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A critical review of a hidden epidemic: examining the occupational and environmental risk factors of chronic kidney disease of unknown etiology (CKDu)

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The global burden of chronic kidney disease (CKD) in terms of mortality and disability-adjusted life years has increased, and this trend is expected to worsen over the next few decades. The primary cause of CKD is known to be due to hypertension and diabetes, however, over the last three decades, a form of CKD has been described in people without any known risk factors. These cases can be described as chronic kidney disease of an unknown etiology (CKDu). Cases of CKDu are rising primarily among rural agricultural communities in affected regions and occur mostly among young male farmers. There is no agreement on whether CKDu in these emerging clusters represents a single disease or a group of different diseases. As such, hypothesized causes of CKDu development include chronic occupational heat stress and dehydration, as well as exposure to environmental contaminants and agrochemicals, such as heavy metals and pesticides. The purpose of this critical review is to bring together the current literature on proposed CKDu etiologies, specifically those related to work in the agricultural sector. This review examines what is known about these occupational and environmental factors and their potential impact on the widespread epidemics of CKDu.

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Environmental significance

This review examines the hidden epidemic of chronic kidney disease of unknown etiology (CKDu), a severe form of chronic kidney disease unrelated to traditional risk factors emerging in clusters predominantly in low- to middle-income countries. There is no agreement on whether CKDu within these clusters represents a single disease or a group of different diseases. Hypothesized causes of CKDu include chronic occupational heat stress and dehydration, as well as exposure to environmental contaminants and agrochemicals. This review examines what is known about these environmental factors and their potential impact on the widespread epidemics of CKDu.

1. Introduction

Chronic kidney disease (CKD) is a potentially life-threatening condition that requires prompt and sustained medical attention.¹ It has become a major public health issue in recent decades, as awareness and recognition of the condition's high burden of disease continues to grow.² In 1990, CKD was ranked as the 17th leading cause of death globally. In 2017 this rose to 12th, indicating that both prevalence and burden of disease are

growing.² Estimates from 2017 found CKD to be responsible for up to 1.2 million deaths worldwide, and this number is projected to reach between 2.2 and 4.0 million deaths by 2040.^{2,3} While chronic kidney disease is commonly related to known risk factors like diabetes, hypertension, and glomerulonephritis,⁴ an epidemic of new CKD cases of unknown cause (CKDu) has been growing across the globe.⁵⁻⁹ In these cases, a patient has none of the known risk factors thought to cause CKD, and the exact cause remains unidentified.^{6,9,10}

The total extent of CKDu worldwide is still unknown due to a lack of quality data from many low-income countries.^{11,12} The first descriptions of CKDu came from Sri Lanka in the 1990s and Costa Rica in the 2000s, when disease hot spots started to appear, but models suggest CKDu in Costa Rica could be traced back to the 1970s.^{13,14} CKDu hotspots, or areas of high disease prevalence, include Central America and South Asia, chiefly Sri Lanka, India, El Salvador, Costa Rica, and Nicaragua.¹⁵⁻¹⁷ Cases there are being recognized particularly in rural areas and regions of agricultural activity.^{5,13} CKDu has also been referred

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to as chronic kidney disease of a nontraditional etiology (CKDnt), mesoamerican nephropathy (MEN), and chronic interstitial nephritis of agricultural communities (CINAC).^{13,18} This variable terminology coupled with varying case definitions may have hindered broader understanding of the condition and its global burden until recent years due to limited literature comparability and data sharing.^{11,13}

With little global acknowledgement of CKDu and varying terminology, tracking and treatment of the condition in certain regions of the world has been a challenge, and much is still unknown about the disease.^{2,19} Many hypotheses have been put forth suggesting potential causes of these unknown cases of CKD, but a definitive etiology is yet to be identified.^{13,20}

The purpose of this critical review is to bring together the current literature on proposed CKDu etiologies, specifically those related to occupational and environmental risk factors. The review followed a structured approach and was conducted by searching the databases PubMed, Scopus and Google Scholar using key words, including “CKDu”, “etiology” “environmental factors”, and “agriculture”. Over 150 publications were reviewed, with 115 cited here. Content reviewed was limited to that published in the last 30 years.

2. Chronic kidney disease

It is estimated that around 10% of the global population is currently affected by CKD, with many unaware of their condition.²¹ In early stages of the disease patients often present with no symptoms,²² but as CKD progresses, scarring and nephron loss can occur, causing the kidneys to gradually lose their ability to function.²³ Further progression can lead to end-stage renal disease (ESRD), which is associated with kidney failure leading to heavy disease burden and premature death.⁴

Recognition of CKD has grown over recent years, as increased awareness of the disease in the research community led to the development of a more specific classification for diagnosis.²⁴ In 2013, a joint international effort involving the U.S. Centers for Disease Control and Prevention (CDC), the Pan American Health Organization, and the Latin American Society of Nephrology and Hypertension led to the development of a set of clinical guidelines for defining and diagnosing CKD. These guidelines have been laid out by an international body that provides recommendations on the evaluation and management of kidney diseases called Kidney Disease: Improving Global Outcomes, or KDIGO.^{25,26} According to the internationally accepted definition, chronic kidney disease is present when “there are abnormalities in the function or structure of the kidney that are present for greater than three months, with implications for health”.²⁵

2.1. Diagnosing CKD

The kidneys serve a variety of purposes in the body. These include key endocrine and metabolic roles but, most importantly, the primary function of the kidneys is to maintain normal internal homeostasis – such as normal electrolyte, fluid, and acid–base balance, and the excretion of water-soluble waste

products from the body.²⁵ Nephrons, the functional units of the kidneys, remove excessive levels of solutes like electrolytes, which prevents blood from becoming toxic.²⁷ Additionally, the kidneys produce important hormones like renin and erythropoietin, which keep the human body functioning properly.²⁴ When the kidneys are not working efficiently, individuals can develop symptoms like weight loss, poor appetite, infections, and impairments in physical and cognitive functioning.^{22,27}

Diagnosing CKD can be a challenge, however, as there is not always a clear indicator of disease. Unlike many other diseases, CKD is often asymptomatic until it is very advanced.^{2,4} The best estimates of kidney function can be determined by testing an individual's glomerular filtration rate and albuminuria levels.^{4,25} The glomerular filtration rate measures the volume of blood that is filtered by the glomeruli, tiny filters in the kidneys, each minute, whereas a test of the albuminuria level measures the amount of the protein albumin that is present in the urine.^{28,29} When the glomerular filtration rate is low and there is evidence of increased albuminuria, the individual is suffering from some level of kidney disease.^{4,25}

After being diagnosed, CKD is classified by cause and then further staged based on rates of eGFR and albuminuria (Fig. 1). These levels are important in determining the risks an individual may have after developing CKD and can help to determine the severity of the disease.^{4,30} To progress between each category a patient must show prolonged elevated albuminuria levels and continual decreases in the estimated glomerular filtration rate (eGFR).^{4,17} Patients can progress rapidly through the stages to kidney failure within months, or it can take years of follow up before a patient will become severely ill with the disease.^{25,26}

2.2. Known CKD risk factors

The most common risk factors for CKD include diabetes and hypertension.²⁴ Of kidney failure cases in the U.S. between 2015 and 2017, 75% had primary diagnoses of diabetes or hypertension.^{31,32} However, factors like age, race, and socioeconomic status are also thought to play a role in determining the risk of developing CKD.² Large scale screening programs implemented in the U.S., Australia, and Norway in the 2000s revealed that over 10% of the population had markers of disease.² It is important to note that despite high prevalence of the disease in the general population, it is not recommended to screen individuals for CKD without a history of known risk factors or symptoms.²⁴ However, most cases of chronic kidney disease begin asymptotically,²² and without a thorough understanding of all of the risk factors that could cause CKD, many CKD patients remain unaware of their condition.²

2.3. CKD treatment and associated costs

In both developed and developing countries, it is estimated that fewer than 10% of patients with CKD know that they have the disease.³³ This results in medical resources being primarily geared towards providing treatments for ESRD and not the earlier stages of CKD. While there is currently no cure for chronic kidney disease, available treatments can prevent the



Prognosis of CKD by GFR and Albuminuria Categories: KDIGO 2012

				Persistent albuminuria categories		
				Description and range		
				A1	A2	A3
				Normal to mildly increased	Moderately increased	Severely increased
				<30 mg/g <3 mg/mmol	30-300 mg/g 3-30 mg/mmol	>300 mg/g >30 mg/mmol
GFR categories (ml/min/1.73 m ²) Description and range	G1	Normal or high	≥90			
	G2	Mildly decreased	60-89			
	G3a	Mildly to moderately decreased	45-59			
	G3b	Moderately to severely decreased	30-44			
	G4	Severely decreased	15-29			
	G5	Kidney failure	<15			

Green: low risk (if no other markers of kidney disease, no CKD); Yellow: moderately increased risk; Orange: high risk; Red, very high risk.

