Cultural influences on physical activity and exercise beliefs in patients with chronic kidney disease: ‘The Culture-CKD Study’ — a qualitative study

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ABSTRACT

Objectives This study used a mixed-method approach to explore cultural and ethnic influences on the perception of, and decision to engage with or not to engage with, physical activity and exercise therapy in patients with chronic kidney disease (CKD).

Design Qualitative research was conducted through the use of semistructured interviews and focus groups. Self-reported physical activity levels were measured using the General Practice Physical Activity Questionnaire (GPPAQ), and self-efficacy for exercise with Bandura’s Self-Efficacy for Exercise Scale.

Setting This study was conducted in a non-clinical setting of a single National Health Service Hospital Trust between April 2018 and July 2019.

Participants Participants >18 years of age with a diagnosis of CKD, from black African, black Caribbean, South Asian or white ethnicity were eligible for the study. 84 patients with a diagnosis of CKD (stages 2–5), aged 25–79 (mean age 57) were recruited. Semistructured interviews (n=20) and six single-sex, ethnic-specific focus group discussions were undertaken (n=36).

Outcomes Primary outcome was to explore the perceptions, attitudes and values about exercise and physical activity in different ethnic groups through qualitative interviews, analysed using an inductive thematic analysis approach. Questionnaires were analysed using Pearson correlation to determine if there was a significant relationship between the self-efficacy and GPPAQ levels.

Results Qualitative analysis provided four primary themes: I am who I am, Change of identity, Influences to physical activity and exercise beliefs in patients with CKD, and Support and education. Quantitative analysis using Pearson correlation revealed a significant correlation between GPPAQ levels of activity and self-efficacy to regulate exercise behaviour (r=−0.40, p=0.001).

Conclusion Understanding the cultural, attitudes and beliefs of individuals with CKD from a variety of ethnic backgrounds is complex. Understanding of patients’ experiences, thoughts and beliefs may be of relevance to clinicians when designing CKD exercise services.

Trial registration number NCT03709212; Pre-results.

BACKGROUND

Chronic kidney disease (CKD) affects 1 in 10 people worldwide. CKD is associated with increased risk of cardiovascular disease (CVD),1 reduced quality of life (QOL),2 decreased physical function and muscle wasting. CVD remains the leading cause of death for individuals with CKD.3

Certain ethnic minorities have been found to be at higher risk of developing CKD.4 5 In the UK, people from ethnic minority groups are more likely to suffer from obesity, type-2 diabetes and hypertension, which are all risk factors for developing CKD.6 As kidney disease progresses, individuals from ethnic minorities are three to five times more likely to start dialysis than Caucasians.7 In the UK, over a third of people waiting for a kidney transplant are from ethnic minority communities, while donors from these groups only make up 7.2% of the National Health Service (NHS) Organ Donor Register.7

There is a broad body of evidence to highlight the benefits of physical activity (PA) and exercise within the general CKD population, including positive outcomes on cardiovascular health, physical function and QOL.8
guidelines recommend that all patients with CKD (patients with CKD not on dialysis, as well as those on haemodialysis and peritoneal dialysis (PD)) perform aerobic exercise training at mild-to-moderate strength (40%–60% of $\text{VO}_{2\text{peak}}$) for 20–60 min/day for 3–5 days/week. These guidelines also recommend resistance exercise training for these patients, at 70%–75% of one-repetition maximum for a minimum one set of 10–15 repetitions on 2–3 days/week. The National Kidney Foundation Kidney Disease Outcome Quality Initiative guidelines recommend that individuals diagnosed with CKD should engage in a PA programme for 30 min, 5 days per week. Typically, people with CKD report high levels of physical inactivity, and a number of barriers have been identified that may influence engagement with exercise training including poor health, fear of injury, lack of guidance from healthcare professionals and a lack of facilities.

A recent study undertaken by our research team, evaluating an exercise intervention with kidney transplant recipients, revealed that 11 of the 18 individuals who did not complete the intervention were from ethnic minorities. Participant feedback identified that potential challenges involved undertaking exercise in a mixed gender group, where culturally appropriate clothing was essential. Given overall low levels of PA in the CKD population, it is imperative that ways are established to engage all individuals with CKD with lifestyle changes that can facilitate an adherence with current guidelines. A number of studies have evaluated the influence of cultural beliefs on engagement with PA in the general population. Others have evaluated the cultural influence on PA in cardiac rehabilitation programmes in the USA. To our knowledge no studies have evaluated the influence of culture and ethnicity on engagement with PA in people living with CKD.

