Global Cardiovascular and Renal Outcomes of Reduced GFR

Due to the number of contributing authors, the authors and affiliations are listed at the end of this article.

ABSTRACT
The burden of premature death and health loss from ESRD is well described. Less is known regarding the burden of cardiovascular disease attributable to reduced GFR. We estimated the prevalence of reduced GFR categories 3, 4, and 5 (not on RRT) for 188 countries at six time points from 1990 to 2013. Relative risks of cardiovascular outcomes by three categories of reduced GFR were calculated by pooled random effects meta-analysis. Results are presented as deaths for outcomes of cardiovascular disease and ESRD and as disability-adjusted life years for outcomes of cardiovascular disease, GFR categories 3, 4, and 5, and ESRD. In 2013, reduced GFR was associated with 4% of deaths worldwide, or 2.2 million deaths (95% uncertainty interval [95% UI], 2.0 to 2.4 million). More than half of these attributable deaths were cardiovascular deaths (1.2 million; 95% UI, 1.1 to 1.4 million), whereas 0.96 million (95% UI, 0.81 to 1.0 million) were ESRD-related deaths. Compared with metabolic risk factors, reduced GFR ranked below high systolic BP, high body mass index, and high fasting plasma glucose, and similarly with high total cholesterol as a risk factor for disability-adjusted life years in both developed and developing world regions. In conclusion, by 2013, cardiovascular deaths attributed to reduced GFR outnumbered ESRD deaths throughout the world. Studies are needed to evaluate the benefit of early detection of CKD and treatment to decrease these deaths.
Understanding the true societal effect of CKD requires evaluating the independent burdens of ESRD and reduced GFR–associated CV disease. Such data would guide national priorities regarding the benefit of early CKD detection. Early CKD detection and management could defray costs related to eventual ESRD development and higher likelihood of CV disease.

**Figure 1.** Higher burden of GFR-attributable deaths and disability-adjusted life years (DALYs) by world region in 2013. (A) Age-standardized deaths per 100,000 in 2013 attributed to reduced GFR. (B) Age-standardized DALYs per 100,000 in 2013 attributed to reduced GFR.
disease development. Thus, we aim to determine the burden of CV disease because of reduced GFR among 188 countries, compare this to the ESRD burden, and evaluate how this combined burden ranks among leading causes of health loss and premature death.

RESULTS

Global Reduced-GFR Mortality and Disability adjusted life-years in 2013

In 2013, 2.2 million deaths were associated with reduced GFR (Figure 1A, Table 1). Nearly 52 million disability-adjusted life years (DALYs) were associated with reduced GFR (Figure 1B, Table 1). These attributable deaths and DALYs account for 3.9% of total global deaths and 2.1% of total global DALYs in 2013, respectively (Table 1).

Changes in Reduced GFR DALY Rates since 1990

At the global level, among all ages, GFR-attributable median DALY counts have increased by 52.0%, respectively, whereas age-standardized rates of DALYs associated with reduced GFR have decreased by 8.1%, respectively (Supplemental Table 1, Tables 1 and 2). Compared with other metabolic risk factors, since 1990, among all ages, DALYs attributed to high fasting glucose (69.6%; 95% uncertainty interval [95% UI], 60.9 to 78.7), high total cholesterol (26.9%; 95% UI, 19.8 to 36.3), and high BP (49.1%; 95% UI, 43.2 to 55.2) have all increased (Supplemental Figure 1). Age-standardized DALY rates for the risk factors high BP (−16.5 mmHg; 95% UI, −20.0 to −12.5) and high total cholesterol (−27.5; 95% UI, −31.5 to −22.1) have decreased, whereas DALY rates associated with high fasting glucose have remained relatively constant (0.2%; 95% UI, −4.8 to 5.6).

Geographic Patterns for Reduced GFR Mortality in 2013

Among super-regions, reduced GFR ranked highest in Latin America and the Caribbean (fifth), with 7.0% of total attributed deaths and outranking metabolic risk factor high total cholesterol (seventh). Within the high-income super-region, reduced GFR ranked eighth and was outranked by all metabolic risk factors except for low bone mineral density (17th). Reduced GFR ranked lowest in sub-Saharan Africa (16th), but still outranked high total cholesterol (25th) with regards to deaths (Figures 2 and 3).