Fig. 1 Prognosis of CKD by eGFR and albuminuria categories, determining level of risk for disease. Reprinted from KDIGO 2012 Clinical Practice Guideline for the Evaluation and Management of Chronic Kidney Disease, Volume 3, Issue 1, Kidney International Supplements, Chapter 1: definition and classification of CKD, 6, 2013, with permission from Elsevier.

disease from progressing to a more advanced stage of CKD.^{2,34} The best approach to managing CKD is early detection so that measures can be put in place to limit disease progression and reduce cardiovascular disease, before the disease has time to greatly impair kidney functions or cause kidney failure.³⁵ Things such as dietary changes, exercise, medications, and treatment of known risk factors (*e.g.* diabetes and/or hypertension) are the primary approaches for CKD treatment by medical professionals.^{25,36} These interventions can improve outcomes for both renal and cardiovascular functions, slowing or preventing the progression to ESRD.²

The treatment costs of CKD started rising after the 1960s when renal replacement techniques (RRT) became available.^{2,36} If CKD progresses and kidney failure occurs, patients will need RRT in the form of regular dialysis or kidney transplants, both of which are costly and not easily obtained in most low- and middle-income countries around the world.^{12,36} This is also the case in high income countries, where the high cost of long-term dialysis is a barrier to many people receiving treatment.³⁷ Estimates show that around 2.5 million patients worldwide currently undergo dialysis or await kidney transplants, and this need is expected to double to 5.4 million by 2030.² While large, this number only represents around 10% of those who currently

require treatment to live, leaving 90% of those with kidney failure without access to necessary lifesaving treatment.²⁴

CKD and its projected growth represent a major public health and economic challenge worldwide. Though many people are not treated, the availability of renal replacement therapies is increasing, and as more people live longer with the disease, the total disease burden of CKD grows.^{2,36} Compared to worldwide rates, the U.S. and Mexico were found to have some of the highest CKD burdens, which can be largely attributed to similarly high rates of diabetes and hypertension.^{38,39} In the U.S., treatment for CKD is thought to exceed 46 billion U.S. dollars each year.^{24,36} Additionally, those with CKD may also develop comorbidities such as cardiovascular disease, anemia, bone disorders, and diseases of the nervous system, which can add to healthcare costs.²⁴

3. Chronic kidney disease of an unknown etiology

Over the last few decades, an increase in cases of CKD and CKD mortality has been observed, particularly in rural areas of Central America and South Asia.^{40,41} As previously discussed,



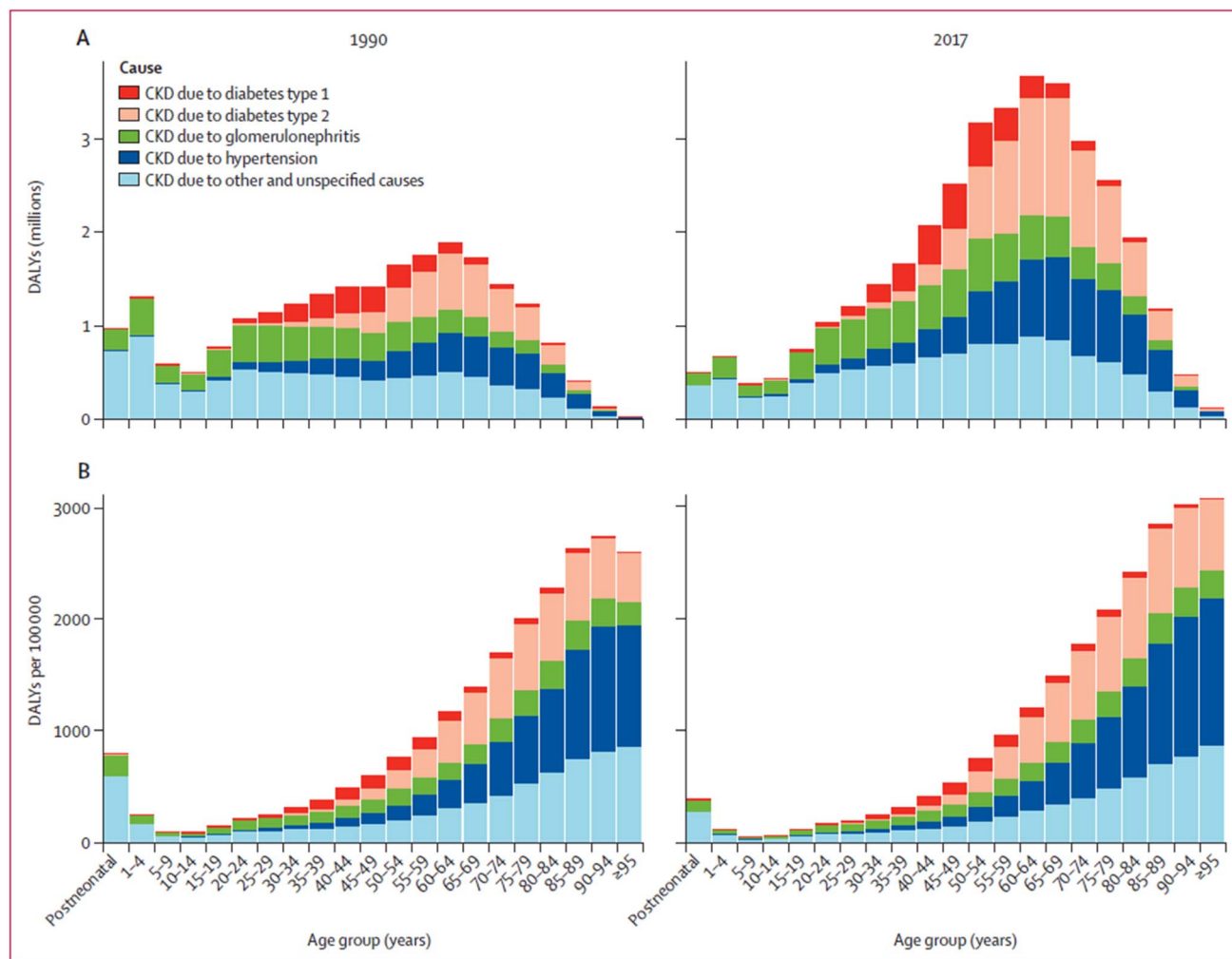


Fig. 2 Number (A) and rate (B) of global disability-adjusted life years (DALYs) for chronic kidney disease (CKD) based on underlying cause in 1990 and 2017. Used with permission of Elsevier Science & Technology Journals, from Global, regional, and national burden of chronic kidney disease, 1990–2017: a systematic analysis for the Global Burden of Disease Study 2017. Bikbov B., Purcell C. A., Levey A. S., *et al.*, *Lancet*, 2020, **395**, 10225; permission conveyed through Copyright Clearance Center, Inc.

many cases of CKD are related to risk factors such as diabetes, hypertension, and aging. However, some cases have an unknown origin, and these instances can be described as cases of chronic kidney disease of unknown etiology, or CKDu. Regional clustering of this disease suggests CKDu is an environmentally induced problem.⁴²

Bikbov *et al.*,² shows the distribution of disability-adjusted life years (or DALYs) of CKD in 1990 and 2017 based on the underlying cause (Fig. 2). DALYs represent the sum of the years of potential life that is lost due to premature mortality or the years of life lost due to a disability, and it is a valuable metric because it exemplifies the overall burden of a disease, not just the case load.² While the largest contribution of CKD DALYs in 2017 was due to traditional risk factors, such as diabetes, a substantial number of DALYs across all age groups were deemed to be caused by 'other and unspecified causes'. These cases were estimated to have the highest age standardized rate of DALYs across all causes of CKD in 2017 at 138.8 per 100 000 in the population.²

CKDu is a global disease affecting low-, middle-, and high-income countries, highlighting the urgency in understanding its risk factors and causes.⁴³ The global burden of CKD and CKDu is increasing faster in low- and middle-income countries, and even within economically developing nations, those affected by CKDu are usually from the most underserved populations, including in the U.S. among immigrant agricultural workers^{38,44} and Aboriginal groups in central Australia and New Zealand.^{11,45} CKDu is a diagnosis of exclusion and characteristics of the disease may vary in different geographic locations.^{43,46} To date, there is no conclusive evidence linking CKDu to any single etiological cause despite consistent association across regions with male sex, rural residence, and farm labor.¹¹