Reducing health inequalities, particularly preventing the development and progression of kidney disease may help alleviate the burden of care to the NHS. One of the research strategies highlighted by the UK Kidney Research Consortium is to address health inequalities by increasing our understanding of the barriers for people in different ethnic groups to accessing high-quality care for CKD, and its risk factors. Understanding the beliefs and cultural influences towards PA and exercise in ethnic minorities with a diagnosis of CKD is essential to ensure that interventions are designed to be culturally sensitive, and are appropriate and accessible for all people living with CKD in the UK. The aim of this study was to explore cultural and ethnic influences on the perception of, and decision to engage with or not to engage with, PA and exercise therapy in patients with CKD.

**METHODS**

**Objectives**

To explore cultural and ethnic influences on the perception of, and decision to engage with or not to engage with, PA and exercise therapy in patients with CKD.

**Study design**

Qualitative data were collected utilising individual semi-structured interviews (SSIs) and focus groups. SSIs were conducted in the first instance. Gender and ethnically specific focus groups were included to further probe how culture shapes beliefs about exercise and PA in order to create a deeper understanding of the perceptions surrounding this. Study protocol can be viewed through online supplemental material S1.

All participants independently completed the General Practitioner Physical Activity Questionnaire (GPPAQ), and Self-Efficacy for Exercise Behaviours (SEE) Scale, prior to attending the SSI or focus groups. This was to establish participants prior engagement in PA, as well as to facilitate purposive sampling, with different levels of PA and exercise represented in the data.

GPPAQ was utilised as a self-reporting measure to evaluate levels of PA and occupation, reported over the past week. It involves three questions, and provides a simple, 4-level PA Index (PAI) categorising patients as active, moderately active, moderately inactive and inactive. Figure 1 describes the scoring algorithm. The GPPAQ is a validated screening tool utilised in primary care settings, as well as being validated within the CKD population in comparison to accelerometry. The GPPAQ can accurately identify those not sufficiently active, identifying individuals most at need of a PA intervention.

The SEE scale, provided information on each participant’s readiness to commence exercise. Self-efficacy scales for eating and exercise behaviours show evidence of being reliable and valid in the literature. The questionnaire consisted of 18 questions utilising a Likert scale between 0 and 100, with 0 being low confidence and 100 high confidence. Score was calculated by adding the total score in order to gain a cumulative total. Scores were categorised into low (0–600), medium (601–1200) and high (1201–1800).
Sample and setting
Recruitment occurred between and July 2019. Individuals aged 18 years of age and over, with a diagnosis of CKD, representative of one of following ethnic groups; BA, BC, SA, and W were considered eligible for inclusion. CKD was defined as abnormalities of kidney function or structure present for more than 3 months. The definition of CKD includes all individuals with markers of kidney damage, or those with an Estimated Glomerular Filtration Rate (eGFR) of <60 mL/min/1.73 m² on at least two occasions 90 days apart (with or without markers of kidney damage). Individuals with an uncontrolled medical or psychiatric condition were excluded.

Participants
Participants were identified using purposive sampling, to ensure that the sample included participants from a range of ethnic backgrounds, age, gender, renal modality and stage of CKD. Individuals were screened and approached during routine nephrology outpatient clinical appointments at a single NHS hospital trust. Ethnicity and gender were established from medical records, and confirmed on initial discussion with participants. Each individual was given 24 hours to review the participant information sheet and following this, provided written consent. Figure 2 displays the Consolidated Standards of Reporting Trials diagram of participant flow. Questionnaire data were translated for non-English speakers, and SSI and focus group interviews were conducted with multilingual facilitators, to ensure that all data collected were diverse and representative of the different ethnic groups of participants within this study. All qualitative interviews were held in a non-clinical setting of a UK NHS hospital.

Data collection
Topic guides were based on a review of the literature and in collaboration with the trial committee (SG, JM, JG, EC, PDH, PO), including the patient advisory group. Focus group topic guide questions were informed by the early individual interview data (online supplemental materials S2 and S3).

SSIs were undertaken in the first instance with 20 participants, and lasted on average 42 min. The primary researcher conducted all SSIs and focus group interviews (JM). Please see online supplemental material S4 for further descriptive details. Focus groups were undertaken with six groups, which were single-gender and specific to ethnicity. These were conducted by a facilitator and a moderator (JM, LH). Each group session consisted of 4–8 participants and lasted on average 58 min. SSI and focus group interviews were undertaken until data saturation was reached.

The primary researcher kept a reflexive diary, to maintain accountability and transparency of the thoughts, feelings and experiences of the interview data. Field notes were made to capture impressions and any contextual information from the SSI and focus group interviews. This was used to inform data analysis.

