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BMJ Open Self-management interventions for adults with chronic kidney disease: a scoping review

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ABSTRACT

Objective To systematically identify and describe selfmanagement interventions for adult patients with chronic kidney disease (CKD).

Setting Community-based.

Participants Adults with CKD stages 1–5 (not requiring kidney replacement therapy).

Interventions Self-management strategies for adults with CKD.

Primary and secondary outcome measures Using a scoping review, electronic databases and grey literature were searched in October 2016 to identify selfmanagement interventions for adults with CKD stages 1-5 (not requiring kidney replacement therapy). Randomised controlled trials (RCTs), non-RCTs, qualitative and mixed method studies were included and study selection and data extraction were independently performed by two reviewers. Outcomes included behaviours, cognitions, physiological measures, symptoms, health status and healthcare.

Results Fifty studies (19 RCTs, 7 quasi-experimental, 5 observational, 13 pre-post intervention, 1 mixed method and 5 qualitative) reporting 45 interventions were included. The most common intervention topic was diet/nutrition and interventions were regularly delivered face to face. Interventions were administered by a variety of providers, with nursing professionals the most common health professional group. Cognitions (ie, changes in general CKD knowledge, perceived self-management and motivation) were the most frequently reported outcome domain that showed improvement. Less than 1% of the interventions were co-developed with patients and 20% were based on a theory or framework.

Conclusions There was a wide range of selfmanagement interventions with considerable variability in outcomes for adults with CKD. Major gaps in the literature include lack of patient engagement in the design of the interventions, with the majority of interventions not applying a behavioural change theory to inform their development. This work highlights the need to involve patients to co-developed and evaluate a self-management intervention based on sound theories and clinical evidence.

Strengths and limitations of this study

- A strength of our study is that it is the first scoping review to apply the principles of patient-oriented research, where patient partners were engaged in determining the research question, advising us on search terms and reviewing the results to ensure we captured and reported the data meaningfully.
- Our scoping review is comprehensive in nature, with inclusion of all study designs and consideration of self-management features that have not been investigated previously.
- ▶ Due to the heterogeneous nature of the literature, it was challenging to synthesise the data. To address this challenge the two reviewers used two standardised tools to independently extract data and independently coded the outcomes into categories using the revised Self- and Family Management Framework.
- A limitation of our scoping review is that we were unable to assess the self-management outcomes in terms of sustained changes in behaviour, physiological and health status.
- We were unable to draw conclusions regarding the most effective self-management intervention for adult patients with chronic kidney disease, keeping in mind that our aim was to review the breadth of the current literature and present the gaps that exist.

INTRODUCTION

Chronic kidney disease (CKD) is associated with adverse health outcomes, poor quality of life and high healthcare costs. Patients with CKD often experience a number of comorbidities including diabetes, cardiovascular disease and depression.² They must balance the medical management of their kidney disease and other chronic conditions with demands of their daily lives, including managing the emotional and psychosocial consequences of living with chronic disease. In a recent CKD research priority setting



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study, individuals with non-dialysis CKD, their caregivers, clinicians and policy-makers identified the need to develop optimal strategies to enable patients to manage their CKD and related comorbidities to slow or prevent the progression to end-stage kidney disease (ESKD).³ International data in research priority setting for kidney disease also highlights self-management as a top priority to prevent progression.⁴

Self-management interventions aim to facilitate an individual's ability to make lifestyle changes and manage symptoms, treatment and the physical and psychosocial consequences inherent in living with CKD and associated comorbidities.⁵ Self-management of CKD involves focusing on illness needs (developing knowledge, skills and confidence to manage medical aspects), activating resources (identifying and accessing resources and supports) and living with the condition (learning to cope with the condition and its impact on their lives as well as the emotional consequences of the illness). Self-management requires patient engagement; however, the degree to which patients are able or willing to participate in self-management can vary, and individual and health system factors may serve as facilitators or barriers to self-management processes.⁷

Despite the high prevalence of CKD and its impact on patient outcomes, there is limited evidence on the effectiveness of self-management interventions. Prior systematic reviews^{8–11} and three integrative reviews^{12–14} found that self-management interventions were variable in their effectiveness for managing and preventing progression of CKD. While these reviews add to the knowledge base, they have restricted inclusion criteria (eg, study type, patient population) and unclear reporting strategies (ie, describing complex self-management interventions in detail and providing structured accounts of the interventions and outcomes). In particular, features of self-management interventions such as person centeredness, applicability to comorbidities associated with CKD, physiological and non-physiological outcomes and application of any behavioural change theories are often lacking. Self-management interventions need to be tailored to suit diverse patient needs and preferences as well as the local healthcare context.⁷ Therefore, investigating the 'who', 'what' and the 'how' of self-management interventions is crucial. We used recognised literature synthesis and reporting guidelines, along with engagement of our patient partners in determining the research question and search terms as well as reviewing the results to ensure we captured and reported the data meaningfully.

To our knowledge, there is no literature synthesis that systematically and comprehensively summarises the breadth of evidence found in primary quantitative, qualitative and mixed methods research regarding self-management interventions for adult patients with CKD. We used a scoping review methodology to understand the range and types of interventions including both educational and support interventions for CKD to inform the future design of a self-management intervention. Specifically,

we conducted a scoping review to identify and describe self-management interventions for adult patients with CKD (stages 1–5; non-dialysis, non-transplant).

MATERIALS AND METHODS

We used a scoping review methodology to enable us to incorporate a broad range of studies and to summarise the knowledge from a variety of sources and types of evidence. 15 Our aim was to identify gaps in literature related to CKD self-management interventions and inform future research. A unique and important aspect was the involvement of 'patient partners'. Through a national initiative, Canadians Seeking Solutions and Innovations to Overcome CKD (Can-SOLVE CKD), patients work side by side with researchers, clinicians and decision makers to address patient-oriented research priorities. 16 Our research team includes Can-SOLVE CKD patient partners with CKD and caregivers. 16 Using the Joanna Briggs Institute framework for scoping reviews, we undertook the following steps: (1) identified the research question, (2) identified relevant studies, (3) completed study selection, (4) charted, collated, summarised and reported the results (5) and consulted with our patient partners. 15 17 These steps were iterative to ensure comprehensive inclusion of the literature and continued meaningful engagement with our patient partners. This work involves identifying, reviewing and categorising data from primary articles and does not involve human participants and is exempt from ethics approval.

Research aim

Our scoping review aimed to determine the available self-management interventions for adults aged 18 years and over and diagnosed with CKD stages 1–5 (not requiring dialysis or transplant).

Search and selection of studies

We worked with an information specialist (DL) to identify key words that represented the population (CKD) and the intervention (self-management). We searched a broad range of information sources including the following online databases: MEDLINE (OVID), EMBASE, PsycINFO, Cochrane Central Register of Controlled Trials, CINAHL Plus and Cochrane Database of Systematic Reviews for published studies, with no limits on date (inception to October 2016), language, age or study design. We also searched Web of Science from 2006 to October 2016 to capture recently published meeting abstracts and summaries. Using the Canadian Agency for Drugs and Technology (CADTH) Grey Matters approach, ¹⁸ we searched Google Canada, Health Technology Assessment (HTA) agencies (Canada, Australia, Ireland, UK and USA) and Clinical Trials databases (Biomed Central—ISRCTN Registry, US National Institutes of Health, ClinicalTrials. gov) during October 2016 with no language restrictions (online supplementary table 1). Our search strategy for grey literature was guided by the specific database (ie, Google search operators, website search filters) and was completed within a single session for each search strategy to ensure consistency due to the dynamic nature of the internet (online supplementary table 2). Two reviewers (BK and MD) also reviewed the reference lists of included studies, along with those identified in past systematic and integrative reviews of our research topic. We contacted authors of relevant protocols and conference abstracts to ascertain if their work and findings were published.