4. Proposed etiologies

While many etiologies have been proposed for CKDu, none have been definitively recognized in any region of the world.⁴⁷ Regional epidemics tend to have common demographics, disease presentation, clinical features, and renal histopathology



(i.e., kidney tissue effects).⁴⁸ However, study designs and outcome measurements differ between studies significantly so that it is difficult to identify underlying etiology or if the clusters of diseases across different geographical locations are a single or a group of different diseases.¹⁵ Studies have noted clinical differences between CKD and CKDu, though there is no agreement on a clinical or histopathological feature that can be diagnostic in CKDu.¹ In CKD caused by hypertension and diabetes, the glomeruli are predominantly affected, thus proteinuria (presence of protein in urine) is an early sign of disease.⁴⁶ Unlike common CKD, CKDu is generally recognized as a disease process affecting the tubulo-interstitial compartment of the kidney, and proteinuria is considered inconsistent with a diagnosis of CKDu, making this commonly used marker ineffective for diagnosis or screening.^{46,49} The Disadvantaged Population estimated Glomerular Filtration Rate (eGFR) Epidemiology Study (DEGREE) definition of CKDu even excludes cases with heavy proteinuria.⁵⁰ In fact, one proposed definition of CKDu is kidney damage with an eGFR of $<60 \text{ mL min}^{-1}/1.73 \text{ m}^2$ in the absence of diabetes, hypertension, heavy proteinuria, and structural renal disease.^{51–53} Nevertheless, proteinuria continues to be used as an early indicator of disease by the Department of Health in Sri Lanka and all other CKDu-affected countries.⁴⁸

Hotspots of CKDu have mostly been reported in agricultural communities in Central America and South Asia, and suspected in locations such as Africa, the Middle East, and North America. Clusters of CKDu have been described in predominantly rural, agricultural communities and appear to affect lower socioeconomic classes that work in farm labor in tropical climates.⁵⁴ CKDu is characterized as slowly progressing and asymptomatic until advanced stages of the sickness, which is described as an incurable and slowly deteriorating disorder.⁵⁵ This can lead to high mortality rates and disease burden, especially in CKDu hotspots. Major hotspots of CKDu disease prevalence include Central American countries such as Nicaragua, El Salvador, Costa Rica, and Mexico, as well as South Asia including India and Sri Lanka.^{13,38} Because there is evidence from some histopathological studies that CKDu patterns are similar (if not the same) across regional epidemics, identification of risk factors would be impactful not only regionally but also globally in understanding chronic kidney disease.

Based on current evidence, there is a clear association between the distribution of CKDu cases and rural and agricultural communities.⁵⁶ Even though some of those who are affected are not agricultural workers, a majority live in communities close to some form of agriculture. This suggests some condition or combination of conditions related to rural agricultural settings is a possible factor contributing to widespread CKDu. Proposed causation typically begins with occupational, social, or environmental exposures, which are then tied directly to secondary factors like heat or exposure to pesticides, heavy metals, or contaminated food.⁵⁷ Based on this premise and by simplifying the web of causation from Gunatilake *et al.*,⁵⁷ three main subgroups of etiologies have been identified that are the most salient in the discussion of the origins of CKDu. These include heat stress, drinking water contamination, and agrochemical exposures. Below, the

relevant literature is synthesized surrounding these three etiologies. Other proposed factors not discussed in this review include genetics, ethnicity, alcoholism, altitude, the use of nonsteroidal drugs, and more.²⁵ While some literature has found associations between these factors and chronic kidney disease of unknown origin, evidence is generally limited.¹ Ultimately, more research is needed to determine whether any are likely drivers of disease.

4.1. Heat stress nephropathy

A significant body of research published on the topic has suggested that heat stress neuropathy due to repeated dehydration and strenuous manual labor is the main risk factor for the development of CKDu.^{58–60} The association between occupational heat exposure with kidney injury has been mainly reported in tropical regions. A combination of heat stress, water shortage, and overexertion can initiate several physiologic processes that lead to kidney injury, and repeated kidney injury may subsequently lead to CKD.⁵⁹ Heat stress occurs when the body cannot dissipate excess heat, leading to a rise in core temperature and increased heart rate.⁶¹ This can happen due to a combination of factors such as high air temperature, humidity, physical exertion, and clothing.⁶¹ When the body absorbs more heat than it can release, it can result in various heat-related illnesses like heat stroke, exhaustion, cramps, and rashes.⁶¹

Experts suggest that CKDu may be a form of heat-stress nephropathy and “may represent one of the first epidemics due to global warming”.⁵⁹ As accelerating climate change brings increased heat extremes, there is an urgent need to identify its impact on climate-related health crises.^{59,62,63} The area of climate and health should be a focus in future research of CKDu to identify, and subsequently prevent and treat, its cause. Future research should also focus on investigation of environmental contaminants and co-occurrence impacts on CKDu.

Central America has been the most studied region surrounding heat stress nephropathy. Researchers have shown that in Costa Rica, CKDu was probably present as far back as the 1970's in the Guanacaste Province on the Pacific Coast.^{16,64} By 2010, the prevalence of CKDu in this area had grown ten-fold in men and four-fold in women.^{16,64} While the study could not definitively attribute the increase in cases to rising temperatures, it noted heat stress and dehydration as possible stressors.¹⁶ Another study suggested a possible link between increase in cases in Costa Rica with climate change.⁵⁹ Over that time (1970s to 2010), maximum temperatures in Central America rose by 0.2 degrees Celsius, with the number of extremely hot days increasing by 30–75%.⁵⁹

For this region, Fig. 3 illustrates that the higher average maximum temperatures fall along the Pacific Coast, the main region of the epidemic in Central America. However, suggesting that a warming climate correlates with a rise in CKDu cases does not necessarily prove that a link exists, and fails to account for other factors that could be at play. CKDu is thought to be most prevalent among sugar cane workers in Central America, occurring more commonly in men,⁶⁵ and thus it could be



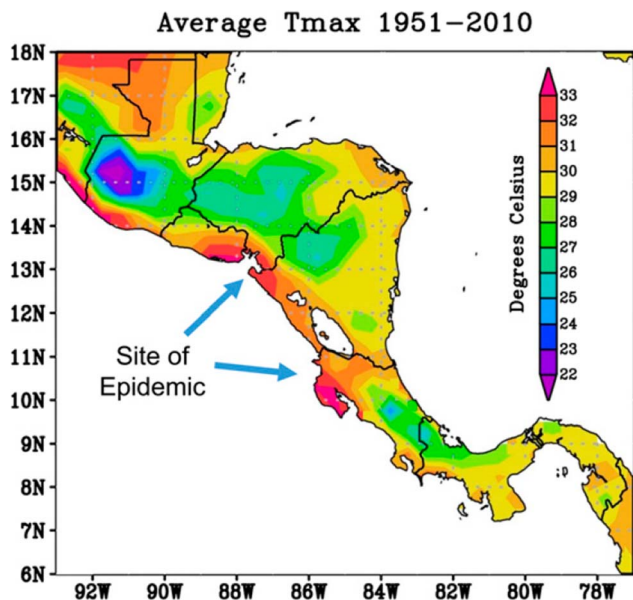


Fig. 3 Average maximum temperatures (T_{max}) in Central America from 1951 to 2010. The higher temperature areas correspond closely with major sites of the CKDu epidemic in this region. Used with permission of American Society of Nephrology, from Climate change and the emergent epidemic of CKD from heat stress in rural communities: the case for heat stress nephropathy. Glaser J., *Clinical Journal of the American Society of Nephrology*, 2016, 11, 8; permission conveyed through Copyright Clearance Center, Inc.

related to occupational heat stress. Nevertheless, women and men who are not sugarcane workers are still suffering from CKDu in Central America,⁵⁶ indicating that there may be co-existent risk factors (e.g., agrochemical exposure or contaminated drinking water). More research targeted to these epidemic sites is recommended to determine possible causation more definitively, and additional contributing risk factors, between heat stress and kidney disease.