Analytical approach
All qualitative interviews were digitally recorded, transcribed and translated verbatim, and analysed using an inductive thematic analysis approach, and strategically followed guidance as demonstrated in online supplemental material S5. SSIs were conducted and analysed in the first instance, to inform the focus group topic guide. Analysis was achieved through qualitative method triangulation. NVivo V.11 software (QSR International Ltd, V.11, 2016) was used to facilitate data management.

JM conducted the primary analysis of data. Transcripts were read a number of times and initial codes generated. Initial codes were then collated into similar groupings in order to create potential themes and thematic maps. Codes and themes were refined to enhance validity and clarity until consensus was achieved across the trial committee (JM, SG, JG, EC). This allowed refinement of codes into primary descriptive themes.
GPPAQ and SEE scale scores were entered into a secure database and analysed using Pearson product–moment correlation coefficient within SPSS V.25.0.

Qualitative and quantitative data were analysed separately. Qualitative and quantitative results were merged using triangulation to provide joint summaries in order to detect areas of convergence/divergence to the research question.28–30 See online supplemental material S6 for Consolidated criteria for Reporting Qualitative research checklist.

**Patient and public involvement**
The involvement of patients has been integral to this research. This involved patients of different ages, stages of CKD and representative of the different ethnic groups involved in this study. Initial meetings revealed several important points which required consideration and informed the study proposal. The need to host the focus groups in a non-NHS environment, and the use of a single-ethnicity, and single-sex focus group structure were identified as important factors to consider in this particular study. The patient advisory group participated in face-to-face meetings to review the protocol and advise on the lay summary, data collection methods, interview topic guides, study documentation and results.

**RESULTS**

**Quantitative**
Pearson correlation revealed a significant correlation between GPPAQ levels of activity and Self-efficacy to regulate exercise behaviour (r=−0.40, p=0.001). This demonstrates that increased self-efficacy was associated with higher levels of PA. Correlation is negative because of direction of comparison between the two surveys.

**Integrated mixed-methods analyses**
The integrated qualitative and quantitative findings suggest that individuals who reported motivations to exercise had higher self-efficacy to regulate exercise behaviours and PAI scores in comparison to individuals presenting with challenges to PA and exercise. Those who identified the need for external support identified a lower self-efficacy, although this was not reflected in a low PAI score. Please refer to online supplemental material S7 for a joint display summarising findings from the integration of qualitative and quantitative results.

**Qualitative**
A total of 20 participants were recruited for SSI and 64 participants recruited for six single-sex, ethnicity-specific focus groups. Please see table 1 for detailed participant characteristics, and online supplemental material S8 for characteristics for each ethnic group. Individuals (n=141) were approached to participate in the study, of which 64 declined to consent. For the individual interviews, there were no refusals to participate. Of the 64 participants who consented to participate in the focus group discussions, 28 were unable to attend the dates organised for discussions. Please refer to online supplemental material S9.

**Major themes and sub-themes**
Thematic analysis revealed four major themes; ‘I am who I am’, ‘Change of identity’, ‘Influences to PA and exercise and ‘Support and Education’. Please see figure 3 for a thematic map.

**Theme 1: ‘I am who I am’**
Theme 1 encompassed individual identity, with varied responses voiced from the participants. Overall,
individuals associated their identity with the environment and location they were raised in, with their family, their education and their work. Examples of quotations can be found table 2.

Perception of cultural identity was voiced by many of the individuals who participated in the research. The importance of family and intergenerational support was particularly evident in SA participants. This frequently arose in data expressed by SA women, associating their role with the home environment, as primary care givers. ‘In the Asian culture, the woman is the home maker, so she’s expected to be at home doing the work’ (F, SA). This contrasted with the BA and BC participant discussions, which focused around associations with activity linking to environment, food preparation and family events. ‘Coming from the Caribbean, I can only speak for myself but I had to do a lot of walking as we have a farm with lots of work and all of that...’ (F, BC).

Previous exposure and experience of PA and exercise varied between participants, as illustrated by table 2. Overall, previous exercise exposure linked into exercise engagement as an adult ‘I don’t come from an exercising, sporty type background...But exercise was never really my thing’ (M, W).