A study was included if the population involved adults with CKD (stages 1–5, non-dialysis, non-transplant). Self-management interventions included strategies, tools or resources in any delivery format (print, electronic, face to face and so on) that facilitated an individual's ability to make lifestyle changes or to manage symptoms, treatment or the physical and psychosocial consequences inherent in living with CKD and other associated comorbidities. Interventions targeted only at selection of treatment for ESKD (ie, dialysis, kidney transplant) were excluded. Other self-management interventions or standard care were considered as a comparison. We included primary studies that used quantitative, qualitative or mixed methods. Systematic and integrative reviews were identified for the purpose of reviewing their included studies for potential relevant studies. We excluded case series, case studies, case reports, clinical practice guidelines, theses and opinion-driven reports (editorials, non-systematic or literature/narrative reviews).

Three reviewers (BK, MD and BH) performed an initial screen of titles and abstracts using a citation screening tool. To determine inter-rater reliability, a calibration exercise was performed by the three reviewers. Pilot testing a random sample of 50 citations achieved good agreement (kappa=0.79) at which point the three reviewers screened the remaining titles and abstracts. Two reviewers (BK and MD) followed a similar procedure for identifying relevant full text studies, with good agreement between the two reviewers (kappa=0.78). Disagreements were resolved by discussion and obtaining consensus between the three reviewers.

Charting, collating and summarising the data

We developed a data extraction form based on the Template for Intervention Description and Replication (TIDieR) checklist. 19 This checklist provides a template to structure accounts of an intervention (eg, goal of intervention, materials used, who delivered the intervention and how, where, when and how much and how well the intervention was delivered). We also used the Effective Practice and Organisation of Care (EPOC) data collection form²⁰ to ensure we were comprehensive in extracting relevant study characteristics as outlined by Cochrane EPOC group. Study characteristics (eg, study design, country of origin, publication year), population characteristics (eg, CKD stage, comorbidities) and self-management intervention characteristics (eg, topics, format, target audience, providers, location, dose, duration and so on) were documented. For the study

outcomes, the two reviewers (BK and MD) independently coded each outcome into categories identified by Grey et al (eg, behaviours, cognitions, physiological measures, symptoms, health status, healthcare and other). We pilot tested the form on a random sample of 10 eligible studies and once consensus between the two reviewers was reached, we independently abstracted data from the remaining eligible studies. Data were categorised and reported descriptively (ie, counts and frequencies). For qualitative studies, we identified the methodology and key concepts presented by the authors.

Consultation with patient partners

Patient partners were engaged throughout this work, specifically to provide input on the research question, search strategies (eg, grey literature sources) and reviewing the final results. The results were presented and discussed at the national Can-SOLVE CKD meeting.

RESULTS

Search results

From 12583 unique citations (figure 1), we included 50 full text studies. ^{21–70}

Description of studies

A summary of the 50 studies included in this review is provided in table 1.

The most common study designs were randomised controlled trials (RCTs) (38%). Non-RCTs consisted of quasi-experimental (14%), observational (10%), pre-post intervention (26%), qualitative (10%) and mixed methods (2%). The studies were conducted in 14 countries, including the USA (20%), UK (14%) and Australia (12%). Most studies were published in the last 5 years (64%).

Patient population characteristics

The target population in most studies was CKD (72%) and 15 studies mentioned CKD plus one or more associated comorbidities. The average ages of participants reported across studies were 50.2 to 74.3 years.

Description of self-management interventions

Table 2 summarises the characteristics of the self-management interventions. Five studies reported the same self-management intervention; therefore, 45 interventions were summarised. The most common intervention topic was diet/nutrition (64%) and the least common topics were symptom management and lifestyle (13% and 11%, respectively). The most frequent modes of delivering the intervention were face to face (80%), multiple (ie, more than one mode) (71%) and print (64%). Electronic was the least common delivery mode (16%). Interventions were administered by a variety of providers. The most common category of providers was 'other' (56%), which was made up of various types of health professionals and lay people. However, the most common identifiable group of providers were nursing

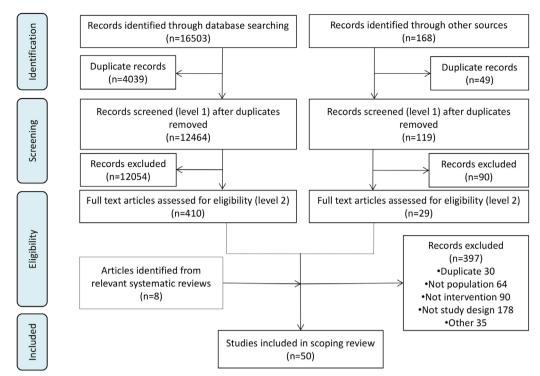


Figure 1 Prisma flow diagram.

professionals (49%). Patient volunteer/mentor was the least common (9%). The outpatient setting was the most common location for providing the self-management intervention (51%), and the inpatient setting was the least popular (2%). Many studies did not report the intervention language (53%), but 12 languages were represented and seven studies reported that they provided the intervention in multiple languages.

In terms of intervention development, only 20% of studies mentioned the use of evidence such as theories or frameworks. These included the transtheoretical model of behaviour change, social cognitive theory and chronic care model. Less than 1% of the studies involved patients in the design of the intervention, where patients were interviewed regarding intervention content. $^{26\,31-33}$

Description of quantitative study outcomes and results

Characteristics of the quantitative study outcomes are presented in table 3. Twenty-three (46%) studies measured physiological outcomes (ie, laboratory tests, body composition and so on). The least common outcomes reported by studies were health status and healthcare (each 10%) and symptoms (ie, fatigue) (4%). Table 4 summarises the details of the quantitative studies. We categorised the overall study results descriptively as improved, unchanged or worse. Many studies had more than one outcome measure (eg, one measure improved, another had no change) and they were reported as mixed results. Based on this method of categorization, 89 outcomes were reported, of which 61% improved, 20% had no change, 1% worsened and 13% had mixed results. Four of the results were reported as not applicable as the outcomes were not relevant. Of the 54 outcome categories

that improved, 15 were cognition, 9 were physiological measures, 8 were behaviours, 8 were individual outcomes, 5 were health status, 4 were healthcare, 4 were intervention specific and 1 was symptom management.

Description of qualitative study outcomes and results

Table 5 summaries the findings from six qualitative studies that explored patient perspectives, one of these being a mixed methods study. All studies used semistructured interviews and one also used a questionnaire. The aims of all these studies were to examine patient perspectives' regarding the self-management interventions they were involved in. Due to the variety of interventions (eg, intervention topics, delivery mode and providers of the intervention), it was difficult to summarise findings into meaningful categories. Overall, patients highlighted that interventions needed to be individualised and tailored to their specific situations and preferences (eg, awareness of having CKD, stage of CKD, knowledge of the disease, access to resources and so on).