CV Disease Attributed to Reduced GFR

In 2013, globally there were 1.2 million CV deaths attributed to reduced GFR, with an age-standardized rate of 20.8 deaths per 100,000 (Table 2). Since 1990, reduced GFR–associated CV deaths have increased by 33.8% (95% UI, 24.6 to 43.8), whereas the age-standardized GFR-attributable rate has decreased by 28.6% (Table 2). The 2013 age-standardized CV mortality rate within the developing world (21.5 per 100,000) was slightly higher than in the developed world (19.2 per 100,000) (Table 2). Notably, the age-standardized CV mortality rate in the developing world has decreased slightly since 1990 (9.5% decline), whereas the developed world has demonstrated a 44% decline in age-standardized CV mortality (Table 2).

Reduced GFR was responsible for 18.7 million CV DALYs in 2013 among all ages, with an age-standardized rate of 304.2 DALYs per 100,000 (Figure 4, Table 3). The developing world had a higher rate, as well as a smaller decrease in rate since 1990, when compared with the developed world (Table 3). Among super-regions, the highest age-standardized GFR-attributed CV mortality rates were estimated for Central Europe, Eastern Europe, Central Asia, and sub-Saharan Africa (Table 2). Since 1990, GFR-attributed CV mortality rates in sub-Saharan Africa and South Asia tended to increase, as opposed to all other super-regions. The high-income super-region demonstrated the greatest decrease in mortality rate since 1990 (Table 2).

ESRD Deaths and CKD DALYs

Globally there were 956,246 (95% UI, 812,896 to 1,034,491) deaths because of ESRD in 2013, with an age-standardized rate of 15.8 per 100,000 (Table 2). Since 1990, both the number of deaths (134.6%; 95% UI, 115.7% to 150.2%) and age-standardized mortality rate (36.9%) have increased (Table 2). The age-standardized mortality rate in the developing world was almost twice that of the developed, and has increased by 44.7% since 1990, compared with the 9.8% increase within the developed world (Table 2).

The super-region with highest ESRD mortality rate in 2013 was Latin America and the Caribbean, with an ESRD mortality rate of 35 per 100,000, which is almost double the global rate (Table 2). South Asia demonstrated the greatest increase in CKD deaths (82.9%) since 1990, although all developing super-regions demonstrated an increase in ESRD mortality (Table 2).

Table 1. 2013 GFR-attributable deaths and DALY counts among all ages

<table>
<thead>
<tr>
<th>World Region</th>
<th>Mortality</th>
<th>DALYs</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Counts</td>
<td>PAF%</td>
</tr>
<tr>
<td>Global</td>
<td>2,163,699 (1,959,711 to 2,387,042)</td>
<td>3.9 (3.6 to 4.3)</td>
</tr>
<tr>
<td>Developed</td>
<td>696,832 (615,822 to 797,400)</td>
<td>5.3 (4.7 to 6.1)</td>
</tr>
<tr>
<td>Developing</td>
<td>1,466,868 (1,256,422 to 1,633,677)</td>
<td>3.5 (3.0 to 3.9)</td>
</tr>
</tbody>
</table>

Data in parentheses indicate 95% uncertainty intervals.
At the global level, age-standardized CKD DALY rates have increased by 12.3%, but this increase largely occurred within the developing world (Table 3). In 2013, CKD DALY rates were highest in the Latin America and the Caribbean region (829 per 100,000), and lowest for Central Europe, Eastern Europe, Central Asia (398 per 100,000) and Southeast Asia, East Asia, and Oceania (393 per 100,000) (Figure 5, Table 3). Age-standardized rates have decreased in three world regions since 1990 (Central Europe, Eastern Europe, and Central Asia [−6.7%]; high-income [−0.7%]; and North Africa and Middle East [−3.1%]) (Table 3). The largest increases in age-standardized CKD DALY rates occurred in Latin America and the Caribbean (39.1%) and South Asia (39.6%), driving the notable increase in the CKD DALY rate within the developing world in comparison to the developed world.

### DISCUSSION

The Global Burden of Disease (GBD) Study 2013 ranks reduced GFR as the 12th leading risk factor for CV and ESRD deaths at the global level, and the 14th leading risk factor for DALYs among 79 risk factors in 2013. Within world regions such as high-income and Latin America and the Caribbean, the mortality ranking was as high as eighth and fifth, respectively. This analysis provides granular detail regarding the contribution of CV disease caused by reduced GFR to these rankings. In 2013, >2 million deaths and 52 million DALYs were associated with reduced GFR. More than half of attributable deaths were estimated to have occurred secondary to CV disease.