The relationship between CKDu and heat has also been found in other studies with specific groups of agricultural workers. In a group of sugarcane workers with normal preharvest kidney function, researchers found that the development of acute kidney injury (AKI) during the harvest season was a common occurrence, developing in 34 of 326 workers.⁶⁵ Of the workers who had developed work-related AKI and participated in at least one follow up visit at 6 and 12 months, 11 experienced a >30% decrease in eGFR, an indication of CKD, while 18 stayed the same or improved.⁶⁵ Researchers attributed the frequency with which CKDu was found in the sugarcane workers to their work in such extreme heat conditions, but of the 326 study participants, all were males who worked in sugar cane fields, and other occupational or socioeconomic factors were not ruled out.⁶⁵ A more recent study published in 2023 focused on a cohort of 569 outdoor workers in El Salvador and Nicaragua and examined how occupation related to heat stress and heat strain.⁶⁶ Of the five industries represented among the cohort (sugarcane, corn, plantain, brickmaking, and construction workers), sugarcane workers and Nicaraguan agrochemical

applicators performed more strenuous work and experienced greater levels of heat strain and subsequent impairment of kidney function (defined by $eGFR < 90 \text{ mL min}^{-1}/1.73 \text{ m}^2$).⁶⁶ Another review analyzed longitudinal data from more than 800 sugarcane cutters and found inflammation biomarkers and fever were associated with kidney injury.⁶⁷

Although few studies have examined the impact of heat stress and strain on kidney function in populations in the U.S., available evidence suggests the presence of CKDu hotspots in North America.^{38,68} In a sample of 471 agriculture workers across 29 farms in California, heavy occupational workload and piece-rate work (i.e., compensation per task completed) were associated with AKI.⁶⁹ Additionally, evidence of AKI across a single work shift was found in 14.9% of participants in this group and increased workload was associated with a nearly two-fold increase in the odds of developing AKI at work.⁶⁹ A study of 192 agricultural workers in Florida found that for each 5 degree (Fahrenheit) increase in the heat index, the odds of AKI increased by 47%.⁶⁸

Recently, researchers have warned against accepting the heat stress neuropathy hypothesis as the leading cause of CKDu because it could be detrimental in the effort to search for other possible etiological factors, such as exposure to environmental toxins.^{46,56,70,71} One research team found that high temperatures did not have any significant impact on CKDu development in El Salvador,⁷² and another concluded that the trends in CKDu in Central America cannot be fully explained by the heat stress hypothesis.⁷³ Herath *et al.*,⁵⁶ based on a review of heat-related literature, found inadequate evidence to support a causal relationship between heat stress and CKDu. They argue that CKDu is not observed in other tropical regions of the world where workers are exposed to heat stress, and that the existence of cases of CKDu in populations that have not been exposed to heat stress would lead to the assumption that other factors are more likely to be at play.^{46,56,70,71,74} A systematic review by Chapman *et al.*, found no consistent evidence that supports an association between CKD and heat-stress or dehydration, though they did find statistically significant associations for the use of pesticides.⁷⁵ We note, as also suggested by Mendley *et al.*,¹³ cases in these other regions of the world may simply remain unreported due to a lack of quality data.

An intervention program study conducted in El Salvador and Nicaragua showed that the implementation of so called water-rest-shade programs was able to reduce the incidence of kidney damage in workers that were high risk.⁷⁶ The program, an intervention adapted from the U.S. Occupational Health and Safety Administration (OSHA) was conducted two months into a five month harvest period. Each worker was provided with a camelback for water and stations for water refills, as well as nearby shade tents and mandatory 10 to 15 minute rest periods every 1 to 1.5 hours. Results showed that those who participated in the intervention program had less of a decrease in their eGFR levels than the control group. This was observed both across a workday and across the entire harvest, though sample size was limited.⁷⁶ While this association does not prove causality for developing the disease, the finding that rest and shade (i.e., less heat stress) leads to a reduction in kidney damage is important



and should be explored further. It also represents a promising, relatively easy to implement best practice for protecting the kidney health of workers in occupations prone to heat stress.

4.2. Drinking water source and quality

The characteristic, unique geographical distribution of CKDu cases suggests the possible involvement of exposure to contaminated drinking water.⁷⁷ Consistent exposures to contaminated drinking water have been found to increase risks of developing cancer and disorders of the kidney, liver, and reproductive organs.⁷⁸ A lack of quality drinking water has also been linked to cases of CKDu.^{14,79–83} Thus, it is an important factor to consider in furthering our understanding of possible etiologies.

A number of studies have postulated a role for groundwater quality in disease incidence and progression, suggesting it is a significant contributor to the etiology of CKDu.^{9,84,85} In Sri Lanka, CKDu is more prominent among communities that consume groundwater as their main source of drinking water, with more than 87% of the population living in CKDu-endemic regions relying on wells.⁸⁰ Additionally, previous research has shown that people who drink well water are more vulnerable to the disease compared to those who consume water from public water supplies, natural springs, rainwater, and surface waters.⁹ In these same CKDu-endemic regions, very low (1.5%) or zero prevalence of CKDu was reported from those who consumed spring water.^{9,86} Data has also shown that CKDu patients tend to cluster toward the lower elevations of local tanks and canal systems, further implying a hydro-geochemical relationship.^{6,86,87} Recently, one study in Central India found CKDu cases had significantly higher surface water usage as their drinking water source (OR 3.178, 95% CI 1.029–9.818).⁸⁸

The most studied environmentally induced causative factor for CKDu is high fluoride (F^-) concentration in drinking water.⁸⁵ Fluorine is a naturally abundant chemical element, and due to water–rock interactions within aquifers fluoride is commonly present in groundwater. Kidneys are one of the most important organs that remove fluoride from the body, with about 60% of the daily fluoride that is absorbed by healthy adults ultimately excreted in urine.⁸⁹ Therefore, the kidney is one of the most exposed soft tissues to fluoride in the body.⁸⁹ A 2006 report by the U.S. National Research Council stressed the dangers of fluoride on renal tissue, stating that “human kidneys concentrate fluoride as much as 50-fold from plasma to urine. Portions of the renal system may therefore be at higher risk of fluoride toxicity than most soft tissues”.⁹⁰ Some of the major adverse effects on kidneys attributed to excessive fluoride consumption include inhibition in tubular reabsorption, changes in urinary ion excretion, and disruption of collagen biosynthesis in the body.^{91,92}

Due to the health risks associated with long-term exposure to high fluoride concentrations, the World Health Organization (WHO) recommends a maximum acceptable limit of 1.5 mg L^{-1} of fluoride in drinking water with an optimal range between 0.5 and 1.0 mg L^{-1} .⁹⁰ The United States began fluoridating drinking water in public waters systems in 1950 to prevent tooth decay.⁹²

The recommended U.S. drinking water fluoride concentration is 0.7 mg L^{-1} , and the current enforceable drinking water standard (*i.e.*, Maximum Contaminant Level or MCL) for fluoride is 4.0 mg L^{-1} .⁹³ However, some evidence suggests that excessive consumption of fluoride may have clinical consequences such as adverse effects on the kidneys, including at concentrations below the U.S. enforceable limit.⁹¹ Studies have noted that increased fluoride consumption among children is a risk factor for renal damage,^{94,95} even at concentrations as low as 2.0 mg L^{-1} in drinking water, due to their ability to retain absorbed fluoride.^{91,96,97}

Fluoride as a causative factor for CKDu in Sri Lanka is of particular interest due to the remarkably confined geographical distribution of cases in the country.^{47,48} Most CKDu-affected areas in Sri Lanka are within the dry zone, an area that receives little rainfall and experiences prolonged dry periods. Additionally, groundwater fluoride concentrations in the dry zone are relatively high (0.5 mg L^{-1}) compared to the wet zone ($<0.5 \text{ mg L}^{-1}$), where incidents of CKDu are minimal.^{6,48} One study analyzed 1304 water samples from wells in all districts in Sri Lanka from 2010 to 2014, finding evidence of multiple drinking water contaminants.⁸¹ Results showed that the Sri Lankan well water was highly contaminated with fluoride, with 20% of wells containing a concentration $> 1.0 \text{ mg L}^{-1}$ (above the Sri Lankan standard of 1.0 mg L^{-1}). Of these wells, 9.9% had concentrations above the WHO's standard of 1.5 mg L^{-1} , with the highest district measuring 7.0 mg L^{-1} . Additionally, a majority of water samples showed high water hardness, with 42.2% of samples having hardness levels (in mg L^{-1} as CaCO_3) greater than or equal to 180 mg L^{-1} . Wells were also found to have elevated levels of aluminum, nitrate, arsenic and manganese, but the researchers did not associate these to possible CKDu risk factors. Based on spatial association, fluoride contamination and water hardness were proposed as suspected risk factors for CKDu.⁸¹