DISCUSSION

To our knowledge, this is the first scoping review involving patients as research partners to identify and summarise self-management interventions for adults with CKD. The scoping review methodology enabled us to systematically summarise a broad range of self-management interventions and describe their features. We identified 50 studies that investigated self-management interventions for adults with CKD, with considerable variation in interventions, outcomes assessed and results obtained (ie, some improved and/or some worsened and/or some did not

Table 1 Characteristics of the studies included in scoping review

Characteristic	Studies (n=50)
Study design	
Randomised controlled trial	19
Pre-post test	13
Quasi-experimental (controlled/non-random)	7
Observational	5
Qualitative	5
Mixed methods	1
Origin of study	
USA	10
UK	7
Australia	6
Canada	5
Taiwan	5
Netherlands	3
Spain	3
Italy	2
Japan	2
New Zealand	2
Sweden	2
Brazil	1
Denmark	1
Korea	1
Year of publication	
2012–2016	32
2007–2011	11
Prior	7

change). We found that self-management interventions for CKD is an emerging area with most studies published within the last 5 years which may be related to the growing recognition of the importance of incorporating patients and their families in managing their disease to improve outcomes.⁷

Our findings are similar to prior reviews reporting that the design of self-management interventions for CKD has not been theoretically driven and they have been predominately designed by healthcare professionals without input from patients. ¹³ ¹⁴ Person-centred care is changing how healthcare professionals deliver care to patients, but more importantly how patients and their families are actively involved in self-managing their chronic conditions. ⁷¹ Engaging patients by having them co-design self-management interventions will ensure that patient preferences based on their values, culture and psychosocial needs will be addressed in the self-management intervention. ^{12–14} Through our current national partnership with patients, researchers and clinicians, we have the opportunity to

Table 2 Overall characteristics of self-management interventions

Variable	Intervention count (n=45)
Intervention topics	
Diet/nutrition	29
General CKD knowledge	18
Other (ie, advanced care planning, meditation)	18
Medication	17
Modalities	13
Physical activity	13
Comorbidities	11
Symptom management	6
Lifestyle	5
Mode of delivery	
Face to face (ie, group, one-on-one)	36
Multiple modes	32
Print	29
Distance (ie, telephone, email)	13
Digital (ie, DVD, PowerPoint, audio recording)	8
Electronic (ie, website, mobile application)	7
Type of providers	
Other*	25
Nurse/nurse practitioner	22
Dietitian	14
Multiple providers	13
Social worker	6
Physician/primary care physician	6
Nephrologist/nephrology fellows	5
Patient volunteer/mentor	4
Pharmacist	1
Location of intervention	
Outpatient	23
Not specified	12
Community (non-clinic)†	10
Patient home	10
Multiple locations	7
Inpatient	1
Intervention languages	
Not Specified	24
English	10
Multiple languages	7
Mandarin	4
Spanish	3
Taiwanese	3

Continued

Table 2 Continued

Tubic 2 Continued	
Variable	Intervention count (n=45)
Dutch	2
Cantonese	1
French	1
Greek	1
Italian	1
Japanese	1
Swedish	1
Vietnamese	1
Intervention development	
Use of framework or theory	9
Codesigned with patients	4

^{*}Other providers: Trained research assistant, lay health worker, Bengali worker, Educators (health, cook, diabetic), online tool, physician assistant, exercise physiologist, technician, psychologist, employment expert, instructor, interpreter, physiotherapist, patient, principal investigator.

obtain patient perspectives, along with incorporating a behaviour change theory to inform the future design of a self-management intervention for CKD.

Only 28% of studies that we identified included patients with CKD plus other comorbidities, despite the common presence of comorbidities in this patient population. Less than one-quarter of included studies provided information on how to manage comorbid conditions such as tracking lab results and symptom management. This highlights the need to consider 'whole person care',

where the self-management intervention needs to encompass the physical, mental and emotional needs of the patient⁷² ⁷³ that are important to them as well as meeting the individuals desires by collaboration between relevant providers.⁷¹

Forty-five different self-management interventions were identified, with one or more topics presented in a variety of formats and by a variety of providers. Symptom management and lifestyle topics were not included in many of the interventions. Based on prior work, non-dialysis patients with CKD have indicated that these were important topics for them in managing their CKD with an aim to slow the progression of CKD and will be important to consider in the development of future interventions. Face to face was the most common delivery format while electronic (internet or mobile application) was least common, with many studies reporting multiple formats (ie, face to face and printed materials). With the expansion of electronic platforms for supporting patients and providers in the uptake of evidence-based care, there is the potential to use an electronic format to support patients in self-managing their CKD and other comorbidities.⁷⁴ It is worth noting that there was variability in duration and frequency of face to face encounters, from a single session to multiple sessions over weeks to months. While varied options for in-person delivery is good if it meets the needs of the patients and their families, it may not be feasible on a larger scale due to the resources required. Only five studies looked at self-management healthcare cost-effectiveness, healthcare utilisation and access, each measuring different end-points with mixed results. Future self-management interventions should include the essential principles to self-management (eg, accessing relevant health information, adhering to multiple treatment

Table 3 Summary of quantitative study outcomes*

Common outcomes	Description	Number of studies	Number of studies in which outcome improved
Physiological measures	Changes in laboratory tests, blood pressure, body composition, functional/performance tests and cardiovascular risk	23	9
Cognitions	Changes in general CKD knowledge, self-efficacy, self-management, motivation, perceived stress, anxiety and fear	21	15
Behaviours	Adherence to diet, medication, physical activity, sleep, blood pressure control	13	8
Individual outcomes	QOL, well-being and general satisfaction	11	8
Intervention specific	Reporting of general concepts regarding feasibility of intervention, enjoyment and interest in intervention	9	4
Healthcare	Measurements of cost effectiveness, healthcare utilisation and access	5	4
Health status	Measurements of morbidity and mortality (ie, time to dialysis, survival, all-cause mortality)	5	5
Symptoms	Changes in overall symptoms (ie, pain, fatigue)	2	1

^{*}Based on primary and distal outcomes from Grey et al.⁶ CKD, chronic kidney disease; QOL, quality of life.

[†]Community: gym, grocery store, "study room". CKD, chronic kidney disease.