Table 2 Theme 1: ‘I am who I am’

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<tr>
<th>Major themes</th>
<th>Subthemes</th>
<th>Examplar quotations</th>
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<tr>
<td>Individual identity</td>
<td></td>
<td>‘Yeah [laughs] ‘cos you know there is a lot of emphasis on errm culture you know I was thinking, well I’m happy, I was born in St Lucia, it is a very beautiful Island. I mean growing up education wise it was the colonial system, so we all followed it all, the books and things in the literature. I have lived here in Britain for a long time now so I have British citizenship, you know.’ (M, SA)</td>
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<td>Cultural identity</td>
<td>SA Female</td>
<td>‘But I think, I mean and I’m not speaking for everyone but in the Asian culture the women is the home maker, so she’s expected to be at home doing the work.’ (F, SA) ‘Asian women are...They’re busy with their children, the housework. You know like you said everything comes first, they come last.’ (F, SA)</td>
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<td></td>
<td>BA and BC</td>
<td>‘Well no I think errm we encouraged my mum to do it. As we were growing up, it’s something with West Indian families they don’t really exercise and stuff like that.’ (M, BA and BC) ‘Coming from the Caribbean, I can only speak for myself but I had to do a lot of walking as we have a farm with lots of work and all of that, you know, so I always did it.’ (F, BC) ‘I was brought up here but my mum would drag me into the kitchen, ‘don’t think you’re lying down, you could spend all day, you getting up.’ Everything she done you would have to be there, even if it was just to pass an onion from here to there, you had to do it so you were moving.’ (F, BA)</td>
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<td>Previous experience of physical activity and exercise</td>
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<td>‘I mean I haven’t been an active person for many years before I even went on dialysis but I remember when I was at school I used to do cross country running. Right, I used to be in the netball team and stuff like that and I think as I got older and I had to go to work and stuff like that that I didn’t do it and then things are not... Things are different to how they are now to how they were when I was growing up and when I started to work and things like that. They wasn’t pushing people to do exercises, it’s only recently in the last ten years they’ve started to get everybody out there.’ (F, BA and BC) ‘Yeah, we, we, we, with that time in their age I can move back with a little bit of history, I grew in the British...I was born in the British Empire. And in the British empire in the school they, they encouraged a lot of activities so by, by those times, our parents were encouraged to dancing. Communal dancing.’ (M, BA and BC)</td>
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Psychologically it was quite hard to come to terms with... (F, and symptoms experienced as renal disease progresses. It's just fatigue and nothing else. They said you must do something, you can do it, not that you know. So it's just fatigue and time. I dialyse in the mornings so I wake up at 7 o'clock and the night before I can't sleep because I'm scared that I ain't going to wake up and the thing... So the next day it's a workout because I'm just too tired. I sleep for 12 hours like you said...’ (F, BC) "Oh when they said to me like you’re going on, we think we’re going to put you on dialysis right, let's say it was Tuesday that day and like Monday next week we're thinking of calling you in to start dialysis. I think I disappeared for a week, I just couldn't get my head round it that I had to do this thing 3 days a week’ (F, BC) ‘But right now my experience is err if there's a dialysis patient right now that's like me. You know the dialysis is sometimes is... You are well and sometimes you are not well. Sometimes for, for example in my experience, 60%–70% is unwell.’ (M, SA) ‘You just look forward to the transplant so you can hopefully stop it. That's one of the reason that you try to exercise to be as fit as you can, so that you can have the transplant.’ (M, W) ‘That's why I hope some day that there is a kidney donor for me, and I have that transplant. And I pray to God that from tomorrow that Guy's Hospital call for me and say (P) come here and have a donor from the kidney and that is important. That's all, that is my experience.’ (M, SA) ‘So sometimes with the machine in the night time, and when it's draining I wake up because I've got drain pain. So I might wake up for 20 min with drain pain and then maybe if I’ve turned to one side it’s 'bleep, bleep, bleep.' I find my sleep is very disrupted with the drain pain and also the bleeps.’ (F, BA) ‘So what they’re saying is that my appetite will get better, and to be honest it hasn’t got better, so I'm just eating for the sake of eating.’ (F, BA) ‘I think with my friends as well I’ve been more anxious post transplant than pre, but everyone is that little bit more sympathetic before when they can see that you’re ill/The minute I had my transplant and I was more you know, I suppose reasonably normal and then that's it you’re one of us now. Sometimes I’ve had to stop and say hold on a sec, don’t include me in that, don't ask me to do that.’ (M, W) ‘Well, you know I started physio after the operation, but before the operation it was bad. I could hardly walk a few distance, you know a few metres and that’s it. And then I get very tired. You know no matter what you do, I had the strength but my legs they kept getting very tired and sometimes if there was a slight inclination or to go up the stairs. You just think of it and then sit down and that’s it. So when you do that you feel pain and you don’t want to go through that again.’ (M, SA) ‘But now I’m feeling tired, most of the day I need to be laying down err even for cooking, daily activities like cooking my...either my sister or my daughter help out.’ (F, SA) ‘Psychologically illness as itself will put you back because it becomes a handicap, illness becomes a handicap at times although you, you know especially if you are working and things because you know that you need to take time off during your working life.’ (M, SA) ‘I don’t know it was a bit strange because I went to my diabetes appointment and I met a doctor and I didn’t think about, I wasn’t thinking about kidney disease or anything like that and I just met a doctor at a diabetes appointment and just out of the blue she said I had kidney disease so I was thinking like I’ve been coming here for months and no one has mentioned that to me until this particular day.’ (F, BC)
For individuals under the ‘Low Clearance Clinic’, most frequently who are preparing for commencement of renal replacement therapy, a number of challenges were raised. This included; disturbance to appetite, fatigue and the psychological impact of preparing for future treatment modalities. ‘Yeah, with kidney [clears throat] failure, you, it comes as a shock when your doctor tells you you’ve got a kidney failure and then ah they say, ‘Okay, you will be on dialysis’ is something which will always be very, very, very new to yourself, your family and you feel phobia for the first time when the announcement comes to you’ (M, BC). Experiences of those individuals undergoing haemodialysis treatment were focused on fatigue and loss of freedom, with attendance at hospital three times per week for treatment. For individuals undergoing PD, complications with infection, particularly peritonitis, were voiced as a challenge. Alongside this, sleep disturbance for those receiving automated PD overnight impacted their ability to continue with regular PA and caused disruption to day to day activities. ‘So I might wake up for twenty minutes with drain pain, and then maybe if I’ve turned to one side it’s “bleep, bleep, bleep”. I find my sleep is very disrupted because with the drain pain and also the bleeps’ (F, BA).

