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EDITORIAL COMMENT

Living well with kidney disease by patient and care partner empowerment: kidney health for everyone everywhere

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ABSTRACT

Living with chronic kidney disease (CKD) is associated with hardships for patients and their care partners. Empowering patients and their care partners, including family members or friends involved in their care, may help minimize the burden and consequences of CKD-related symptoms to enable life participation. There is a need to broaden the focus on living well with kidney disease and re-engagement in life, including an emphasis on patients being in control. The World Kidney Day (WKD) Joint Steering Committee has declared 2021 the year of 'Living Well with Kidney Disease' in an effort to increase education and awareness on the important goal of patient empowerment and life participation. This calls for the development and implementation of validated patient-reported outcome measures to assess and address areas of life participation in routine care. It could be supported by regulatory agencies as a metric for quality care or to support labeling claims for medicines and devices. Funding agencies could establish targeted calls for research that address the priorities of patients. Patients with kidney disease and their care partners should feel supported to live well through concerted efforts by kidney care communities including during pandemics. In the overall wellness program for kidney disease patients, the need for prevention should be reiterated. Early detection with a prolonged course of wellness despite kidney disease, after effective secondary and tertiary prevention programs, should be promoted. WKD 2021 continues to call for increased awareness of the importance of preventive measures throughout populations, professionals and policymakers, applicable to both developed and developing countries.

Keywords: care partner, health policy, low-middle-income countries, patient empowerment

PATIENT PRIORITIES FOR LIVING WELL: A FOCUS ON LIFE PARTICIPATION

Chronic kidney disease (CKD), its associated symptoms and its treatment, including medications, dietary and fluid restrictions, and kidney replacement therapy, can disrupt and constrain daily living and impair the overall quality of life of patients and their family members. Consequently, this can also impact treatment satisfaction and clinical outcomes [1]. Despite this, the past several decades have seen limited improvement in the quality of life of people with CKD [1]. To advance research, practice and policy, there is increasing recognition of the need to identify and address patient priorities, values and goals [1].

Several regional and global kidney health projects have addressed these important questions including the Standardized Outcomes in Nephrology (SONG) with more than 9000 patients, family members and health professionals from over 70 countries [2, 3]. Across all treatment stages, including CKD, dialysis and transplantation, SONG participating children and adults with CKD consistently gave higher priority to symptoms and life impacts than health professionals [2, 3]. In comparison, health professionals gave higher priority to mortality and hospitalization than patients and family members. The patient-prioritized outcomes are shown in Figure 1. Irrespective of the type of kidney disease or treatment stage, patients wanted to be able to live well, maintain their role and social functioning, protect some semblance of normality, and have a sense of control over their health and wellbeing.

Life participation, defined as the ability to do meaningful activities of life including, but not limited to, work, study, family responsibilities, travel, sport, social and recreational activities, was established a critically important outcome across all treatment stages of CKD [1, 2]. The quotations from patients with kidney disease provided in Box 1 demonstrate how life participation reflects the ability to live well with CKD [4]. According to the World Health Organization (WHO), participation refers to 'involvement in a life situation' [5]. This concept is more specific than the broader construct of

quality of life. Life participation places the life priorities and values of those affected by CKD and their family at the center of decision-making. The World Kidney Day (WKD) Steering Committee calls for the inclusion of life participation, a key focus in the care of patients with CKD, to achieve the ultimate goal of living well with kidney disease. This calls for the development and implementation of validated patient-reported outcome measures, which could be used to assess and address areas of life participation in routine care. Monitoring of life participation could be supported by regulatory agencies as a metric for quality care or to support labeling claims for medicines and devices. Funding agencies could establish targeted calls for research that address the priorities of patients, including life participation.

