population demonstrate that CKD is seen in up to 15.3%, including 39% of those over the age of 65 years [1-3]. Previously restrictive selection criteria for commencement of dialysis in a patient have over the years been relaxed, for a number of reasons. As a consequence

prevalent dialysis numbers are increasing with its associated considerable cost [1,4].

In addition to the cost of dialysis, earlier stages of CKD are associated with increased morbidity and mortality [5,6]. A common complication, even with mild to moderate CKD, is anaemia [7,8]. This has significant effects on the quality of life of an individual that can be improved through the use of epoetins [9].

There is a perception that medical professionals caring for the elderly are potentially biased against referring patients to Nephrologists [10,11]. This may be partly due to the strict selection criteria that existed, when renal replacement therapy initially began, for the commencement of dialysis which often included an age limit [11]. In addition, some health care professionals may

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view that quality of life for their elderly patients may be better served by not going onto dialysis.

Our aim was to examine the prevalence of CKD in a tertiary Geriatrics outpatient clinic, in addition to its association with anaemia and mortality, and the referral patterns of Geriatricians to Nephrologists.

Methods

This was a retrospective study that included all patients over the age of 65 years who were reviewed in the outpatient clinics of the Department of Aged Care and Rehabilitation in Sir Charles Gairdner Hospital (SCGH) during the first quarter of 2006. This is the second largest teaching hospital within Western Australia with clinics including general geriatrics clinics, and specialised memory and falls clinics. The Department of Aged Care and Rehabilitation included 6 Specialists. There were 5 Specialists in the Department of Renal Medicine in SCGH with 20 overall in Western Australia. Patients with end-stage kidney disease were excluded. This study was approved by the hospital Human Research and Ethics Committee.

The hospital laboratory database (Pathwest) was then examined for the results of blood tests performed on these patients. In addition, medical records were reviewed. Variables recorded included demographic data, serum creatinine, haemoglobin, and iron studies. Referral to a nephrologist within the next 18 months was then recorded through review of the patients' medical records, CKD databases, and review of individual nephrologist databases. Survival after 18 months was also recorded.

Creatinine was measured using a kinetic colourimetric assay (Jaffe method) analysed on the Roche Hitachi 917 analyser (Roche Diagnostic GmbH, Mannheim, Germany). Total CV's for this assay are 6.6% at 0.8 mg/dL, and 4.1% at 5.5 mg/dL. Glomerular filtration rate (eGFR) was estimated from serum creatinine utilising the 4-variable Modification of Diet in Renal Disease study equation [12].

All variables are reported as mean \pm standard deviation, unless otherwise stated. Normality of variables was examined and if required transformation to improve normality was performed. Patients were grouped based upon their age into one of 3 categories – 65–75 years old; 75–85 years old; and over 85 years old. Simple linear regression was utilised to assess the relationship between haemoglobin and eGFR. Cox proportional hazards regression were used to examine the association between mortality and kidney function and haemoglobin on univariate and multivariate analysis. Other variables were examined to detect an association with mortality including age, gender, and iron status. Variables with a p-value < 0.10 on univariate analysis was added in the multivariate model. STATA 11.0 (Stata Corporation, College Station, Texas, USA) was used for the analysis.

Results

439 patients were reviewed during the first 3 months of 2006 and had biochemical testing performed (Table 1). A total of 867 patients were reviewed in this clinic during the baseline period. The mean age of the total 867 patients was 80.2 \pm 8.5 years, compared to 80.7 \pm 8.9 years for those with biochemical data. 38.6% of the total

Table 1 Baseline demographics

	65-75 years old (n = 87)	75-85 years old (n = 221)	>85 years old (n = 131)	All (n = 439)
Male (%)	42 (47.1%)	89 (40.3%)	44 (33.6%)	175 (39.9%)
Estimated Glomerular Filtration Rate (eGFR) (mL/min/1.73 m ²)	74.0 \pm 26.7	69.4 \pm 31.8	59.7 \pm 24.2	67.4 \pm 29.1
Chronic Kidney Disease (CKD) stage (eGFR) (%)				
1 (>90 mL/min/1.73 m ²)	33.3	17.2	11.5	18.7
2 (60–90 mL/min/1.73 m ²)	34.5	44.8	26.7	37.4
3 (30–60 mL/min/1.73 m ²)	26.4	33.9	56.5	39.2
4 (15–30 mL/min/1.73 m ²)	2.3	3.2	5.3	3.6
5 (<15 mL/min/1.73 m ²)	3.5	0.9	0	1.1
Haemoglobin (g/dL)	13.0 \pm 1.7	13.0 \pm 1.8	12.6 \pm 1.9	12.9 \pm 1.8
Mean Cell Volume (fL)	91.3 \pm 4.9	91.5 \pm 6.7	92.8 \pm 6.1	91.9 \pm 6.2
Vitamin B12 (pmol/L)	410.9 \pm 203.1	409.3 \pm 320.0	436.8 \pm 360.2	418.1 \pm 318.3
Red cell folate (nmol/L)	1087.9 \pm 653.8	1477.6 \pm 940.9	1082.1 \pm 871.3	1294.2 \pm 898.3
Ferritin (μ g/L)	251.9 \pm 491.6	221.3 \pm 342.9	169.3 \pm 228.2	212.4 \pm 345.9
Transferrin saturation (%)	21.6 \pm 9.0	17.2 \pm 10.2	16.9 \pm 8.8	17.9 \pm 9.7