	Study results							
	Study outcomes		Health status: Duration between session and dialysis initiation—patient in E group survived 4.6 months longer w/o requiring RRT	Oognitions: Patient reliance on dietician's feedback, support and modelling strategies—decreased over time in E group Individual outcomes: Top rated interventions by patients—counselling, self-monitioning, protein counter	Health status: ▼ Time to dialysis—E group had 3 month delay in dialysis compared with C group	Health status: Survival predialysis and after dialysis initiation—significantly longer in the E group (2.25 years and 8 months, respectively)	Individual outcomes: QOL—many components of KDQOLSF V1.3 improved: CKD symptoms, cognitive function, vitality Physiological measures: ▼ Nutritional assessment PG-SGA—in E group those who were malnourished at baseline improved, in C group mainourished from 12.5%—25%	Intervention specific: Feasibility (recruitment, retention, patient satisfaction, patient access of additional support)—findings suggest delivening/evaluating an effective structured group educational intervention to promote better BP control would be challenging
	Description of intervention		Finhanced education: 22-page booklet Individual slide presentation (75 min) Duration – one session Comparator: standard care	"Wodification of diet in renal disease": * Keeping Track' booklet * Monthly meeting with dietician * Protein Wise Counter (lists protein content of foods) * Shopping Wise': a guide to convenience and fast foods * Visited restaurants and shops * Visited restaurants and shops * Duration – 26 months * Comparator: standard protein diet	Psychoeducation': ▶ 60-page booklet ▶ 90min interactive educational intervention personalised for each patient ▶ Supportive (10min max) phone calls Q3 weeks Duration — 18 months or initiation of RRT Comparator: standard care	Psychoeducation session': ► 22-page booklet ► Individual slide presentation (60–75 min long) Duration — one visit Comparator: standard care	'Individual nutritional counselling': ► Initial individual consultation with dietician ► Then phone follow-up Q2 weeks x 1 month then Q1 month Duration: 12 weeks Comparator: standard care	Structured education session': ► Leaflet on HTN management ► CHEERS patient education intervention and standard care ► 2.5-hour group session ► Phone supporf from nurse Duration: one session Comparator: standard care
	Delivery format		► Print ► Face to face ► PowerPoint slides	▼ Print Face to face	▶ Print ▶ Face to face ▶ Telephone	► Print Face to face FowerPoint slides	► Face to face Telephone	► Print ► Face to face Telephone
	Provider(s)		Trained research assistant	Dietician	Social worker	Health educator	Dietician	Nurse
	Intervention topic(s)		 ▶ General CKD knowledge ▶ Diet/nutrition ▶ Modalities 	■ DieVnutrition	► General CKD knowledge ► Diet/nutrition ► Medication ► Modalities ► Lifestyles	► General CKD Knowledge ► Diet/nutrition ► Modalities	► Diet/nutrition ► Other (ie, self-management principles)	▼ Comorbidities (le, HTN management)
studies	Study size Age (years)		204 (E=87, C=92, not part of education=25) Age: 50.2	840 (unclear) Age: NR	297 (E=149, C=148) Age: 58.6	335 (E=172, C=163) Age: 47.4–53.9	47 (F=24, C=23) Age: 68.5–72.6	81 (E=40, C=41) Age: 62.8–65.4
Summary of guantitative studies	Target population		Pre-RRT CKD (creatinine>350 µmol/L and rising rapidly)	OKD 35	CKD (creatinine<300 µmo/L and deemed to need RRT in 6-18 months)	OKD with progressive reduction in kidney function	OKD 4-5	CKD 1–4+HTN
Sum	Design		RCT	RCT	RCT	RCT	RCT	RCT
Table 4	Study and year (Reference)	RCT	Binik <i>et al</i> (1993) ³⁴	Gillis et al (1995) ³⁵	Devins <i>et al</i> (2003) ³⁶	Devins <i>et al</i> (2005) ³⁷	Campbell et al (2008) ³⁸	Byrne <i>et al</i> (2011) ²⁶

	Study results									
	Study outcomes	Physiological measures: • GGFR change—higher in E group • GGFR reduction of >50% less in E group in E group cause mortality—no significant difference between groups	Health status: ► # of hospitalisations in 1 year of follow-up—less in E group	Physiological measures: ► Improvement in 4/5 of the following: urinary protein, total cholesterol, eGFR decline, BP, urinary sodium—was considered a success—61% in E group vs 12% in C group	Behaviours: ► SM score—some changes in some components in both groups	Cognitions: ▶ Self-efficacy	Behaviours: Medication adherence Adherence to BP and weight measurements Limiting salt intake Alcohol consumption Smoking	Physiological measures: ▶ BP	Physiological measures: ▶ BP—no difference between groups	Behaviours: ► Medication adherence—no difference between groups
	Description of intervention	'Self-management Support': Individual monthly health education Weekly telephone based support Aid of support group twice monthly (5-10 patients) Duration: 12 months Comparator: standard care		Cooking and exercise class: Standard care and: Group CKD nutrition class (with dietician and cook educator: 2 hour sessions over 4 weeks) plus one shopping tour led by a dietician CKD cookbook 12-week exercise programme (3 × 1 hour	sessions/week, led by a certified exercise physiologist and nurse <i>Duration:</i> 12 weeks <i>Comparator:</i> standard care	'EASE (encourage autonomous self-enrichment) programme: Nurses Isten to what patients have difficulties	and discuss how they will try to improve P ace to face interview monthly Telephone or email contact every 2 weeks Duration: 12 weeks Comparator: standard care		'Multifactorial intervention': ► Individual medication review (draw chart) ► Daily self-monitoring of BP × 3 months	Q2 week motivational interviewing follow- up via phone x 12 weeks to support BP management and optimise medication SM Duration: 12 weeks Comparator standard care
	Delivery format	▶ Print ▶ Face to face ▶ Telephone		▼ Print Face to face		► Face to face ► Telephone ► Fmail			Print Face to face Telephone	
	Provider(s)	Nurse, dietician, nephrologist, peers, volunteers		Nurse, exercise physiologist, dietician, cook educator		Nurse			Nurse	
	Intervention topic(s)	■ General CKD knowledge ■ Diet/nutrition ■ Medication ■ Lifestyle ■ Modality information for stage IV		▶ Diet/nutrition ▶ Physical activity		▼ Other			▼ Medication ▼ Comorbidities	
	Study size Age (years)	54 (E=27, C=27) Age: 68.2		40 (E=23, C=17) Age: 63.4		31 (E=19, C=12)	Age: 69.8		75 (E=39, C=41)	76e: 0.7
pen	Target population	CKD 3-5		OKD 3-4+HTN		CKD			CKD 2–4 (diabetic kidney disease)+DM+HTN	
Continued	Design	RCT		RCT		RCT			RCT	
Table 4	Study and year (Reference)	Chen <i>et al</i> (2011) ³⁸		Flesher <i>et al</i> (2011) ⁴⁰		Joboshi <i>et al</i> (2012) ⁴¹			Williams et al (2012) ⁴²	

	Study results		•		<u>a</u>			b wer		Φ	00),
	Study outcomes	Intervention specific: ► Attrition rate to assess feasibility of study—high attrition	Cognitions: ► Medication self-efficacy—no difference between groups	Healthcare: ► Health care utilization—no difference between groups	Physiological measures: ▶ Routine clinical lab surrogate measures—no difference	Behaviours: ► Medication adherence—no difference	Individual outcomes: ▶ General well-being—no difference	Physiological measures: BP-decreased by 8mm Hg in E group 24 hours urinary salt excretion—decrease in E group eGFR—no difference between groups	Behaviours: P Reduction in protein intake— decreased Adherence to low protein diet—effective	Physiological measures: ■ Body composition: waist circumference, body fat, BMI, mid-am muscle circumference—no change ■ Serum albumin—no change	Cognitions: ▶ Positive and active engagement in life (heiQ)—no difference between groups	Physiological measures: ▶ BP control—better BP maintenance in Egroup	Individual outcomes: Health related QOL (EuroQoL EQ-5D index)—higher in E
	Description of intervention		Individual slide presentation (¿Umin) via interpreter (Greek, Italian, Vietnamese) ▼ Q2 week motivational interviewing follow-up via phone × 12 weeks	Duration: 12 weeks Comparator: standard care				Uplet advice: ▶ Practical cooking and education sessions in the community facilitated by a Bengali worker the community facilitated by a Bengali worker. ▶ Followed by Q2 week phone calls to reinforce advice and set new targets. Duration: 6 months. Comparator: standard care	Nutrition education programme: Standard dietary counselling AND Education folder with recipes to replace saft with sodium free seasoning blends Individual 15–20 min class	Hands on session about protein rich food Hands on session using test tubes with the amount of salt in different foods Telephone call to address any doubts with dietary plan Duration: 4-7 months Comparator: standard care	'Information and telephone-guided access to community services: Widney Information Guidebook Patient-Led Assessment for Network Support	tailored access to community resources Telephone guided help from a lay health worker	Duration: 6 months Comparator: standard care
	Delivery format	Print 'Se Face to face Pleiphone PowerPoint sildes						► Face to face Telephone	► Print ► Face to face ► Telephone		▼ Print ▼ Website ▼ Telephone		
	Provider(s)	Nurse, interpreter						Dietician and Bengali worker	Dietician		Lay health worker		
	Intervention topic(s)	■ Medication ■ Comorbidities ■ Other (le, self-efficacy)					▶ Diet/nutrition	▶ Diet/nutrition		▼ General CKD knowledge Comorbidities ▼ Comorbidities ▼ Other (ie, community resources			
	Study size Age (years)	78 (E=40, C-38) Age: 74.31						56 (E=28, C=28) Age: 55.7–60.7	89 (E=43, C=46) Age: 63.4		436 (E=215, C=221) Age: 72.1		
pel	Target population	CKD 2-4+DM+ cardiovascular disease				CKD 3-5+HTN (BP>130/80) +Bengali population	CKD 3-5		CKD 3				
Continued	Design	RCT						RCT	RCT		RCT		
Table 4	Study and year (Reference)	Williams <i>et al</i> (2012) ⁴³						de Brito- Ashurst et al (2013) ⁴⁴	Paes-Barreto et al (2013) ⁴⁵		Blakeman <i>et al</i> (2014) ⁴⁶		