In another study, Balasooriya *et al.*⁸⁰ compared hydro-geochemical data from wells of consumers affected and unaffected by CKDu in Sri Lanka. A total of 63 groundwater well samples were taken, with one third (19) taken from wells used by CKDu patients. Significantly higher values of pH, total hardness, electrical conductivity, alkalinity, and calcium (Ca^{2+}), magnesium (Mg^{2+}), and F^- were observed for wells used by CKDu patients when compared to wells used by study participants without CKDu. Mg^{2+} was found to be the most prominent factor for water hardness in the wells of CKDu affected consumers, and the combination of fluoride and hard water was suggested as a potential risk factor for CKDu in the region. The median F^- concentration for non-CKDu regions was 0.49 mg L^{-1} compared to 0.63 mg L^{-1} for CKDu regions, with CKDu wells ranging from concentrations of 0.23 mg L^{-1} to 5.00 mg L^{-1} . Similarly, another study by Shi *et al.*⁹⁸ compared 52 groundwater samples from CKDu-impacted wells to 18 groundwater samples from non-CKDu impacted wells, based on official reports on CKDu well distribution in Sri Lanka. This study found significantly higher Si^{4+} and F^- concentrations in CKDu groundwater sample.⁹⁸



Gobalarajah *et al.*²⁰ collected 38 water samples in Thunukkai Division in Mullaitivu District, Sri Lanka and investigated the association between physicochemical characteristics of the drinking water and serum creatinine levels (a measure of kidney function) of CKDu patients. Of the 38 samples, 35 were taken from areas CKDu patients lived and 3 were collected as controls where there were no records of CKDu patients. The authors found that six explanatory variables accounted for 51% of variation in serum creatinine concentration of the CKDu patients: F^- , phosphate, TDS, total hardness, and arsenic ($R^2 = 0.5109$). Serum creatinine levels of the CKDu patients were significantly and negatively correlated with phosphate ($R = -0.628, p < 0.001$), as likelihood of CKDu occurrence decreased by 0.211 times for every unit increase in phosphate content. This negative influence of phosphate ions in water on serum creatinine of CKDu patients had been noted in an earlier study.⁹⁹ The study also found that TDS ($R = 0.271$) and As ($R = 0.304$) concentrations were significantly positively correlated with creatinine levels ($p < 0.05$).²⁰ Although the authors found F^- was not significantly correlated to serum creatinine levels, they noted multiple studies in the past had found significant relationships between F^- content and CKDu occurrence.⁸⁰

While fluoride has been linked to incidence of CKDu, fluoride as a single causative factor would not explain the absence of CKDu in areas located in endemic regions with similar geographic and socio-economic characteristics. Additionally, CKDu has been diagnosed in individuals who used to drink low fluoride water. Other groundwater constituents, such as hardness⁹ and more recently, silica,^{100,101} have been linked to the occurrence of CKDu. For example, one study found groundwater in CKDu endemic regions had higher average Si compared to non-CKDu regions in Sri Lanka ($p < 0.05$).⁹⁸

Correlation between high water hardness and CKDu occurrence has been extensively reported in the literature.^{20,42} Imbulana *et al.*¹⁰² found higher numbers of CKDu cases in areas using finished (*i.e.*, after treatment) drinking water with high levels of total dissolved solids, major ions (calcium, magnesium, sodium, potassium, chloride, and sulphate) and high hardness. Many earlier studies have reported a relationship between water hardness and CKDu occurrence, with one study demonstrating that 96% of patients who developed CKDu had been exposed to hard or very hard water over a period of 5 years.⁸² Hard water is attributable to the presence of polyvalent cations including calcium, magnesium, strontium, and iron, which will be present together with carbonate, bicarbonate, sulfate, and chloride anions.¹⁰³ These hardness-causing minerals enter groundwater during naturally-occurring hydro-geochemical processes, along with other elements commonly found in the geogenic environment.^{20,104} Research has suggested that hard water in combination with other elements or contaminants may increase the risk of CKDu incidence, as the presence of hardness may modify chemical forms of fluoride and trace elements in the water, acting as a secondary factor for disease etiology.^{9,83,105,106} A recent study published in 2024 suggested Mg hardness combined with excess F^- could produce nephrotoxic complexes that may trigger kidney damage in CKDu endemic areas of Sri Lanka.¹⁰⁷ Another found exposure to

Sri Lanka's groundwater caused kidney damage in zebrafish, and the authors concluded these findings suggested high water hardness and fluorine could be the inducible environmental factors for CKDu.¹⁰⁶

Fig. 4 from Wasana *et al.*⁹ shows such a correlation between high hardness, high fluoride, and CKDu prevalence. For this study, researchers divided a larger, high-risk region for CKDu in Sri Lanka into three sub zones. Zone 1 describes areas of high CKDu prevalence with a majority of residents being affected by the disease, and the main water resources are either dug or tube well water. Zone 2 refers to areas of low CKDu prevalence with the majority unaffected by disease, and the main water resource in this zone is mostly dug well water. Zone 3 encompasses areas that are within the larger high-risk region, but there is no prevalence of disease and residents here use spring water. After plotting fluoride and hardness of drinking water against prevalence of CKDu, Fig. 4 reveals a strong correlation between the high-prevalence areas of zone 1 and high levels of hardness and fluoride. Zone 2, which has less CKDu prevalence, also exhibits lower levels of hardness and fluoride, and in zone 3 where the disease is not present, these levels decrease even further.

A follow up study with mice corroborated the possible combined effects of fluoride and hard water on CKDu.⁷⁸ The authors suggested that synergic effects of cadmium, fluoride, and water hardness could lead to kidney damage in mice, even at WHO maximum recommended levels. Toxic synergistic effects of cadmium, fluoride, and hardness were more pronounced when present in water at relatively higher concentrations (*e.g.*, double the WHO maximum levels). Notably, when the mice were treated with drinking water excluding one or two of the elements, evidence of renal injury was absent.

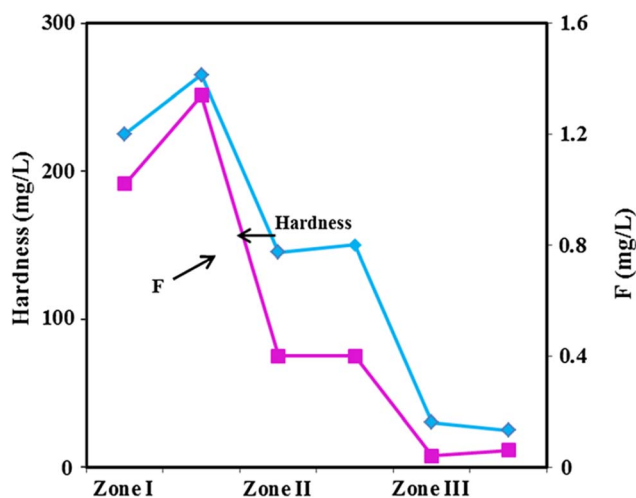


Fig. 4 Variation in levels of total hardness and fluorine in potable waters, related to prevalence of CKDu in Sri Lanka. Zone 1: high CKDu prevalence, zone 2: low CKDu prevalence, zone 3: no CKDu prevalence. Used with permission of Springer Nature BV, Drinking water quality and chronic kidney disease of unknown etiology (CKDu): synergic effects of fluoride, cadmium and hardness of water. Wasana H. M. S., Aluthpatabendi D., Kularatne W. M. T. D., Wijekoon P., Weerasooriya R., Bandara J., *Environ. Geochem. Health*, 2016, **38**(1); permission conveyed through Copyright Clearance Center, Inc.