There were contrasting views of transplantation expressed from those pretransplant, and those who had received a kidney transplant. Of those with earlier stages of CKD, renal transplantation was often perceived as a solution to symptom burden. ‘You just look forward to the transplant so you can hopefully stop it’ (M, W). However, those with a renal transplant continued to report challenges, primarily with medication burden, time with medical appointments and the lack of understanding of this modality with friends and family. ‘I think with my friends as well I’ve been more anxious post transplant, everyone is a little bit more sympathetic before when they can see that you’re ill’ (M, W).

A common concern of participants was the impact of CKD on lifestyle choices and the adaptations that individuals felt were required. ‘The time comes when you’re feeling depressed with your kidney condition and your ill health…I used to teach and work in a school for 18 years. Now I’m not doing anything but staying at home’ (F, SA).

**Theme 3: influences to PA and exercise**

A number of participants voiced the perceived benefits to PA and exercise. Primary codes emerging from the data highlighted motivators and challenges to partaking in PA or exercise training across the participants. Please see table 4 for further information. The majority of individuals identified the role of exercise in maintaining muscle strength and preserving functional independence. A small number of those participants taking part in focus group discussions reported an improvement in health outcomes associated with a change in lifestyle, including reduction in blood pressure and decreased dependence on diabetic medication use. ‘My GP is very pleased with the results of that daily walking…My blood pressure medication, I’ll probably be able to get off this very soon’ (F, SA).

There were a number of challenges to PA and exercise engagement. A lack of time due to numerous medical appointments, childcare and work commitments were the most commonly reported challenges. Access to community exercise services was voiced as a barrier, alongside the cost implications often associated with accessing local gym and community centres. Fear avoidance behaviours were evident, particularly concerns about injury or aggravating their kidney condition through exercise. ‘It’s very easy to get things wrong if you don’t know what you’re doing’ (M, W).

Ethnic-specific influences to PA and exercise provided insightful information. Those of BA and BC ethnicity reported a primary motivator to exercise being weight management. ‘Exercise and diet go really well together, it would be good for both teams to combine to help’ (F, BA and BC). The importance of diet, including providing food for family, social gatherings and events was evident. Consensus within focus group discussions highlighted the conflict between cultural dietary choices, and a limitation of dietary choices that were associated with a reduction in renal function and weight management. ‘What causes the problem for us African’s is our food. Because all the food we are eating in Africa is carbohydrates...’ (M, BA).

Those of SA background showed a preference for PA and exercise outside of the home environment, including within community centres. Providing exercise classes at flexible times of day was voiced as important to allow individuals to access services around other working or childcare commitments. Participants identified a strong cultural value to provide care for those unwell. Analysis also revealed a cultural view to avoid exercise or PA if someone is unwell. ‘So in India if you are ill, they will say sit. Don’t do anything’ (M, SA).

Overall PA was described as incidental activity, for example cleaning or gardening. ‘So physically active would be, say you know using the stairs instead of the lifts, it would be running around the house...’ (F, W). Exercise was perceived as more intensive activity, often through a formal sporting activity such as running or football. ‘Exercise can be anything as I say from swimming, running, that sort of thing’ (M, W).

Individuals of BA and BC ethnicity reported greater incidental activity, linked to the environment and access to the community. Individual interviewees who immigrated to the UK, identified a change in activity behaviour and an increase in sedentarism due to a greater convenience of amenities. ‘When you come from different countries, as he said, when you are from Africa, you walk, yeah you know you walk, because there is no two ways about it, when you come into a comfortable environment I think we all get lazy, and you just start falling off’ (M, BA).