Table 4 (Continued	pe							
Study and year (Reference)	Design	Target population	Study size Age (years)	Intervention topic(s)	Provider(s)	Delivery format	Description of intervention	Study outcomes	Study
McManus et al (2014) ⁴⁷	RCT	HTN (BP>130/80) +CKD3 or DM or CHD	555 (E=277, C=278) Age: 69.3–69.6	■ Medication Comorbidities	General practitioner, patient	▶ Print Face to face	Self-monitoring of BP and self-titration of medications: Self-monitoring of BP Self-tration of medications following a 3-step plan designed by general practitioner and patient Duration: 12 months Comparator standard care	Physiological measures: SBP at 12 months − no difference difference Healthcare: Prescription of antihypertensive medications increased in both groups but greater significance in Egroup Sumptom month.	
								Adverse effects— no significant difference between groups Individual outcomes: QOL—no significant difference between groups	
Park e <i>t al</i> (2014) ⁴⁸	RCT	CKD3+HTN+ African-American	15 Age: 58.7	▶ Other (ie, meditation)	Principle investigator, patient	► Face to face Audio recording	'Mindfulness meditation (MM)' 14 min of preecorded guided MM using MP3 player and headphones Duration: one session Comparator: BP education	Physiological measures: P BP-decrease in SBP/ DBP/HR/ MAP Muscle sympathetic nerve activity—decreased	
Howden et al (2015) ¹⁸	RCT	CKD 3-4 and 72 >-1 uncontrolled (E=36, cardiovascular risk factor C=36) Age 66	72 (F-36, C-36) Age 60.2-62.0	▶ Physical activity	Nurse practitioner, social worker, exercise exprisse physiologist, dietician, psychologist, diabetes educator diabetes educator	► Print Face to face	Exercise training and lifestyle intervention: Standard care AND Detailed medical/surgical history taken by nurse practitioner Education about exercising safely: maintaining hydration, signs/symptoms of abnormal response to exercise parecription on hypoglycaemia Exercise prescription individualised on patient's commontal conditions. Goal=150 min/week of moderate intensity exercise plus resistance training exercise plus resistance training booklet Patients contacted regularly to monitor adherence to training booklet Patients contacted regularly to monitor adherence to training Duration: 12 months Comparator: standard care	Physiological measures: ► METS-improved ► 6 min walk distance-improved ► BMI-improved	
Leehey <i>et al</i> (2016) ⁵⁰	RCT	CKD 2-4+DM2+BMI>30+ persistent proteinuria	36 (Exercise+dlet = 18, Diet=18) Age: 66	► Diet/nutrition	Personal trainer	▼ Face to face ▼ Telephone	*Structured exercise programme": Dietary counselling=baseline nutritional counselling with nine follow-up phone calls (both groups) AND Supervised exercise programme 3x week (60 min cardio plus 25-30 min resistance training) Followed by home exercise phase: 3x/week x 60 min with weekly follow-up phone calls and patient encouraged to meet trainer G1 month Duration: 12 months Comparator: diet counselling only	Physiological measures: Vuine protein to creatinine ratio—no change at 52 weeks Symptom limited and constant work rate treadmill time—significant increase in diet+exercisegroup at 12 but not 52 weeks Vuine albumin to creatinine ratio—no change Fratio—no change Inflammation—no change Inflammation—no change Fratio—no change CERF—no change Fratio—no change CERF—no change CERF—no change CERF—no change Fratio—no change	

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Table 4	Continued	per							
Study and year (Reference)	Design	Target population	Study size Age (years)	Intervention topic(s)	Provider(s)	Delivery format	Description of intervention	Study outcomes	Study
Montoya et al (2016) ³⁰	RCT	OX 4	30 (F=16, C=14) Age: 67.9-68.3	General CKD knowledge Diet/nutrition Medication Modalities Other (e, putting affairs in order)	Nephrologist, nurse practitioner, dietician, social worker	► Print Face to face FowerPoint slides	'Nurse practitioner facilitated CKD group visit: Binder with section on individual labs, another section for topics of groups visitis Six 15-2-hour long monthly group visits of 8 patients (-1/2 had family members with them) Three visits done in conjunction with nephrologist's examinations (first half-apt, second half-aducation) Interedive discussion and each visit Side presentation (30-45 min) Duration: 9 months	Cognitions: Cognitions: CKD knowledge—improved in both groups Self-efficacy/disease SM— upward trend in E group Individual outcomes: Satisfaction—high	
Non-RCT							Comparator. standard care		
Robinson <i>et al</i> (1988) ⁵¹	sqO	CKD	25 Age: NR	□ General CKD knowledge □ Diet'nutrition ■ Medication □ Other (le, self-care activities Modalities	Ω.	► Face to face	Henal Bingo: ■ Bingo game format/group gaming technique ■ Provision of refreshments & prizes for motivation ■ Refreshments made with dietician consultation consultation consultation controlled.	Cognitions: Information was gained or reinforced—desirable outcome Met a variety of learning needs—desirable outcome	
							Comparator: none	Intervention specific: Participation was enjoyed—desirable outcome Interest expressed for repeating the exercise—desirable outcome	
Klang e <i>t al</i> (1998) ⁵²	ä	OKD 4-5	56 (E=28, C=28) Age: 54–58	General CKD knowledge Diet/nutrition Physical activity Modalities Chter (le, psychosocial— impact of CKD on economy, family and social life)	Nurse, physician, social worker, dietician, physiotherapist	Face to face	'Pre-dialysis patient education: ► Four 2-hour sessions of group teaching with a classroom approach ► Individual support follow-up by nephrology team member • Duration: four sessions Comparator: standard care	Individual outcomes: ► Functional and emotional wellbeing—better in Egroup*	
Cupisiti <i>et al</i> (2002) ⁵³	<u>Ф</u>	CKD 3b-5	20 Age: NR	▶ Diet/nutrition	E N	▶ Print	Vegetarian diet: ► Alternate between animal based conventional low protein diet and a vegetable-based low-	Individual outcomes: ▼ Opinions on diet−90% enjoyed	
							protein diet Booklets explaining general guidelines and features of the diet Duration: one session Comparator: conventional protein diet	Physiological measures: Creatinine—no change Albumin—no change I cital protein—no change Lipids—decreased Electrolytes—no change Haematorit—no change Uninary protein excretion—decreased Uninary urea excretion—decreased	
									7