population was male compared with 39.9% in those with a serum creatinine measurement.

CKD

44% of patients had a eGFR <60 mL/minute/1.73 m², with 20%, 5% and 1.1% having a eGFR <45 mL/min/1.73 m², <30 mL/min/1.73 m² and <15 mL/min/1.73 m² respectively. The mean eGFR for people aged 65–75 years was 74.0 ± 26.7 mL/min/1.73 m², 75–85 years was 69.4 ± 31.8 mL/min/1.73 m², and for those over 85 years was 59.7 ± 24.2 mL/min/1.73 m².

Anaemia

11.8% of this sample had a haemoglobin less than 110 g/L, with 4.3% having a haemoglobin less than 100 g/L. Of those with a eGFR <60 mL/min/1.73 m², 17.2% had a haemoglobin <110 g/L and 6.5% had a haemoglobin <100 g/L. Haemoglobin was not significantly associated with eGFR on linear regression in the total population (p = 0.065), but it did reach significance on examining those with a eGFR <60 mL/min/1.73 m² (Figure 1, p = 0.0092). Mean haemoglobin was similar between those aged 65–75 years, 75–85 years and over 85 years at 130 ± 17 g/L, 130 ± 17 g/L and 126 ± 19 g/L respectively. The mean ferritin was 94 ± 154 ng/mL with a transferrin saturation of 17.9 ± 9.7%.

Mortality

After 18 months of follow-up, 16.0% of the sample had died, compared to 10.3% of the total 867 patients that attended the clinics. Death was significantly associated with eGFR and anaemia on univariate analysis, p = 0.0074 and p = 0.019 respectively (Table 2). On multivariate

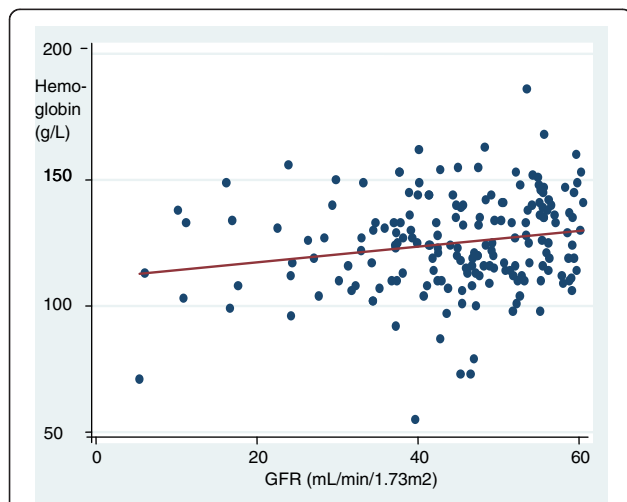


Figure 1 Scatter plot and regression line for estimated Glomerular filtration rate (eGFR) versus haemoglobin in those participants with a eGFR <60 mL/min/1.73 m².

Table 2 Univariate and multivariate logistic regression of factors associated with death.

Variable	Univariate analysis (p-value)	Multivariate analysis (p-value)
Haemoglobin (g/L)	0.02	0.03
Glomerular filtration rate (mL/min/1.73 m ²)	0.01	0.03
Age (years)	0.009	0.36
Gender	0.03	0.06

logistic regression, both eGFR and haemoglobin remained independently associated with mortality, p = 0.0047. Survival analysis indicated that CKD stage 4 was associated with a 19.7 fold increased hazard of mortality than CKD stage 1, p = 0.012.