PATIENT EMPOWERMENT, PARTNERSHIP AND A PARADIGM SHIFT TOWARD A STRENGTHS-BASED APPROACH TO CARE

Patients with CKD and their family members including care partners should be empowered to achieve the health outcomes and life goals that are meaningful and important to them. The WHO defines patient empowerment as 'a process through which people gain greater control over decisions or actions affecting their health', [6] which requires patients to understand their role, to have knowledge to be able to engage with clinicians in shared decision-making, skills and support for self-management. For patients receiving dialysis, understanding the rationale for a lifestyle change, having access to practical assistance and family support promoted patient empowerment, while feeling limited in life participation undermined their sense of empowerment [7].

The WKD Steering Committee advocates for strengthened partnership with patients in the development, implementation and evaluation of interventions for practice and policy settings that enable patients to live well with kidney diseases. This needs to be supported by consistent, accessible and meaningful communication. Meaningful involvement of patients and family members across the entire research process, from priority setting and planning the study through to dissemination and

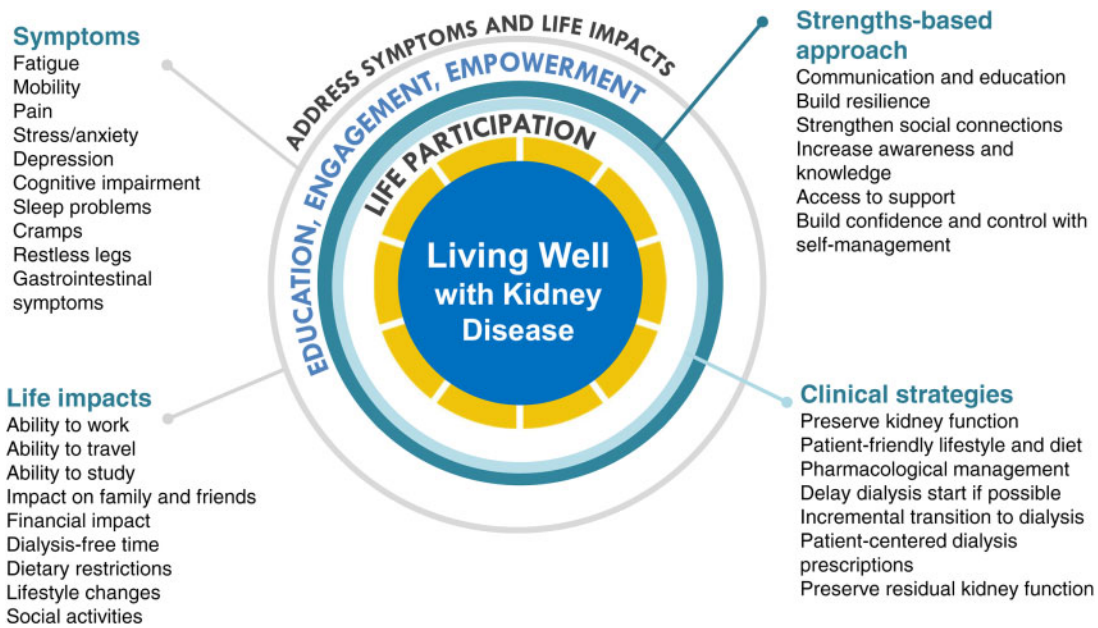


FIGURE 1: Conceptual framework of 'Living Well with Kidney Disease' based on patient centeredness and empowering patient with focus on effective symptom management and life participation.

Box 1. Quotations from patients with CKD related to priorities for living well

'I don't want to think about dying from my disease. I want to be able to live well with my disease.'—Patient with CKD

'Life participation is most important because without it, you can't do anything.'—Child with CKD

'Maybe it's as simple as asking patients whether, how well they are able to participate in the life that they want to lead because it's going to be different for different people.'—Kidney transplant recipient

'Everyone has to face death, what I would like to have is a good quality of life rather than to face death.'—Kidney transplant recipient

'So, it doesn't actually really matter what the numbers say, and some of my numbers should have suggested that I should be feeling a lot worse than what I actually was, it's about how much I feel I can do and participate in my life and feel normal.'—Patient with CKD

'I'm still living. I get out of bed, and I'm still living and still breathing. As long as I can do that, I'm going to carry on and be positive because life is short.'—Patient with CKD