	Study results							
	Study outcomes	Cognitions: Improvement in knowledge of CKD Behaviours: Modified lifestyle, diet	Intervention specific: Reduction of stress, fear Improvement in therapeutic relationships with healthcare providers, companions and multi-disciplinary team.	Oggnitions: Participation, self-care and disease related knowledge Intervention specific: Use of diary Suitability for teaching purposes	Cognitions: QOL (WHOQOL-BREF Taiwan version)—global increase Knowledge of renal function protection (checklist made by investigators)—no change	Physiological measures: Creatinine—no change BUN—no change GRR—no change Muscle weight—no change Muscle weight—no change Waist-to-hip ratio—significant decrease in E group BMI—significant decrease in E group BPI—no change	Cognitions: A Anxiety—decreased Fear—more control of fear response Stress—decreased	Health status: ESRD warranting RRT—13.9% in Egroup vs 43% in C group All cause mortality—1,7% in E group vs 10.1% in C group Healthcare: Hospitalisation—2,8% E group versus 16.4% in C group
	Description of intervention	Feducation intervention, Eight 2-hour classes Didactic and discussion Duration: 6 months Comparator none		► A diary to promote disease related knowledge, involvement and self-care ability and to promote cooperation between patient and nurse Duration: 12 months Comparator: none	Educational intervention': Handouts One 180 min workshop Individual consults Q6 month with nurse Phone number provided to participants for questions	Deserts recommended by detaining year at workshop for educational purposes, lunch boxes designed by dietician given out at the end of the workshop Duration: 12 months Comparator, none	Teaching group': ► Six 2-hour monthly group education sessions ► Booklet for future reference Duration: 6 months Comparator: none	Multidisciplinary predialysis education (MPE): Individual lectures, content-based on CKD stage Polietary counselling biannually Duration: 12 months Comparator: standard care
	Delivery format	Print Face to face PowerPoint slides		▶ Print	► Print Face to face Telephone		▶ Print▶ Face to face	▶ Face to face
	Provider(s)	Nurse, patient volunteers		Nurse	Nephrologist, nurse, dietician, social worker		Nurse, physician, technician, three expert patients	Nurse, social worker, dietician, HD/PD patient volunteers, physicians
	Intervention topic(s)	General CKD knowledge Diet/nutrition Modalities Other		■ General CKD knowledge	General CKD knowledge Diet/nutrition Physical activity Medication		General CKD knowledge Diet/nutrition Physical activity Modalities Other (e, psychosodal-impact of CKD family, finances, social life)	General CKD knowledge Diekrutrition Medication Lifestyle
	Study size Age (years)	24 Age: 64.5		58 Age: 65	66 Age: 67.4		41 Age: 60.56	573 (E=287, Cohort=286) Age: 63.4
Continued	In Target population	Q O		CKD	OKD 3		CKD 4-5	OKD 3-5
	e) Design	PP tal		Ops	<u>d</u>		PP	9 B
Table 4	Study and year (Reference)	Gutiérrez Vilaplana et al (2007) ⁵⁷		Pagels <i>et al</i> (2008) ⁵⁵	Yen et al (2008) ⁵⁶		Gutiérrez- Vilaplana e <i>t al</i> (2009) ⁵⁴	Wu et al (2009) ⁵⁸

	Study results				
	Study outcomes	Cognitions: ▶ LTMBSES—difference in mean self-efficacy score at post-test	Cognitions: ► Knowledge ► Anxiety— <i>increased</i> Individual outcomes: ► Satisfaction in group therapy	Cognitions: Knowledge of CKD scale— increases-in Egroup Behaviours: Self-care practice scale for patients with CKD—no difference between Egroup and C group but did increase over time for both groups Physiological measures: BUN/Creatinine—no change CaPO4—no change Haemoglobin—no change GFR—no change	Behaviours: ► Exercise behaviour—improved in E group Cognitions: ► Depression—score decreased (fe, improved) in E group Symptom management: ► Fatigue—score decreased in E group
	Description of intervention	*Motivational interviewing": Counselling by nurse practitioner (in addition to care by nephrologist) using motivational interviewing interviewing. Using the "Long-Term Medication Behaviour Self-Efficacy Scale (LTMBSES)" – areas with score-5 were identified and then up to five areas (picked by patient) were discussed and solutions and goals were set Duration; 6 months. Comparator; standard care	Escuela ERCA: ▼ 7 1.5 hour multidisciplinary group education ▼ 8 1.5 hour multidisciplinary group education ▼ 1.5 hour multidisciplinary group education ▼ 1.5 hour held biweekly ■ Up to 10 patients per group with family members ▼ Didactic plus discussion format Duration: not reported Comparator: none	Face-to-face SM programme: > 90min lecture with slides (3–5 people/group) > 20min individual consult > 1 week later individual reinforcement education and consultation Duration: two sessions Comparator: general maintenance	**Exercise education intervention': **Manual **1.5-hour exercise/health education course **Prishour exercise contract and exercise **Programmes **Follow-up phone calls **Ix/month for patients in maintenance phase **Ex/month for patients in action/prep stages **Ex/month for patients at precontemplation/ **Contemplation stages **Goal: workout 3-5x/week x 30 min for 3 months **Duration: 4 months **Comparator: standard care
	Delivery format	► Face to face Print	► Face to face	► Face to face ► PowerPoint slides	▼ Print Face to face ▼ Talephone
	Provider(s)	Nurse practitioner	Nurse, physiotherapist, dietician, pharmacist, psychologist, coordinators, nephrologist, patient mentors	Physician, nurse, dietician	Instructor
	Intervention topic(s)	■ Medication	General CKD knowledge Diet/nutrition Madication Symptom management Physical activity Modalities Other	■ General CKD knowledge ■ Diet/nutrition ■ Modefalties ■ Other (e, understanding and compliance with SM) Other (e) Other (e) Other (e) Other (e) Other (e)	■ General CKD knowledge ■ Physical activity
	Study size Age (years)	54 (E=28, C=26) Age: 55–59	19 Age: 58	61 (E=31, C=30) Age: 53.93-58.33	94 (E=45, C=49) Age: 73.17
pa	Target population	CKD	CKD	OKD 1-5	OKD 1-4
Continued	Design	O G	SqO	oe.	ë
Table 4	Study and year (Reference)	Wierdsma <i>et al.</i> QE (2011) ⁵⁹	Aguilera Florez et al (2012) ⁸⁰	Choi <i>et al</i> (2012) ⁶¹	Kao <i>et al</i> (2012) ²⁷