Males from a W ethnicity described past experiences and associations with exercise linked to social events and formal sporting activities for example, football and rugby. This formed an important aspect of exercise and over half the participants voiced a preference to exercise within a group environment. ‘For me it’s definitely the camaraderie.'
You know, football with your mates, then your pub cricket teams and stuff” (M, W).

**Theme 4: support and education**

Multifaceted support was identified as a key influence to engagement in PA and exercise. Examples of this theme are highlighted in table 5. Education and awareness of the role of exercise within the management of CKD was highlighted as a need across the participants. This included both raising the profile of specialist renal physiotherapy services, as well as increasing the presence of physiotherapists within the clinical setting to form part of routine care for individuals with CKD. ‘When I used to have every 3 months a check up, I wasn’t given any guidance to do exercise. Or how exercise could make a difference in renal failure.’ (F, SA). Individuals identified the requirement for education and guidance early within their treatment
Table 5  Theme 4: ‘support and education’

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<thead>
<tr>
<th>Major theme</th>
<th>Subtheme</th>
<th>Quotation</th>
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<tr>
<td>Peer support</td>
<td>Online support platforms</td>
<td>‘As I mentioned before we started this, that my biggest support is the Facebook kidney care group where I can reach out to other patients, even though I don’t know them, I can put a question out there on the page and just say ‘has anyone experienced this, or what do you think about that?’. Sometimes I will get a hundred comments of different tips and tricks and I think that’s my best support, kidney Care UK on Facebook.’ (F, W)</td>
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<tr>
<td>Peer support</td>
<td>Peer support platforms</td>
<td>‘So yeah I definitely think you’re touching on a point there, that on the medical side of it you have to go for the days learning, don’t you before you go on dialysis or before you have your transplant and that, I just think both need counselling. There should be people in a room like this discussing it with other people who say they’ve been through it (exercise), it’s brilliant, it’s free, go for it. You’re only finding out about yourself, it is nothing to worry about and again, you may have picked up three new people here today, so if they are arranging informal meetings for patients then I think it would be a blessing for a lot of people don’t you?’ (M, W)</td>
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<tr>
<td>Peer support</td>
<td>Group exercise</td>
<td>‘Coming to classes it just encourages you to do more. I think. And erm, I’m lacking that, I’m lacking the motivation. I’m lacking the motivation of that football team, if you like. Because you create the programme. I’m lacking that.’ (M, SA)</td>
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<td>Group exercise</td>
<td>Inclusivity</td>
<td>‘Personally, I think it’s good with a mixture of ethnicities and age, and men and women because I think we are all in it together sort of thing. The commonality is we’ve got an illness and I think that there’s a kind of empathy that goes with that. Personally I’m not in favour of any segregation.’ (M, W)</td>
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<td>Family support</td>
<td>Family providing motivation</td>
<td>‘Erm, very much so, because like my sister we used to go, when I was able to so we used to go to the gym. Sometimes when I didn’t want to, ‘cause when she was going she would kind of like pick me up and say, ‘oh I’m going today, are you coming?’ (F, BC)</td>
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<tr>
<td>Ethniciy-specific</td>
<td></td>
<td>‘I mean, well according to our cultural values we are very family based. We live in a family group, so I think if I lived on my own then I would have to face all those situations and I think that would have been difficult for me.’ (F, SA)</td>
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<tr>
<td>Renal specialist Physiotherapy support</td>
<td>Specialist physiotherapy as part of routine care</td>
<td>‘When I used to have every 3 months a check up, I wasn’t given any guidance to do exercise. Or how exercise could make a difference with renal failure.’ (F, SA)</td>
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<tr>
<td>Renal specialist Physiotherapy support</td>
<td>‘Prehabilitation’</td>
<td>‘I think before that I had the operation I wanted to be as fit as possible because the fitter I was going in the quicker I recover coming out, which I think is probably quite true because I was walking quite well within a couple of days after the operation up and down the hospital aisles and things.’ (F, W)</td>
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<td>Awareness and exposure to specialist physiotherapy services</td>
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<td>‘I think you you promote, promote as this male said promote more in the hospital in the environment in the renal centres, in the clinics get people interested involve there are so many people good people here that will spread the word as well to say yeah look exercise is really good, you know.’ (M, BC)</td>
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<tr>
<td>Patient education around the benefits of exercise in CKD</td>
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<td>‘Erm I think education is key. I think I’ve learnt from my experience that I need to be healthy, it’s not just you know one sided. I can’t just rely on doctors to treat me I have to help myself so I think education is important.’ (F, SA)</td>
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CKD, chronic kidney disease.

pathway, to improve physical function and to aid with self-management. ‘But I think it is something that should be introduced early so that you’re prepared for it when the time comes and you need it properly’ (F, BC).