'I put life participation because I know that looking from the outside, I know [his kidney disease] stops [him] from thinking bigger. . . Although that's really big, there's this life that has to happen at the same time.'—Family member

'Amazed at comments from professional (sic) about travel, free time, etc. they seem to think the mechanics of dialysis far more important. Dialysis is a treatment which keeps us alive to live a life, not just to wait for death.'—Patient receiving dialysis

'I prefer to be above ground, then below ground. So why not enjoy life whilst being above ground.'—Adam Martin*

'Over the years, I have learned to worry less, control my emotions, and not fear death. I keep my mind active. I follow the advice of the philosopher-emperor Marcus Aurelius to 'love the hand that fate (has dealt me) and play it as (my) own'. Living well with CKD means to live the best life I can in the time I have available. . . Living well with CKD is the same as living well.'—Tess Harris*

'While CKD brings me some limitations, I can maximize the possibility to live well. I kept working when I was doing hemodialysis. After transplant, I could live: study, work, travel, marry, have children, and service the community.'—Maggie Ng*

*Personal communication; quotations are identified by name with permission.

implementation, is now widely advocated [8]. There have also been efforts, such as the Kidney Health Initiative, to involve patients in the development of drugs and devices to foster innovation [9].

We urge for greater emphasis on a strengths-based approach as outlined in Table 1, which encompasses strategies to support patient resilience, harness social connections, build patient awareness and knowledge, facilitate access to support, and establish confidence and control in self-management. The strengths-based approach is in contrast to

the medical model where chronic disease is traditionally focused on pathology, problems and failures [10]. Instead, the strengths-based approach acknowledges that each individual has strengths and abilities to overcome the problems and challenges faced, and requires collaboration and cultivation of the patient's hopes, aspirations, interests and values. Efforts are needed to ensure that structural biases, discrimination and disparities in the healthcare system also need to be identified, so all patients are given the opportunity to have a voice.

Table 1. Suggested strategies for living well with CKD using a strengths-based approach

Strengths-based approach	Suggested strategies
Build resilience	<ul style="list-style-type: none"> Identify or provide strategies and resources to manage stress and functioning when encountering challenges, adversity and trauma (e.g. commencement of dialysis)
Harness social connections	<ul style="list-style-type: none"> Facilitate connections with other patients to learn coping strategies and for support Support family members/caregivers
Build awareness and knowledge	<ul style="list-style-type: none"> Provide education (including practical advice) on diet and lifestyle modifications Understand, identify and address the potential impacts of CKD (e.g. cognitive function) Encourage patients to ask questions Encourage the use of knowledge to empower and prepare for the future
Facilitate access to support	<ul style="list-style-type: none"> Refer to allied healthcare professionals (e.g. dietitian, social worker, mental health professionals, occupation therapists) Provide support that enables the patient to participate in important life activities, e.g. work
Establish confidence and control in self-management	<ul style="list-style-type: none"> Support informed and shared decision-making (including dialysis, kidney transplantation, conservative or nondialytic care) Encourage patients to learn to 'get in tune' with what works well for them and to voice any concerns, and work together to develop better management strategies to enable patients to feel better Provide strategies to prevent or manage complications (e.g. infection) Support open communication regarding goals, concerns and priorities

THE ROLE OF CARE PARTNER

A care partner is often an informal caregiver who is also a family member of the patient with CKD [11]. They may take on a wide range of responsibilities including coordinating care (including transportation to appointments), administration of treatment including medications, home dialysis assistance and supporting dietary management. Caregivers of patients with CKD have reported depression, fatigue, isolation and also burnout. The role of the care partner has increasingly become more important in CKD care given the heightened complexity in communicative and therapeutic options including the expansion of telemedicine under the coronavirus disease 2019 (COVID-19) pandemic and given the goal to achieve higher life expectancy with CKD [12]. The experience of caring for a partially incapacitated family member with progressive CKD can represent a substantial burden on the care partner and may impact family dynamics. Not infrequently, the career goals and other occupational and leisure aspects of the life of the care partner are affected because of CKD care partnership, leading to care partner overload and burnout. Hence, the abovementioned principles of life participation need to equally apply to care partners as well as all family members and friends involved in CKD care.