Referral to Nephrologist

After 18 months of follow-up, 17 patients were referred to a nephrologist. This represents 8.8% of those patients with known/identified CKD on blood test. 34 patients had, at baseline, either a haemoglobin <100 g/L or a eGFR >60 mL/min/1.73 m². The mean eGFR of those patients referred to a nephrologist was 27.1 ± 14.2 mL/min/1.73 m², with 10 (58.8%) having a eGFR <60 mL/min/1.73 m² (see Table 3). Haemoglobin, age and eGFR did not predict referral to a nephrologist, p = 0.93, 0.95 and 0.055 respectively.

Discussion

Chronic kidney disease is associated with increased morbidity, mortality, haematological and biochemical abnormalities. Increasing age is associated with an increased prevalence of CKD [3]. There is evidence however that there may be a pattern of restricted referral to Nephrologists from medical practitioners caring for the elderly [10,11]. Our study demonstrated that CKD is common in patients over 65 attending a tertiary geriatrics out-patient clinic. CKD was associated with anaemia and mortality but despite this few patients were referred to a nephrologist.

Mortality has been demonstrated to be inversely associated with kidney function, with the risk increased once the eGFR is below 60 mL/min/1.73 m² [5]. Interventions that can slow progressive renal failure exist but sometimes require input from a nephrologist [13,14]. In addition a nephrologist can assist with the optimal management of mineral bone disease that also occurs with increasing prevalence once eGFR <60 mL/min/1.73 m² [15–17].

Anaemia can be directly related to CKD, with its increased risk commencing once the eGFR reaches 60 mL/min/1.73 m² or less [7,8]. This was demonstrated

Table 3 Age and Chronic Kidney Disease (CKD) stage of patients referred to a nephrologist

CKD stage (eGFR)	Age = 65–75 years	Age = 75–85 years	Age > 85 years
1 (>90 mL/min/1.73 m ²)	0%	0 %	0%
2 (60–90 mL/min/1.73 m ²)	0%	0%	0%
3 (30–60 mL/min/1.73 m ²)	0%	29.4%	23.5%
4 (15–30 mL/min/1.73 m ²)	5.9%	17.6%	5.9%
5 (<15 mL/min/1.73 m ²)	11.8%	5.9%	0%

in our study with an 11.3% prevalence of anaemia in our population. Quality of life can be improved considerably, and blood transfusions avoided, with the prescription of erythropoiesis stimulating agents [18,19]. In Australia, government subsidised prescription of these agents is restricted to Nephrologists and Haematologists.

The prevalence of CKD with a eGFR < 60 mL/min/1.73 m² in our geriatric outpatient population was 44%, with an increased rate with increasing age. This is consistently greater than the prevalence of CKD in other cohorts - 14.7% of over 65 year old residents of long-term care facilities, 15.6% in outpatient clinics, 17.5% in a community-based laboratory, and 22.4% over 65 year olds in a primary care setting [5,20-22]. Other studies have shown that a considerable proportion of patients do not receive evidence based intervention for CKD in non-nephrology practices [23]. Suggesting that referral of this population to a Nephrologist may assist in the diagnosis of the aetiology of the CKD and to ensure that optimal management was being prescribed.

Contrary to the belief of some medical practitioners, commencement of dialysis in the elderly may be associated with an improved quantity and quality of life [24]. As a consequence of multiple factors, the greatest increase in new dialysis patients in Australia and the UK are in the over 65 age group [1,25]. Late referral to a nephrologist may delay optimal preparation for dialysis and is associated with worse outcomes, prompting national and international Guidelines for early referral of CKD patients to Nephrologists [24,26,27].

Despite this there was a surprisingly low referral rate of patients with CKD to a Nephrologist, 50% of the recommended numbers based upon Kidney Health Australia guidelines of eGFR < 30 mL/min/1.73 m² or Haemoglobin < 100 g/L with eGFR = 30-60 mL/min/1.73 m² [26,28,29]. It is unclear why this practice pattern exists [28]. It may be related to a previous bias towards not dialysing the elderly or the belief that dialysis is not appropriate or in the best interests of the elderly.

The Nephrology workforce in SCGH and in the State is similar to the rest of Australia, with 72.7 end-stage kidney disease (ESKD) patients per full-time equivalent nephrologist in WA compared with 60.3 for Australia [30]. In addition the demographics of ESKD patients in WA is similar to the rest of Australia with 453 Dialysis

patients per million population, compared with 471 in Australia [31]. The peak in the age group for prevalent dialysis patients in WA and Australia is also the same at 65 to 74 years old [31]. This suggests that results from this paper may be generalizable to the rest of Australia.