Participants voiced the importance of peer support, involving access to online digital support platforms, as well as peer support groups within their local community. Overall consensus revealed that individuals felt a group exercise environment to be beneficial, to aid motivation, increase social interaction and enjoyment. Access to renal-specific exercise services were also highlighted, to increase confidence with exercise, and for access to peer support. ‘It is therapeutic because you talk to people with the same problems. Discuss your feelings and talk and it’s nice to share. Always helps you’ (F, SA).

SA participants emphasised the importance of involving family within their care, and the value of this support to encourage and facilitate exercise behaviour. ‘Mind you I find that family makes a lot of difference…Everyday ringing me to find out how I am doing. That’s a big support for me’ (F, SA). One participant raised a concern of exercising within a mixed gender exercise group, due to concerns
with appropriate clothing however, this was not felt to be a challenge for the majority of participants. Overall cross-sectional analysis revealed the preference for inclusivity with exercise interventions. Individuals with different cultural backgrounds and genders exercising together was seen as a benefit to engagement in exercise.

**DISCUSSION**

In this qualitative study, we found the influences to PA and exercise in individuals with CKD from different ethnic backgrounds to be complex. In accordance with other research, participants within this study felt that the main motivators to engage in PA and exercise were taking ownership of health, health maintenance and professional support and education. Challenges varied depending on renal modality but some common barriers included symptoms associated with CKD and treatment undertaken, comorbidities, time, lack of confidence and fear avoidance behaviours associated with lack of specialist education and support. These themes, have been emphasised in other recent qualitative research. Similar barriers have also been demonstrated across a number of other long-term conditions, including chronic obstructive pulmonary disease and heart failure.

This study suggests that individuals from across the representative ethnic groups were more likely to report motivators to exercise if they had higher levels of self-reported self-efficacy and PA levels, in comparison with individuals who reported a number of barriers to exercise. This finding resonates with previous research into self-efficacy behaviours in relation to exercise in CKD, which demonstrate that higher cardiorespiratory fitness and self-efficacy levels were associated with an increased odds of being active. The identification of the requirement for support was indicated by more individuals with low self-efficacy to regulate exercise behaviour. The role of exercise and PA interventions and support for people living with CKD may be warranted in order to facilitate exercise behaviours. This was demonstrated in a recent study, which indicated that although a structured exercise intervention did not improve self-reported PA levels, it increased patients’ self-efficacy for PA.

There was variation between ethnic groups regarding the reason they engaged in, or do not participate in PA and exercise. Data from individuals of BA and BC ethnicity indicated cooking and diet to form a large part of their cultural identity. Consensus in focus group discussions identified the need for specialist and culturally sensitive dietary advice, and the need for combined education on both diet and exercise. Weight management was a key driver expressed in this group for engagement in exercise interventions.

Data gathered from the SA focus groups highlighted the strong family bond that exists in this culture. Involvement of family members in healthcare was evident, and the strong family bond that exists in this culture. Involve-

<table>
<thead>
<tr>
<th>Ethnicity</th>
<th>Practical applications</th>
</tr>
</thead>
<tbody>
<tr>
<td>White</td>
<td>► Include a peer and social support element to exercise programmes.</td>
</tr>
<tr>
<td>Black African and black</td>
<td>► Consider providing a variety of exercise options, for example, the opportunity to engage in sports as well as individually.</td>
</tr>
<tr>
<td>Caribbean</td>
<td>► Consider location, access and the environment when setting up renal exercise services.</td>
</tr>
<tr>
<td>South Asian</td>
<td>► Consider the exercise environment, particularly the choice for individuals to be able to exercise away from their home environment in order to dedicate protected time to exercise.</td>
</tr>
<tr>
<td>Non-English speakers</td>
<td>► Provide education to clinicians to increase understanding around cultural perceptions of illness and exercise within the SA population.</td>
</tr>
<tr>
<td>Overall</td>
<td>► Include patient education around the benefits of exercise in CKD in patient consultations.</td>
</tr>
<tr>
<td></td>
<td>► Increase the presence and use of specialist renal physiotherapists within routine clinical care.</td>
</tr>
<tr>
<td></td>
<td>► Clinical appointments should be undertaken initially 1:1 to ensure that individual patient needs are understood and exercise interventions are tailored.</td>
</tr>
</tbody>
</table>

CKD, chronic kidney disease; SA, South Asian.
exercise. Jepson et al.,37 emphasised the centrality of cross-generational family relationships and how this played an important role as to how individuals of SA background socialised.37 These relationships may need to be taken into consideration when developing exercise services or PA interventions for this population.