While there is no consensus on any of these factors as a definitive cause of CKDu, sufficient evidence has established a direct or indirect relationship between disease occurrence and groundwater use. Additionally, some studies have suggested that providing water that meets drinking water standards and guidelines helps reduce the occurrence of CKDu. For example, low incidence of disease has been noted among people who consume more pristine natural spring water and treated water from large municipal water systems while living in hotspot regions of CKDu.^{102,108} One study in Sri Lanka found the highest risk for CKDu was in participants who consume well water (OR 2.52, 95% CI 1.12–5.70).⁸² Reducing excessive contents of fluoride, hardness, and other dissolved ions in groundwater has been effective in reducing the incidence of CKDu and protects residents from other health impacts related to consumption of drinking water with high fluoride and hardness.¹⁰²

4.3. Nephrotoxic heavy metals in drinking water

Trace heavy metals are common environmental pollutants that have been associated with impaired kidney function in epidemiological studies. Many such metals are nonessential for human health, toxic at low doses, and persistent in the environment. Potentially nephrotoxic heavy metals include cadmium (Cd), mercury (Hg), lead (Pb), chromium (Cr), uranium (U), gold (Au), platinum (Pt), cobalt (Co), nickel (Ni), vanadium (V), and thallium (Tl); and metalloids such as arsenic (As) and antimony (Sb).⁶ Heavy metals can bioaccumulate in the kidney, which increases the risk of kidney damage from both high-level acute and low-level chronic exposures.^{109,110} Some studies have identified heavy metals and metalloids (notably, arsenic, cadmium, and lead) as causative factors for CKDu, although some others have found conflicting results.¹¹⁰

In Western Australia, CKD has become endemic for Aboriginal people living in the area and researchers believe a high proportion of these cases are CKDu.⁴⁵ In an examination of drinking water quality in the affected communities, exceedingly high levels of both nitrate and uranium in the groundwater were observed.⁴⁵ Each of these components are thought to be nephrotoxic, but they can also combine to form a highly nephrotoxic compound called uranyl nitrate. Uranyl nitrate [$\text{UO}_2(\text{NO}_3)_2$] is a complex of the common uranium ion uranyl (UO_2^{2+}) and nitrate (NO_3^-), and it is highly soluble in water and is thus absorbed by the gastrointestinal tract more readily than low soluble uranium compounds.⁴⁵ Uranium's main target is the kidneys, and kidney damage has been seen in humans and animals after inhaling or ingesting uranium compounds.¹¹¹ However, ingesting water-soluble uranium compounds (*e.g.*, uranyl nitrate) is known to result in kidney effects at lower doses than following exposure to insoluble uranium compounds.¹¹¹

Other contaminants thought to be linked to CKDu include cadmium, lead, and arsenic, which have each been found in drinking water, foods, and soils in areas of high CKDu prevalence.¹¹² Although the United States has not yet reported hot spots of CKDu incidence, low levels of lead in community water systems have been associated with lower hemoglobin levels and higher erythropoietin stimulating agent (ESA) use among

patients with end stage kidney disease (ESKD), suggesting that even low levels of lead can have impacts on kidney health in vulnerable patients.¹¹³ Arsenic has received considerable attention over the years as a potential driver for CKDu due to widespread groundwater contamination. One study in the US investigated the relationship between urine arsenic levels and the presence of albuminuria, a marker of kidney damage, in American Indian adults from rural areas of Arizona, Oklahoma, and North and South Dakota.¹¹⁴ In a cross-sectional study, the researchers found urine arsenic concentrations were associated with increased albuminuria in a rural US population with a high burden of diabetes and obesity.¹¹⁴

Conversely, many research groups have reported large variations in concentrations of heavy metals in drinking water sources used by CKDu patients, and nearly all studies detected amounts of contaminants below the maximum acceptable levels by the WHO.^{6,80,110} Additionally, concentrations of heavy metals and metalloids have generally not shown significant variation between endemic and non-endemic areas. For example, when analyzing spatial relationships between high disease prevalence and drinking water concentrations of arsenic, lead, cadmium, or uranium, Rango *et al.*¹¹⁵ found no significant relationship linking any contaminant to CKDu. Thus, although most evidence suggests that heavy metals and metalloids do not cause CKDu on their own, given their ability to bioaccumulate in tissues, more work is needed in order to understand or eliminate heavy metals as a cause of CKDu.

4.4. Agrochemical exposures

Given the prevalence of CKDu in agricultural communities and among rural agricultural workers, multiple studies have aimed to determine the effect that pesticides and other agrochemicals can have on CKDu development. Experimental studies have suggested a relationship between pesticide exposure and impaired renal function, but few high-quality epidemiological studies exist that directly examine this relationship. One such study is the Agricultural Health Study (AHS), a longitudinal cohort study of agricultural workers and their families from Iowa and North Carolina.¹¹⁶ Based on data from the Agricultural Health Study, a smaller cohort study of licensed pesticide applicators was evaluated. ESRD risk was associated with the highest tertile of intensity-weighted use of the herbicides atrazine, metolachlor, alachlor, paraquat, and pendimethalin when compared to no use.¹¹⁶ Researchers observed a significant exposure-response trend with increasing use levels for all of these herbicides. Additionally, ESRD risk was associated with the highest tertile of metalaxyl use, a fungicide (HR = 1.92; 95% CI: 1.01, 3.66). The risk of developing ESRD was greater for study participants who had to undergo multiple doctor visits (HR = 2.13; 95% CI: 1.17, 3.89) or hospitalizations due to pesticide use (HR = 3.05; 95% CI: 1.67, 5.58).¹¹⁶ These findings support an association between ESRD and chronic exposure to certain pesticides.

The authors conducted another study investigating the relationship between ESRD among wives of licensed pesticide applicators in the same AHS cohort.¹¹⁷ The rate of ESRD was



significantly elevated among women who reported the highest *versus* the lowest cumulative pesticide use (HR: 4.22; 95% CI: 1.26, 14.20). Among women who never applied pesticides, ESRD was associated with whether their husband ever used paraquat (HR = 1.99; 95% CI: 1.14, 3.47) and butylate (HR = 1.71; 95% CI: 1.00, 2.95). These findings suggest ESRD may be associated with direct and/or indirect exposure to pesticides.

Additional studies have corroborated the connection between pesticide use and diminished kidney function. A study of a community in Mexico found that kidney function tests of pesticide-exposed farmworkers were worse than tests of workers in an unexposed control group.¹¹⁸ Another found lower eGFR among pesticide applicators who had ever used the herbicides pendimethalin, atrazine, and dicamba compared with applicators who never used each pesticide. Prior use of pendimethalin (OR = 1.6, 95% CI: 1.1, 2.2) and atrazine (OR = 1.8, 95% CI: 1.0, 3.0) was associated with elevated odds of CKD.¹¹⁹ In a 2019 study, patients with Mesoamerican nephropathy (a form of CKDu) with exposure to any agrochemical (OR = 4.86; 95% CI: 1.82, 12.96) or paraquat (OR = 12.25; 95% CI: 1.51, 99.36) was associated with kidney failure compared to non-exposed patients.¹²⁰

Glyphosate has been identified as a possible agrochemical that could lead to kidney damage.⁵⁷ Some have suggested this could be because of its unique metal chelating properties, which allow it to form a complex with nephrotoxic metals.⁷⁹ Glyphosate is one of the most widely used herbicides in the

modern world, and has often been reported as widely used in hot spot areas of the CKDu epidemic. Indeed, one study in Sri Lanka found that participants had a higher risk of CKDu if they had a history of working with spray glyphosate (OR 5.12, 95% CI 2.33–11.26).⁸²

Co-exposure to the pesticide paraquat along with glyphosate was also cited as a possible risk factor for developing chronic kidney disease.^{57,116} Indirect exposure to the pesticides paraquat and butylate were found to be associated with increased ESRD rates, with those exposed developing ESRD at more than twice the rates of those who had no exposure. The synergistic effects of both glyphosate and paraquat have also been described as playing a role in renal damage.^{57,105,121}

Another widely studied chemical class proposed as a causative factor in CKDu development is organochlorine pesticides (OCPs). Ghosh *et al.*²¹ compared levels of pesticides in the blood in healthy controls, CKDu, and CKD of known etiology participant groups. They detected nine OCPs in both groups of CKD patients and in healthy controls. In CKDu patients, increased levels of four pesticides, α -HCH, aldrin, β -endosulfan, and *p,p'*-DDE were observed when compared to levels in healthy controls. Two pesticides exhibited significant association with CKDu when using patient with CKD of known origin as the reference group: β -endosulfan (OR = 2.16) and *p,p'*-DDE (OR = 3.20).²¹ This indicates that there is a potential association between certain OCP exposures and the development of CKDu. Notably, these study participants were not directly involved in