	Study results					
	Study outcomes	Intervention specific: First entry into website30% of participants entered within 365 days (total follow-up period) Average dwell time on the website—7 min Modules were ranked by frequency of selection—The tree most frequently visited pages were "Penal function calculator," Pills to avoid" and 'Foods to avoid" and	Individual outcomes: ▶ QOL—self-efficacy and SM behaviours improved Physiological measures: ▶ Renal function—no change ▶ Haemoglobin A1c—decreased postintervention	Cognitions: ► CKD self-efficacy—increased Behaviours: ► CKD SM— no change Physiological measures: ► Creatinine—marginally significant decrease ► GFR—remained stable	Cognitions: Change in patients' self-effracy to adhere to KDDQI GL after single exposure to the tool-three worsened, three improved, six no change Intervention specific: Tool acceptability -well acceptabl Congruence of patient and provider attitudes—incongruence in 4/10 cases where provider states used report but patient doesn't verify	Cognitions: ► Self-efficacy—limited effectiveness Behaviours: ► SM—limited effectiveness
	Description of intervention	 'Disease-specific safety information: Safe kidney care website – patient/family member and provider portals Education modules displayed in circular distribution to avoid prioritisation of topics Duration: not applicable Comparator, none 	'SM skills programme': Textbook Daily journal Four I hour face-to-face sessions Q2 weeks at outpatient clinic or in home Two 30 min phone or email sessions Q1 month Then Q1 month phone calls Duration: 6 months Comparator none	'SM programme': ▶ Self-monitoring workbook ▶ 5 week SM programme ▶ Weekly 90min face- to-face group sessions (6-8 patients) ▶ CKD SM video about self-regulation Duration: 5 weeks Comparator none	Dietary assessment and evaluation tool': ➤ Self-administered ➤ Obtains 24 hours food history Then evaluates diet based on KDOOI GL. ➤ Then share general tips for success ➤ A report is generated for the nephrologist to guide discussion with patients Duration: single exposure Comparator: none	Lifestyle management tool: ▼ 33-page quick start guide provided ► Patients had access to site for 4 months—patient choice to frequency of visits to website ► Duration: 4 months ► Comparator: none
	Delivery format	▼ Website	▶ Print ► Face to face ▼ Telephone ► Email	► Print ► Face to face ▼ Video	▼ Website	► Print Website
	Provider(s)	Online tool	Nurse	Nurse	Online tool	Online tool
	Intervention topic(s)	► Diet/nutrition ► Medication	■ Diet/nutrition ■ Medications ■ Physical activity ■ Comorbidities ■ Other (ie, stress management, identify supporters (family) & how they can contribute, goal setting)	▼ Other (ie, self-regulation/ selfmanagement topics)	■ Diet/nutrition	► Diet/nutrition ► Physical activity ► Lifestyle
	Study size Age (years)	108 Age: 64	30 Age: 67	37 Age 67.42	12 Age: 68	22 Age: 55.2-59.8
per	Target population	OKD 3-5	CKD 3-4 (diabetic nephropathy)	OKD 1-3a	CKD 4	CKD
Continued	Design	94 B	<u>a</u>	<u>a</u>	<u>a</u>	<u>a</u>
Table 4	Study and year (Reference)	Diamantidis et PP al (2013) ⁸²	Kazawa et al (2013) ³¹	Lin et al (2013) ⁶³	Murali <i>et al</i> (2013) ²⁸	Nauta <i>et al</i> (2013) ³²

	Study results							
	Study outcomes	Physiological measures: ▶ BP – no statistically sig difference ▶ Haemoglobin A1c–no change ▶ BMI – no change	Behaviours: ► SM (Partners in Health (PIH) instrument)—had change in certain domains	Cognitions: Nichey specific knowledge— associated with increase in knowledge Intervention specific: Peasibility of intervention— physicians found it useful and efficient but had concern regarding some of the talking points	Physiological measures: Albuminuria – <i>improved</i> GFR – <i>no change</i> 5 year absolute cardiovascular risk – <i>improved</i> PB – <i>improved</i> PB – <i>improved</i> PTotal cholesterol – <i>improved</i> Paemoglobin A1c – <i>improved</i> Cognitions: Knowledge of medications/ conditions: Pahaviours:	Medication adherence, adherence to healthy lifestyle improved	Physiological measures: GRA decline—slower in E group Heamoglobin—E group maintained more stable level compared with non-KDE group who lost 1 g/L from baseline Cognitions:	► Kidney disease knowledge (KiKS survey) – no change
	Description of intervention	'SM package': ▼ Written materials ▼ 20 min DVD ► Self-monitoring diary ► Fridge magnet with key messages ► Promitor if needed Duration: one session Comparator: standard care	Nurse practitioner intervention in primary care setting: ► SM booklet ► Initial assessment of lifestyle behaviours, SM practice, health/medication knowledge ► Individual education ► Individual	Physician-delivered education too' 1-page intervention worksheet delivered during clinic visits—take 1–2 min to administer. Duration: one session Comparator: historical group'—who developed sheet	See Walker <i>et af</i> ⁶⁴		'Kidney Disease Education (KDE)' ▶ six education classes on one on one or group basis Duration: unclear Comparator: no KDE	
	Delivery format	▼ Print DVD	▼ Print ► Face to face	Print Face to face	▶ See Walker et af ⁶⁴		Nurse practitioner, P Face to face physician assistants, clinical nurse specialist	
	Provider(s)	Z Z	Nurse, nurse practitioner	Nephrology fellows	Nurse, nurse practitioner		Nurse practitioner, physician assistants, clinical nurse specialist	
	Intervention topic(s)	 ▶ General CKD knowledge ▶ Comorbidities ▶ Lifestyle 	► Dievnutrition ► Medication ► Symptom management ► Physical activity ► Physical activity ► Other (ie, compliance)	General CKD knowledge Distriction Medication Physical activity Lifestyle Comorbidities Comorbidities	► See Walker et a ^{p4}			
	Study size Age (years)	176 (E=116, C=60) Age: NR	52 Age: 57.5	556 (F=155, Cohort-401) Age: 57	52 Age: 57.5		49 (F=25, C=24) Age: 73	
pen	Target population	Diabetic nephropathy (DM+microalbuminuria)	CKD with high risk of Progression+DM2+HTN + albuminuria	CKD 1-5	CKD with high risk of Progression+DM2+HTN + albuminuria		CKD 1-4	
Continued	Design	Σ	<u>d</u>	B	<u>a</u>		B	
Table 4	Study and year (Reference)	Thomas and Bryar (2013) ³³	Walker <i>et al</i> (2013) ⁹⁴	Wright Nunes et al. (2013) ⁶⁵	Walker <i>et al</i> (2014) ²⁴		Enworom et al (2015) ⁶⁶	

Table 4	Continued	pen							
Study and year (Reference)	Design	Target population	Study size Age (years)	Intervention topic(s)	Provider(s)	Delivery format	Description of intervention	Study outcomes	Study results
Vann et al (2015) ²⁸	<u>&</u>	CKD 3b-4	9 Age: mean NR	General CKD knowledge Diet/nutrition Symptom management Modalities Comorbidities Other (le, self-care management strategies and behaviours)	Nurse practitioner	Print Website Face to face White board	'CKD Education Programme' CKD education sessions Assessment of readiness to change CKD toolkit individualised for each participant Collaborative goal setting between rurse practitioner and patient Information booklet with websites listed Patients met with rurse practitioner for 60 min Duration: over six visits Comparator: none	Cognitions: CKD-related knowledge— improved Behaviours: Self reported behaviour change—improved	
Cupisiti et al (2016) ⁶⁷	Obs	OKD 3b-5	823 (E=305, O=518) Age: 69–74	▶ Diet/nutrition	Dietician	Face to face	Nutritional Treatment Penal dietician assessed dietary habits using 3-day dietary recal & performed an intervention tailored to the needs/clinical features of the patient Progressed from 'normal' diet → low protein diet → very low protein diet depending on needs Duration: at least 6 months Comparator: standard care	Physiological measures: Phosphaturia—lower in E group Healthcare: Furosemide use—lower in E group Calcium free phosphate bind use—lower in E group ESA use—lower in E group SSA use—lower in E group Individual outcomes: Dietary satisfaction are—lower in regroup Individual outcomes: Dietary satisfaction are majority of E group outcomes:	
Ong et al (2016) ⁶⁸	đ.	CKD 4-5	45 Age: 59.4	Medications Symptom management Comorbidities Other (e, tracking lab results)	Mobile application	Smart phone application	'Smartphone based SM system' Application generated personalised patient messages based on prebuilt algorithms Duration: 6 months Comparator: none	with the date of the property	
Penaloza- Ramos <i>et al</i> (2016) ²⁵	Obs	HTN (BP>130/80)+CKD stage three or CVA/TIA or DM or MI or angina or CABG	NR e Age: NA I G	► See McManus et al ⁴⁷	General practitioner, patient	See McManus et af?	► See McManus et al ⁴⁷	Healthcare: ► Cost effective—yes	

Not applicable.