LIVING WITH KIDNEY DISEASE IN LOW-INCOME REGIONS

In low- and lower-middle-income countries (LICs and LMICs) including in sub-Saharan Africa, South East Asia and Latin America, patients' ability to self-manage or cope with the chronic disease vary but may often be influenced by internal factors including spirituality, belief system and religiosity, and external factors including appropriate knowledge of the disease, poverty, family support system and one's grit and social relations network. The support system comprising healthcare

providers and caregivers plays a crucial role as most patients rely on them in making decisions, and for the necessary adjustments in their health behavior [13]. In LIC regions, where there are often a relatively low number of physicians and even lower number of kidney care providers per population especially in rural areas, a stepwise approach can involve local and national stakeholders including both nongovernmental organizations and government agencies by (i) extending kidney patient education in rural areas, (ii) adapting telehealth technologies if feasible to educate patients and train local community kidney care providers and (iii) implementing effective retention strategies for rural kidney health providers including adapting career plans and competitive incentives.

Many patients in low-resource settings present in very late stage, needing to commence emergency dialysis [14]. The very few fortunate ones to receive kidney transplantation may acquire an indescribable chance at normal life again, notwithstanding the high costs of immunosuppressive medications in some countries. For some patients and care partners in low-income regions, spirituality and religiosity may engender hope: when ill they are energized by the anticipation of restored health and spiritual well-being. For many patients, informing them of a diagnosis of kidney disease is a harrowing experience both for the patient (and caregivers) and the healthcare professional. Most patients present to kidney physicians (usually known as 'renal physicians' in many of these countries) with trepidation and apprehension. It is rewarding therefore to see the patient's anxiety dissipate after reassuring him or her of a diagnosis of simple kidney cysts, urinary tract infection, simple kidney stones, solitary kidneys, etc., that would not require extreme measures like kidney replacement therapy. Patients diagnosed with glomerulonephritis who have an appropriate characterization of their disease from kidney biopsies and histology, who receive appropriate therapies and achieve remission, are relieved and are very grateful. Patients are glad to

discontinue dialysis following resolution of acute kidney injury or acute on CKD.

Many CKD patients who have residual kidney function appreciate being maintained in a relatively healthy state with conservative measures, without dialysis. They experience renewed energy when their anemia is promptly corrected using erythropoiesis-stimulating agents. They are happy when their peripheral edema resolves with treatment. For those on maintenance hemodialysis who had woeful stories from emergency femoral cannulations, they appreciate the construction of good temporary or permanent vascular accesses. Many patients in low-resource settings present in very late stage needing to commence emergency dialysis. Patients remain grateful for waking from a uremic coma or recovering from recurrent seizures when they commence dialysis.

WORLD KIDNEY DAY 2021 ADVOCACY

WKD 2021 theme of 'Living Well with Kidney Disease' is deliberately chosen to have the goals to redirect more focus on plans and actions toward achieving patient-centered wellness. 'Kidney Health for Everyone, Everywhere' with emphasis on patient-centered wellness should be a policy imperative that can be successfully achieved if policymakers, nephrologists, healthcare professionals, patients and care partners place this within the context of comprehensive care. The requirement of patient engagement is needed. In 2016 the WHO put out an important document on patient empowerment (WHO 2016): 'Patient engagement is increasingly recognized as an integral part of health care and a critical component of safe people-centred services. Engaged patients are better able to make informed decisions about their care options. In addition, resources may be better used if they are aligned with patients' priorities and this is critical for the sustainability of health systems worldwide. Patient engagement may also promote mutual accountability and understanding between patients and health care providers. Informed patients are more likely to feel confident to report both positive and negative experiences and have increased concordance with mutually agreed care management plans. This not only improves health outcomes but also advances learning and improvement while reducing adverse events.' In the International Society of Nephrology (ISN) Community Film Event at World Congress of Nephrology 20 (ISN Community Film Event 2020), it is good to see a quote in the film from patients: 'Tell me. I will forget; Show me. I will remember; Involve me. I will understand.' ISN Global Kidney Policy Forum 2019 included a patient speaker Nicki Scholes-Robertson from New Zealand: 'Culturally appropriate and sensitive patient information and care are being undertaken in New Zealand to fight inequities in kidney health, especially in Maori and other disadvantaged communities.'