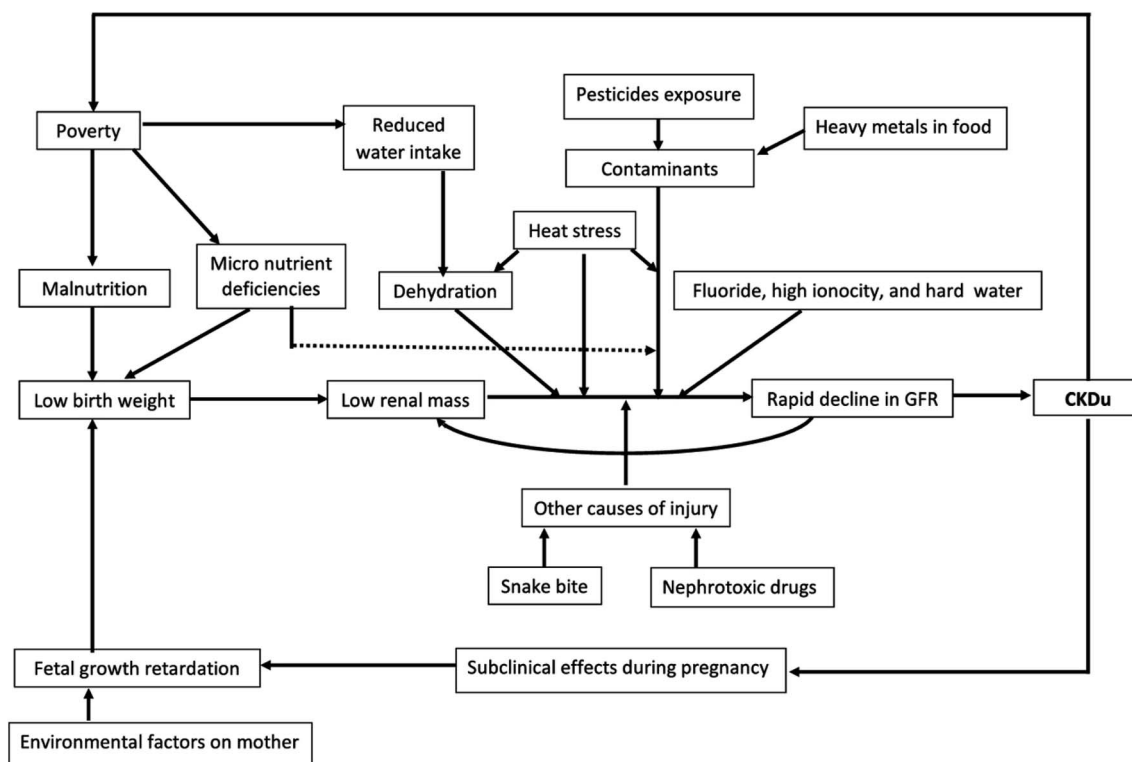


Fig. 5 Graphic visualization of a system dynamics approach to conceptualize the possible causes of the Sri Lankan CKDu epidemic. Used with permission of Elsevier Science & Technology Journals, from Chronic kidney disease of unknown etiology (CKDu): using a system dynamics model to conceptualize the multiple environmental causative pathways of the epidemic. Jayasinghe S., Zhu Y. G., *Science of the Total Environment*, 2020, 705, 135766; permission conveyed through Copyright Clearance Center, Inc.



agricultural activity or manufacturing, but the authors suggest they were likely exposed through environmental contamination.

As with each of the previously discussed proposed etiologies, the literature is not conclusive nor extensive enough to determine if agrochemicals are a definitive cause of CKDu and conflicting opinions exist. A review of 25 epidemiological studies from Meso-America found no significant associations between CKDu and pesticide exposure.¹²² Another determined that while agrochemicals may contribute to the genesis of CKDu, they are unlikely to be the sole or main cause.¹²³ Valcke *et al.*¹²⁴ found that existing studies did not provide enough evidence for an association between pesticides and regional CKDu epidemics. However, due to poor pesticide exposure assessment, the authors stated that the role of nephrotoxic agrochemicals cannot be discarded, and future research should assess lifetime exposures to relevant pesticides while also looking at potential interactions with other major risk factors.¹²⁴

4.5. Multifactorial causes

A growing number of studies have suggested that multiple factors combine to cause CKDu, and refer to CKDu as chronic kidney disease of multi-factorial origin.^{125–127} Some studies suggest a multifactorial approach because no single cause has been determined and the strong association with agricultural occupations. For instance, a system dynamics model (Fig. 5) would incorporate exposures from pesticides and heavy metals, drinking hard water with high levels of fluoride, poverty, low birth weight, micronutrient deficiencies and heat stress. Instead of using linear models, Jayasinghe and Zhu⁸ also incorporate feedback loops into their system dynamics model, which they propose is a more effective way of analyzing disease. The authors hypothesized that by using this approach, researchers can gain a better understanding of how various factors impact the disease and predict disease hot spots and system vulnerabilities. For instance, factors may overlap, such as exposure to heat stress, contaminated drinking water, and dehydration. In this way of thinking, it may be easier to quantify the interactions between different factors and unravel the potential etiologies.

5. Discussion: knowledge gaps and research needs

Based on our review of the literature, a summary of proposed etiologies, supporting evidence for each etiology, and associated areas of future research are presented in Table 1 and discussed below. Since a standardized definition for CKD was established, use of this definition has been largely accepted across the world. One of the major, remaining challenges of studying CKDu is there is no universally accepted definition, and differences remain in how CKDu is diagnosed, leading to large regional disparities in research approaches.^{128,129} While traditional CKD treatment involves control of diabetic state, blood pressure, and other risk factors, the efficacy of these therapies slowing CKDu progression is unknown and, in some cases, could potentially exacerbate it.¹²⁸ In order to effectively

treat and prevent CKDu, a consensus definition is crucial to identify the underlying cause or causes. In this review, we investigate three broadly identified potential etiologies of CKDu.

5.1. Heat stress

The only variable currently known to be constant across all regions with CKDu is the presence of heat, and therefore the ability to undergo heat stress. Collectively, research has demonstrated that a combination of extended heat exposure coupled with episodes of dehydration has the potential to cause damage or at least stress to the kidneys. If CKDu cases are a type of heat stress nephropathy, it would explain the similar epidemics occurring among agricultural workers laboring in high-heat conditions. However, if this is the case, it could be assumed that the same types of epidemics that are emerging in South Asia and Central America should be happening in equally warm areas of Africa, the Middle East, and Australia. There is some evidence of small disease clusters in these areas, like in the Aboriginal communities of Australia, but large-scale CKDu hotspots have not yet been observed. Of course, the most at-risk subjects in these areas are usually from impoverished communities where medical care may be scarce, and diagnosis would be hard to confirm.

Most data to date related to heat stress and CKDu seems to be concentrated in specific locations (*e.g.*, Central America), and more research is needed in other areas of the globe establishing heat stress as a potential risk factor. Additionally, few large-scale epidemiological studies have been conducted that isolate heat stress as a potential risk factor, and more are needed. One study that conducted a longitudinal assessment of kidney function in migrant farm workers in 2021 stated most studies related to CKDu and AKI focus on workload, heat stress, and hydration and that there is a need for studies that can evaluate occupational risk factors in combination with heat stress and dehydration.⁵⁸

5.2. Contaminated drinking water

Some research has suggested that by providing drinking water that is compliant with water quality standards and/or access to advanced drinking water treatment (*e.g.*, reverse osmosis), CKDu occurrence is reduced. Additionally, lower prevalence of CKDu has been shown among those who consume water from areas with natural springs. Accordingly, this has led many to conclude that CKDu is linked to drinking water quality and the presence of contaminants of either natural (*e.g.*, hardness causing elements) or anthropogenic (*e.g.*, agrochemical) origin.

There is need to develop a priority list of drinking water contaminants that should be the focus of future studies looking at potential associations with CKDu. From the existing literature, fluoride, hardness-causing elements, as well as select metals (*e.g.*, cadmium, arsenic) and agrochemicals (*e.g.*, nitrate and glyphosate) are high-priority contaminants that should be included in future studies focusing on links between drinking water quality and CKDu. These should be the focus of both fundamental toxicological studies establishing the mechanism,



Table 1 Summary of proposed etiologies including the supporting evidence for each, and areas where further research should be pursued