Outcome improved post intervention.

Outcome unchanged post intervention.

Outcome worsened post intervention.

Outcome had mixed results (some improved and/or some worsened and/or some did not change).

BMI, body mass index; BP blood pressure; C, control; CALD, culturally and linguistically diverse; CHD, coronary heart disease; CHEERS, Controlling Hypertension: Education and Empowerment Renal Study; CKD, chronic kidney disease; CVA, cerebrovascular acident; DBP diastolic blood pressure; DM, diabetes melitus; E, experimental; eGFR, estimated glomenular filtration rate, ESA, erthropolesis stimulating agents; ESRD, early stage renal disease; HTM, hypertension; MM, mixed methods; and reported; Dbs. observational; PP, pre-post intervention; QE, quasi-experimental; QDL, quality of life; RCT, randomised controlled trial; RRT, renal replacement therapy; SBP, systolic blood pressure; SM, self-management; TIA, transient ischaemic attack.

Table 5 Summ	Summary of qualitative studies	ies			
Study (Reference)	Number of Target population participants	Number of participants	Aim/Intervention	Methods	Summary of findings
Blickem <i>et af</i> ²¹	CKD stage 3	20	'To explore the experience of patient-led assessment for network support (PLANS) from the perspectives of participants and telephone support workers.' (p. 1) Intervention: see table 4 Blakeman et al 46	Interviews and focus groups: no analytic methodology discussed	 Mixed reception from participants Formulation of 'health' in everyday life (ie, participants unaware of having CKD or its significance—confused about relevance of PLANS) Trajectories and tipping points (ie, engagement in PLANS depended on participants' stage of life—either could influence trying new things or disrupt routines) Trust in networks (ie, unwillingness to seek support, intrusive, others saw improved awareness/access to local resources; tailored support)
Heiden <i>et al</i> ⁶⁹	CKD predialysis, dialysis, transplant	വ	To identify participant's perspective regarding a 'web application prototype to help make decisions regarding diet restrictions and phosphate binder dosage.' (p. 544) Intervention: Website tool for patients that included three components—diet/fluid education; diet registry and phosphate binder decision support tool.	Interviews: no analytic methodology discussed	 ▶ Benefits: Education tool increased insight and understanding Assisted in tracking and choosing best food alternative Decision support for binder dosage Limitations:
Jansen e <i>t al</i> ⁷⁰	CKD stages 4–5	_	Feasibility of 'a psychosocial intervention to assist ESRD patients and their partners in integrating renal disease and treatment into daily activities, primary work and thereby increasing autonomy.' (p. 280) Intervention: group teaching and handbook regarding coping strategies and goals based on self-regulation, social learning and self-determination theories.	Interviews: no analytic methodology discussed	 ▶ Benefits: Group included predialysis and dialysis patients Leaders addressed individual needs, situations and questions Limitations:

Study Number of Reference Target population participants Aim/Intervention			
Type 1 or 5 (3 face-2 DM with to-face microalbuminuria interviews) CKD stages 2-4 26 with diabetes and cardiovascular disease CKD stages 2-4, 39 with coexisting diabetes and hypertension		Methods	Summary of findings
CKD stages 2–4 26 with diabetes and cardiovascular disease CKD stages 2–4, 39 with coexisting diabetes and hypertension	~	Questionnaire and interview: no analytic methodology discussed	 Mixed responses DVD—content distressing and took effort to use Written material useful, but need to elaborate on seriousness of disease Package helped change behaviour—stop smoking, monitoring DM
CKD stages 2–4, 39 with coexisting diabetes and hypertension	d p b	Interviews: Ritchie and Spencer thematic approach	 ► Attitudes towards taking medications (le, appreciate importance of taking; medication burden; concern with the number of medications, effectiveness and side effects of medications. ► Having to take medications (le, behaviours and family support to assist taking medications; forgetting and non-adherent; motivation to take to prevent becoming worse) ► Impediments to chronic illness medication self-efficacy (le, lack of knowledge regarding medication; strong faith in physician's advice; multiple medications too overwhelming; cost)
Williams et al	onal n.'	Interviews: Ritchie and Spencer thematic approach	 Importance of health (ie, determined the degree of health behaviour; altered medications or use of complementary medicine to control health) Perceived seriousness of disease (ie, thinking about mortality; comorbidities complicate care; acute illness with chronic conditions) Perceived threat of disease (ie, want to learn about disease control earlier; symptom management; looking for reasons to explain why ill)

CKD, chronic kidney disease; DM, diabetes mellitus.

protocols, changing health behaviours, shared decision making with healthcare providers), ^{7 75} along with evaluation of the cost-effectiveness and resource utilisation.

The majority of studies did not identify a single primary outcome but rather multiple outcomes. We found that physiological outcomes (ie, blood pressure) were the most commonly reported and symptoms were the least mentioned. These findings demonstrate the lack of patient-driven outcomes that may be important to them, for example, a patient's individual health goals across a variety of dimensions (ie, symptoms, mobility, social and role function in the family or community) that could possibly maximise their quality of life. Work by Tong et al (2015) highlights this concept, where patients with CKD are more interested in treatment choices that influence non-traditional clinical outcomes such as impact on family and lifestyle.⁷² A holistic approach should be considered where mental and psychosocial outcomes are investigated, rather than just physiological endpoints.

Our findings from the qualitative studies looking at patient perspectives are inconclusive because of the limited number of studies and the heterogeneity of the interventions. Havas *et al*¹² similarly reported a lack of research related to patient perspectives on self-management in CKD.¹² There is also a lack of qualitative studies overall, which could provide valuable information regarding attitudes and challenges of self-management interventions from the perspective of both providers and patients.

Strengths of our study include the comprehensive nature of our search, inclusion of all study designs and consideration of self-management features that have not been investigated previously. We also engaged patient partners in determining the research question, advising us on search terms, grey literature sources and reviewing the results to ensure we captured and reported the data meaningfully. One of the main limitations was the challenge in synthesising the data given its heterogeneous nature. To address this challenge, the two reviewers used two standardised tools TIDieR¹⁹ and the EPOC tool²⁰ to independently extract data and independently coded the outcomes into categories using the revised Self-and Family Management Framework . 6 Also, we were unable to assess the self-management outcomes in terms of sustained changes in behaviour, physiological and health status. A final limitation was our inability to draw conclusions regarding the most effective self-management intervention for adult patients with CKD, keeping in mind that our aim was to review the breadth of the current literature and present the gaps that exist.

Overall, we found considerable variation in self-management interventions for adults with CKD with respect to their content and delivery as well as the outcomes assessed and results obtained. Major gaps in the literature include the lack of patient engagement in the design of the self-management intervention, along with the lack of a behavioural change theory to

inform their design. Our future research will incorporate intervention frameworks to codevelop and evaluate a self-management intervention based on a sound behavioural theory involving our national patient partners, specialists, primary care providers and decision makers.

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Data sharing statement The following data will be available, study protocol and analysis plan, to anyone who wishes to access them and will be available immediately following publication from the corresponding author.

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