Limitations include its retrospective nature. While a complete list of people attending the geriatrics outpatients clinic during the baseline period was obtained, data was only obtained on those patients that were having blood tests performed for clinical reasons. This bias is likely to overestimate the true prevalence rates. Additionally, an examination was not made to see if the CKD patients identified were already on optimal treatment to slow progressive kidney failure. Regardless, those with anaemia and advanced CKD would probably benefit from referral to a nephrologist. Other metabolic complications of CKD were not examined due to the lack of information as measurement of these parameters (e.g. serum phosphate) were not part of usual clinical practice in these clinics.

Conclusions

We have documented a high prevalence of CKD in patients attending a tertiary hospital geriatrics outpatient clinic. The CKD was associated with an increased risk of mortality and anaemia. The referral patterns of the supervising practitioners demonstrated restricted consultation with nephrologists, despite the potential benefits to the patient. This requires further investigation to examine the reasons behind these referral patterns in order to be able to implement interventions to increase referral if it is deemed appropriate.

Competing interest

The authors declare that they have no competing interests.

Authors' contributions

NB was involved in the design of this study, analysis of the dataset and drafted this manuscript. KM assisted with the design of this study, collected the data from medical records and administrative databases and reviewed the manuscript. CI assisted with the design of this study and reviewed the manuscript. All authors read and approved the final manuscript.