Proposed etiology	Supporting evidence	Areas of need for future study
Heat stress	<ul style="list-style-type: none"> • Warmer temperatures have been correlated with increases in CKDu cases in Central America, and regions that were hottest had the highest prevalence of disease • Sugarcane workers, who are objectively exposed to the highest levels of heat stress, were identified as the most at risk for CKDu • Cross sectional studies have identified changes in kidney function during harvests, with the most change coming from those with the highest workloads • Heat stress interventions in Central America have shown promise in reducing kidney injury 	<ul style="list-style-type: none"> • Determine if there are other hot regions of the world experiencing similar CKDu prevalence that have been previously overlooked due to limitations in medical care and diagnosis • Improve understanding as to why some hot regions of the world do not experience high levels of CKDu • Most supporting evidence for the heat stress hypothesis comes from Central America, but relatively less research has been done elsewhere (<i>e.g.</i>, Sri Lanka or India). A broad scale harvest and non-harvest study comparing kidney function should be completed to attempt to isolate heat as a variable • Why has CKDu prevalence emerged recently? Is this emergence due to increased awareness, warming induced by climate change, or other factors
Contaminated drinking water	<ul style="list-style-type: none"> • There are direct adverse effects on the kidneys from excess fluoride, and the kidney is exposed to higher concentrations of fluoride than most other soft tissues • Fluoride and high-water hardness have been shown in multiple studies to correlate to high levels of CKDu • Women and children who are not exposed to certain contaminants occupationally are still becoming ill, making drinking water a possible pathway • Strong associations exist between drinking water from abandoned wells and developing CKDu • CKDu was found significantly less or was not present in areas that used water from natural springs • Australian Aboriginal communities and those in proximity had much higher rates of renal disease, with drinking water as a common environmental factor 	<ul style="list-style-type: none"> • For often suggested contaminants linked to CKDu (<i>e.g.</i>, fluoride, hardness causing elements), studies are necessary to establish concentration thresholds and exposure durations associated with CKDu onset. Additional studies should also aim to establish a physiological mechanism by which these contaminants lead to kidney disease • Larger longitudinal studies that measure levels of contamination should be conducted to determine how contaminants might affect CKDu prevalence over time • More studies are needed to understand the synergistic effects that contaminant mixtures may exhibit. Testing more combinations of contaminants, as has been done to some extent with simultaneous measurement of fluoride, cadmium, and high hardness, could provide insightful results • Broader spectrum analytical techniques (<i>e.g.</i>, non-targeted methods for both organic and inorganic contaminants) should be applied to water samples to look for previously overlooked drinking water contaminants in CKDu affected populations
Agrochemical exposure	<ul style="list-style-type: none"> • Many pesticides and herbicides have been linked to kidney damage; for example, glyphosate and paraquat are very widely used and have been linked to kidney damage through epidemiological and animal studies • CKDu is occurring distinctly in agricultural regions, which could correlate with high interaction with pesticides 	<ul style="list-style-type: none"> • Biomonitoring studies should continue to focus on the degree of occupational exposure to pesticides for agricultural workers with CKDu. Such work will help to determine if there is a causal mechanism at play based on duration or quantity of agrochemical exposure • Occupations outside of farming have also been affected by CKDu, thus it is important to investigate other routes for agrochemical exposure beyond occupational (<i>e.g.</i>, through drinking water contamination or airborne drift to nearby communities)



Table 1 (Contd.)

Proposed etiology	Supporting evidence	Areas of need for future study
	<ul style="list-style-type: none"> • Large cohort studies have found a correlation between increased use of certain pesticides and end stage renal disease 	<ul style="list-style-type: none"> • Agrochemicals or pesticides encompass a very large group of chemical contaminants, including both active ingredients and other chemicals in more complex formulations. More definitive research is needed linking certain commonly used active ingredients and agrochemical formulations to kidney damage, with further epidemiological studies conducted with at risk populations • There is need for improved understanding as to why certain agricultural regions of the world are not experiencing high levels of CKDu despite comparable patterns in agrochemical use
Multifactorial approach	<ul style="list-style-type: none"> • The emerging CKDu epidemics of the 1990s coincide with a global increase in use of agrochemicals • The higher risk of developing CKDu in men <i>versus</i> women has been related to occupational pesticide exposure • The low, but present, risk for women and children in proximity to pesticides could be through indirect contamination or pesticide drift • Chronic diseases are commonly multifactorial (<i>i.e.</i>, diabetes, cancer) • Multiple of the suggested etiologies may be working in combination (<i>e.g.</i>, heat stress and agrochemical exposure) to drive kidney disease in certain populations 	<ul style="list-style-type: none"> • Large epidemiological studies testing the weight of multiple factors, including social determinants, on disease prevalence could help elucidate how factors may work in combination to cause kidney disease and those populations most at risk • To support such work, there is need to make more readily available and better coordinate the use of existing datasets on global CKDu incidence and associated data relevant to exposure to proposed risk factors (<i>e.g.</i>, drinking water quality, pesticide use) • Protocols for studying CKDu should be standardized so that future work consistently builds off existing literature toward a more definitive understanding of CKDu root causes

dose–response behavior, and relevant exposure timescales for onset of kidney disease, as well as larger longitudinal studies focused on population health over time. There is also the need to better account for complex mixtures of contaminants in drinking water. Though acceptable drinking water quality has been defined through guidelines and standards from the WHO and other regulatory bodies, water quality standards focus on individual contaminants, and do not account for possible synergistic effects of trace metals and other constituents, like hardness-causing elements, present in mixtures. Additional research is needed to better characterize the composition of mixtures using broader spectrum, non-targeted analytical approaches and the adverse impact on kidney health of such mixtures.

5.3. Agrochemical exposure

CKDu is often observed in agricultural workers and in areas of intense agricultural activity. Moreover, several pesticides widely used around the globe are known to negatively impact kidney function. For example, glyphosate is regulated in US drinking

water for its potential to cause kidney damage. Accordingly, many have suggested exposure to agrochemicals like pesticides can explain CKDu occurrence in certain, but not all, communities (*e.g.*, CKDu in populations that are unrelated to farming, such as mining and construction occupations).

Most studies to date that have suggested links between pesticides and CKDu cases contained unspecific and unquantified exposure measurements, thereby limiting the ability to make a significant conclusion about whether agrochemicals play a role in CKDu development. Future research should prioritize biomonitoring in affected populations, particularly agricultural workers, to strengthen possible associations between CKDu and pesticide exposure. Moreover, pesticide active ingredients are applied in formulations, which are mixtures of both the active and inert ingredients. Additional research should be careful to distinguish between potential exposure to active ingredients and formulation mixtures when assessing potential links to CKDu incidence. To address CKDu incidence beyond agricultural workers, more studies are needed establishing non-occupational exposure routes to pesticide active ingredients such as contaminated drinking water or



inhalation of airborne drift. Studies should prioritize those agrochemicals known or suspected to exhibit nephrotoxicity but should not overlook the potential for previously unrecognized toxicity for other agrochemicals, highlighting the need for ongoing toxicity studies to explore adverse health outcomes resulting from long-term exposure to widely used agrochemicals and agrochemical mixtures.

5.4. Other research needs

It is unclear if the cause for CKDu is the same across all affected regions or if they are different from region to region. No studies have found a definitive association with any single biological, agrochemical, or hydrogeochemical etiological factor. Few studies have considered the interplay of multiple risk factors or the potential role of social determinants of health. A significant challenge is that there is often little to no overlap in CKDu study designs, limiting attempts to better understand commonalities and differences of disease between regions. Finding causative risk factors across regions is challenging when using study data at a single point in time, further complicating efforts to associate any given risk factor with clinical and pathological findings. There are also limitations in data availability from region to region, which further limits comparison of global CKDu occurrence.

6. Conclusion

Chronic kidney disease is treatable and deserves more attention globally. Unlike the global decline seen in other non-communicable disease over the past few decades, CKD has not seen the same extent of improvement. An active public health response would include effective management of risk factors at both the individual and primary care levels. However, for a large portion of those with chronic kidney disease, their primary risk factor is not known. This complicates intervention, which is predicated on a thorough understanding of all risk factors and how exactly they cause disease. Despite this uncertainty, public health measures could be implemented using interventions known or suspected to reduce CKDu incidence such as providing clean drinking water or implementing a shade-rest program for workers vulnerable to heat stress. While the current CKDu epidemics remain unexplained, it is still likely that an etiology will be confirmed with more research and global awareness of the disease. For example, two other forms of nephropathy, Itai-Itai disease in Japan and Balkan endemic nephropathy (BEN), had etiologies that were uncovered in 1968 and 1993, respectively, following multiple decades of research and funding.⁸⁵

There is ample evidence that CKDu is caused by common but not necessarily singular factors across the globe. This is supported by the fact that regional epidemics are occurring simultaneously on opposite quadrants of the world, with many having similar occupational, geographical, and socioeconomic characteristics. Based on reasonably consistent findings in terms of pathologies, it is likely that some sort of toxin or contaminant is a factor in each of these epidemics. However, it

appears that no single hypothesis that has been put forth thus far can explain in totality the findings and distribution of the disease since its initial discovery. Moreover, there is little agreement between researchers as to the combination of exposures and environmental toxins that might lead to the manifestation of CKDu. The research that exists is insufficient, failing to strongly support any single etiology, and ultimately more research is needed to further establish root causes of CKDu.

Data availability

No new data were created or analyzed during this study. Data sharing is not applicable to this article.

Conflicts of interest

There are no conflicts to declare.

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