White males generally associated exercise as something that happened as a social event. Preference for choice of exercise was largely within a group setting, and the importance of peer support was also highlighted to aid motivation. Clarke et al.12 expressed the need for practitioners to consider renal-specific exercise programmes as a potential benefit to aid engagement in PA and exercise for individuals with CKD.12

All interviews and focus groups identified a need for further specialist support and education to provide information with regards to exercise. Challenges with the lack of specialist knowledge of PA and exercise for people with CKD in the community were identified as a barrier, alongside a need for renal physiotherapists as part of the multidisciplinary care delivered during routine outpatient appointments. Table 6 summarises practical applications to engagement in PA and exercise interventions.

Overall, participants felt that exercise within a group setting with other individuals with CKD would provide an environment for peer support, facilitate an increase in confidence and provide accountability with exercise. Participants felt that exercise should foster inclusivity and should be diverse across ethnicity, age and renal modality. This relates to the quantitative data analysis, revealing a significance between self-efficacy to regulate exercise behaviours and self-reported PA levels. This affords a link to a number of psychological theories, including the Social Cognitive Theory (SCT) and the Self-Determinism Theory.38 39 SCT demonstrates that the reproduction of an observed behaviour is influenced by environmental and behavioural factors. This highlights the process through which efficacy beliefs, outcome expectations and goals influence behaviour.36

**Limitations**

Some potential limitations should be considered. This study was conducted in a single centre in South-East London. This area has a diverse ethnic population, and therefore cultural and ethnic views may not be generalisable to the wider CKD population. Only a small number of ethnicities were included in this study. Findings are therefore specific to those individuals and ethnicities included in focus groups and interviews that are relevant to our centre population. The focus groups included individuals from a BC background, and individuals from BA background together within the discussions. This is acknowledged as a limitation.

It is important to acknowledge that the quantitative data are cross-sectional; findings may not be similar if looked longitudinally. This has been explored in previous research which demonstrated a change in self-efficacy but no change in PA levels.38 Future research would benefit from including additional analysis to look at modifiers of relationship between Self-efficacy and PAI scores.

At this NHS Hospital Trust, there is an existing culture within the renal department to promote exercise behaviours. While attempts were made to maintain reflexivity and rigour during the research, prior understanding of rehabilitation held by the researchers may have shaped the research process. To mitigate this, participants were recruited using purposive sampling. Participants with prior engagement in exercise services were limited to minimise sample bias. The main interviewer also used a reflective journal to ensure themes were data driven. Themes were agreed with discussions with an external qualitative researcher.

Lastly, there were a number of participants who were unable to attend the focus group discussions. Attempts were made to create a suitable time and date which would be inclusive for the participants. Due to the challenges on time that patients with CKD face, as well as variability in symptoms day to day it was difficult to accommodate all participants at a single date and time.

**CONCLUSION**

Thematic analyses suggest the understanding, attitudes and beliefs to PA and exercise among different ethnic groups are complex. CKD-specific group exercise interventions may facilitate peer support, improve self-efficacy and health maintenance in individuals with CKD. It is imperative that exercise interventions foster a supportive and inclusive environment, and are designed alongside service users. Raising the profile of specialist renal physiotherapists within the outpatient clinical setting, to provide individualised education and exercise prescription for individuals across the spectrum of CKD may allow for personalised and culturally sensitive exercise and PA education and support. Future research is needed to design exercise interventions that are sensitive to different ethnic groups needs and to evaluate its impact on patient outcomes.

**Acknowledgements** We would like to thank the renal exercise and rehabilitation research team at King’s College Hospital for their support with this study. We also thank Lynda Haggis for her contribution as moderator for the focus group sessions, as well as supporting with the administrative work to bring the focus groups together.

**Contributors** All the authors made significant contributions and were involved in the conception, design, and in drafting the article or revising it critically for important intellectual content; and approved the version of the manuscript submitted (JM, SG, JG, PO, PDH, EC). JM, SG, JG and EC were involved in acquisition of the data, and analysis and interpretation of the data. Manuscript preparation: JM, final version of manuscript and as the guarantor: all. Supervision and mentorship: JM, SG, PO, PDH.

**Funding** This study was funded by Kidney Care UK & BRS Joint Grants Programme 2017. We also thank Kidney Research UK for providing funding for EC through the KRUK AHP PhD fellowship (AHPF_001_20171122). JM is supported through an NIHR HEE/NIHR ICA Pre-doctoral Clinical Academic Fellowship (NIHR301893).

**Competing interests** None declared.

**Patient consent for publication** Not applicable.