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References

1. Feest TG, Rajamahesh J, Byrne C, Ahmad A, Ansell D, Burden R, Roderick PJ: Trends in adult renal replacement therapy in the UK: 1982–2002. *QJM* 2005, **98**:21–28.
2. White SL, Polkinghorne KR, Atkins RC, Chadban SJ: Comparison of the prevalence and mortality risk of CKD in Australia using the CKD Epidemiology Collaboration (CKD-EPI) and Modification of Diet in Renal Disease (MDRD) Study GFR estimating equations: the AusDiab (Australian Diabetes, Obesity and Lifestyle) Study. *Am J Kidney Dis* 2010, **55**:660–670.
3. Whaley-Connell AT, Sowers JR, Stevens LA, McFarlane SJ, Shlipak MG, Norris KC, Chen SC, Qiu Y, Wang C, Li S, et al: CKD in the United States: Kidney Early Evaluation Program (KEEP) and National Health and Nutrition Examination Survey (NHANES) 1999–2004. *Am J Kidney Dis* 2008, **51**:S13–S20.
4. Olsen J, Bonnevie B, Palmhøj-Nielsen C, Povlsen JV: Economic consequences of an increased number of patients on outgoing dialysis. *Scand J Urol Nephrol* 2010, **44**(6):452–458.
5. Go AS, Chertow GM, Fan D, McCulloch CE, Hsu CY: Chronic kidney disease and the risks of death, cardiovascular events, and hospitalization. *N Engl J Med* 2004, **351**:1296–1305.
6. McCullough PA, Li S, Jurkovic CT, Stevens L, Collins AJ, Chen SC, Norris KC, McFarlane S, Johnson B, Shlipak MG, et al: Chronic kidney disease, prevalence of premature cardiovascular disease, and relationship to short-term mortality. *Am Heart J* 2008, **156**:277–283.
7. Levin A, Djurdjev O, Duncan J, Rosenbaum D, Werb R: Haemoglobin at time of referral prior to dialysis predicts survival: an association of haemoglobin with long-term outcomes. *Nephrol Dial Transplant* 2006, **21**:370–377.
8. Cumming RG, Mitchell P, Craig JC, Knight JF: Renal impairment and anaemia in a population-based study of older people. *Intern Med J* 2004, **34**:20–23.
9. Soni RK, Weisbord SD, Unruh ML: Health-related quality of life outcomes in chronic kidney disease. *Curr Opin Nephrol Hypertens* 2010, **19**:153–159. 110.1097/MNH.1090b1013e328335f328939.
10. Hansberry MR, Whittier WL, Krause MW: The elderly patient with chronic kidney disease. *Adv Chronic Kidney Dis* 2005, **12**:71–77.
11. Campbell K, Dale W, Stankus N, Sachs G: Older Adults and Chronic Kidney Disease Decision Making by Primary Care Physicians: A Scholarly Review and Research Agenda. *J Gen Intern Med* 2008, **23**:329–336.
12. Levey AS, Greene T, Kusek JW, Beck GJ, Group MS: A simplified equation to predict glomerular filtration rate from serum creatinine. *J Am Soc Nephrol* 2000, **11**:155A.
13. Trivedi H: Cost implications of caring for chronic kidney disease: are interventions cost-effective? *Adv Chronic Kidney Dis* 2010, **17**:265–270.
14. Ruilope LM: Angiotensin receptor blockers: RAAS blockade and renoprotection. *Curr Med Res Opin* 2008, **24**:1285–1293.
15. Ogata H, Koiwa F, Kinugasa E, Akizawa T: CKD-MBD: impact on management of kidney disease. *Clin Exp Nephrol* 2007, **11**:261–268.
16. KDIGO: inical practice guideline for the diagnosis, evaluation, prevention, and treatment of Chronic Kidney Disease-Mineral and Bone Disorder (CKD-MBD). *Kidney Int Suppl* 2009, **S1**:130.
17. Levin A, Bakris GL, Molitch M, Smulders M, Tian J, Williams LA, Andress DL: Prevalence of abnormal serum vitamin D, PTH, calcium, and phosphorus in patients with chronic kidney disease: results of the study to evaluate early kidney disease. *Kidney Int* 2007, **71**:31–38.
18. Hirakata H, Tsubakihara Y, Gejyo F, Nishi S, Iino Y, Watanabe Y, Suzuki M, Saito A, Akiba T, Inaguma D, et al: Maintaining high hemoglobin levels improved the left ventricular mass index and quality of life scores in pre-dialysis Japanese chronic kidney disease patients. *Clin Exp Nephrol* 2010, **14**:28–35.
19. Agnihotri P, Telfer M, Butt Z, Jella A, Cella D, Kozma CM, Ahuja M, Riaz S, Akamah J: Chronic anemia and fatigue in elderly patients: results of a randomized, double-blind, placebo-controlled, crossover exploratory study with epoetin alfa. *J Am Geriatr Soc* 2007, **55**:1557–1565.
20. Akbari A, Swedko PJ, Clark HD, Hogg W, Lemelin J, Magner P, Moore L, Ooi D: Detection of chronic kidney disease with laboratory reporting of estimated glomerular filtration rate and an educational program. *Arch Intern Med* 2004, **164**:1788–1792.
21. Garg AX, Mamdani M, Juurlink DN, van Walraven C: Identifying individuals with a reduced GFR using ambulatory laboratory database surveillance. *J Am Soc Nephrol* 2005, **16**:1433–1439.
22. Garg AX, Papaioannou A, Ferko N, Campbell G, Clarke JA, Ray JG: Estimating the prevalence of renal insufficiency in seniors requiring long-term care. *Kidney Int* 2004, **65**:649–653.
23. Philipneri MD, Rocca Rey LA, Schnitzler MA, Abbott KC, Brennan DC, Takemoto SK, Buchanan PM, Burroughs TE, Willoughby LM, Lentine KL: Delivery patterns of recommended chronic kidney disease care in clinical practice: administrative claims-based analysis and systematic literature review. *Clin Exp Nephrol* 2008, **12**:41–52.
24. Joly D, Anglicheau D, Alberti C, Nguyen AT, Touam M, Grunfeld JP, Jungers P: Octogenarians reaching end-stage renal disease: cohort study of decision-making and clinical outcomes. *J Am Soc Nephrol* 2003, **14**:1012–1021.
25. McDonald SP, Russ GR, Kerr PG, Collins JF: ESRD in Australia and New Zealand at the end of the millennium: a report from the ANZDATA registry. *Am J Kidney Dis* 2002, **40**:1122–1131.
26. Kazmi WH, Obrador GT, Khan SS, Pereira BJ, Kausz AT: Late nephrology referral and mortality among patients with end-stage renal disease: a propensity score analysis. *Nephrol Dial Transplant* 2004, **19**:1808–1814.
27. Thomas MC: The CARI guidelines. Prevention of progression of kidney disease: early referral of patients with pre-end-stage kidney disease. *Nephrology (Carlton)* 2007, **12**(Suppl 1):S41–S43.
28. Navaneethan SD, Kandula P, Jeevanantham V, Nally JV Jr, Liebman SE: Referral patterns of primary care physicians for chronic kidney disease in general population and geriatric patients. *Clin Nephrol* 2010, **73**:260–267.
29. Chronic kidney disease (CKD) management in general practice. In *Book Chronic kidney disease (CKD) management in general practice*. 2nd edition. Melbourne: Kidney Health Australia; 2012.
30. *The Australian Nephrology Workforce Survey*. 2007 <http://www.nephrology.edu.au/workforce/documents/AustralianNephrologyWorkforceReportforANZSN.pdf>.
31. *Chapter 1 - Stock and Flow & Appendix II*. <http://www.anzdata.org.au/anzdata/AnzdataReport/32ndReport/Ch01.pdf>.

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