WKD 2021 would like to promote to the policymakers increasing focus and resources on both drug and non-drug programs in improving patient wellness. Examples include funding for erythropoiesis-stimulating agents and anti-pruritic agents for managing anemia and itchiness, respectively, to name but a few [15, 16]. Home dialysis therapies have been consistently found to improve patient autonomy and flexibility, and quality of life in a cost-effective manner, enhancing life participation. Promoting home dialysis therapies should tie in with appropriate 'assisted dialysis' programs to reduce patient and care partner fatigue and burnout. Also, examples like self-management programs, cognitive behavioral therapy and group therapies for managing depression, anxiety and insomnia should be

promoted before resorting to medications [17]. The principle of equity recognizes that different people with different levels of disadvantage require different approaches and resources to achieve equitable health outcomes. The kidney community should push for adapted care guidelines for vulnerable and disadvantaged populations. The involvement of primary care and general physicians, especially in LICs and LMICs, would be useful in improving the affordability and access to services through the public sector in helping the symptom management of CKD patients and improving their wellness. In the overall wellness program for kidney disease patients, the need for prevention should be reiterated. Early detection with a prolonged course of wellness despite kidney disease, after an effective secondary prevention program, should be promoted [18]. Prevention of CKD progression can be attempted by lifestyle and diet modifications such as a plant-dominant low-protein diet and by means of effective pharmacotherapy including administration of sodium-glucose co-transporter 2 inhibitors [19]. WKD 2021 continues to call for increased awareness of the importance of preventive measures throughout populations, professionals and policymakers, applicable to both developed and developing countries [18].

CONCLUSIONS

Effective strategies to empower patients and their care partners strive to pursue the overarching goal of minimizing the burden of CKD-related symptoms in order to enhance patient satisfaction, health-related quality of life and life participation. The WKD 2021 theme of 'Living Well with Kidney Disease' is deliberately chosen to have the goals to redirect more focus on plans and actions toward achieving patient-centered wellness. Notwithstanding the COVID-19 pandemic that has overshadowed many activities in 2020 and beyond, the WKD Steering Committee has declared 2021 the year of 'Living Well with Kidney Disease' in an effort to increase education and awareness on the important goal of effective symptom management and patient empowerment. Whereas the WKD continues to emphasize the importance of effective measures to prevent kidney disease and its progression [18], patients with preexisting kidney disease and their care partners should feel supported to live well through concerted efforts by kidney care communities and other stakeholders throughout the world, even during a world-shattering pandemic such as COVID-19 that may drain many resources [20]. Living well with kidney disease is an uncompromisable goal of all kidney foundations, patient groups and professional societies alike, to which the ISN and the International Federation of Kidney Foundation World Kidney Alliance are committed at all times.

CONFLICT OF INTEREST STATEMENT

K.K.-Z. reports honoraria from Abbott, Abbvie, ACI Clinical, Akebia, Alexion, Amgen, Ardelyx, AstraZeneca, Aveo, BBraun, Cara Therapeutics, Chugai, Cytokinetics, Daiichi, DaVita, Fresenius, Genentech, Haymarket Media, Hospira, Kabi, Keryx, Kissei, Novartis, Pfizer, Regulus, Relypsa, Resverlogix, Dr Schaefer, Sandoz, Sanofi, Shire, Vifor, UpToDate and ZS-Pharma. P.K.-T.L. reports personal fees from Fibrogen and AstraZeneca. G.S. reports personal fees from Multicare, Novartis, Sandoz and AstraZeneca. V.L. reports nonfinancial support from Genesis Pharma.

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