The GBD Study indicates that in 1990, the developed world demonstrated notably higher rates of total CV disease among all ages than in developing world regions. Over time, the burden of age-standardized fatal CV disease has markedly declined in the developed world, whereas it has largely remained stable within developing regions. These shifting patterns of total CV disease activity in the developed and developing worlds are driving the overall pattern of GFR-attributable CV DALY and death activity. There are likely three factors contributing to the change in prevalence of total CV disease within the developing world. First, improved success in treating leading causes of premature mortality within resource-limited nations, mainly related to infectious diseases and maternal–perinatal mortality, has allowed individuals to reach more advanced age and thus develop conditions related to aging.

Second, the shift in diet toward greater intake of animal fat and high-caloric food has been documented within the developing world for at least the past decade, facilitating CV disease development. Third, limited implementation of CV risk factor detection, treatment, and disease management within the developing world in the setting of a growing burden has led to sustained mortality rates. The differences between age-standardized and all age rates merit further research.

Our results showing the attributable CV burden within the developed world may illustrate the success within regions such
as North America and Western Europe in addressing risk factors for incident CV disease over the past decades.\(^\text{22}\) Importantly, when comparing the reduced GFR risk factor to other CV risk factors, such as high blood glucose, high BP, elevated total cholesterol, and smoking, reduced GFR and elevated fasting glucose had the lowest decrease in DALY rates within the past two decades.\(^\text{17}\) This finding suggests that building on the success of CV risk factor detection and treatment will require paying equal attention to more novel CV risk factors, such as reduced GFR, in the coming years, as well as their treatments.

Regarding ESRD deaths and CKD DALYs, recent meta-analyses have estimated >2 million individuals to have died prematurely in 2010 because of lack of access to dialysis, and that the majority of these deaths occurred within Southeast Asia and Africa.\(^\text{23,24}\) Similar to these estimates, our results indicate a high burden of CKD DALYs to have occurred in sub-Saharan Africa, but our results also highlight the very high burden of CKD DALYs in Mexico, most likely related to their extraordinarily high rate of diabetes mellitus.\(^\text{25}\) Despite high death rates to ESRD within world regions with no ESRD care access, the literature also describes the encouraging statistic of growth in dialysis provision over the past two decades. Studies indicate that this growth relates more to government expansion of dialysis programs rather than the increase in causative diseases of ESRD, such as diabetes and hypertension.\(^\text{24}\) Considering this, attention needs to be paid to ESRD prevention strategies alongside continued development of ESRD programs.

The potential public health benefits of CKD screening have been evaluated in the past.\(^\text{15,24–26}\) Cost-effectiveness CKD screening studies that fully incorporated fatal and nonfatal CV outcomes as well as incident ESRD have concluded CKD screening within the general population to be cost-effective, in opposition to results of studies that only incorporated ESRD outcomes and fatal CV disease.\(^\text{16,29,30}\) These results are the first to quantify and compare ESRD and reduced GFR attributable CV outcomes across developed and developing nations, and illustrate that ESRD represents less than half of health events attributable to reduced GFR, even within the developing world. This reiterates the importance of CV fatal and nonfatal disease incorporation into evaluations of the health and economic benefits of population CKD screening. Within low- and middle-income countries, it is possible that CKD screening is of even higher importance in forestalling ESRD development, because RRT is limited in more than half of the world’s countries.\(^\text{24}\)

Evidence exists from randomized controlled trials of the benefit of statins in lowering CV morbidity.\(^\text{33}\) A recent major meta-analysis described that intensive BP control tended to reduce the relative risk of some CV outcomes, although not CV mortality or ESRD development.\(^\text{34}\) There is need for similar evidence demonstrating that diagnosing CKD early and slowing CKD progression lowers CV disease incidence and mortality. There are strong arguments that population-level CKD screening using urinary albumin and GFR measurement could potentially complement screening for hypertension and...
diabetes, the treatment of which would forestall CKD progression.\textsuperscript{33–35}

In developing countries, access to care is a barrier to chronic disease detection and treatment.\textsuperscript{26} Literature suggests that the prevalence of undiagnosed hypertension and diabetes is higher in developing than developed nations.\textsuperscript{26} Yet there are examples of efforts to detect and treat chronic diseases even within resource-limited settings. A nonprofit organization within Cambodia uses a community health worker approach to screen rural Cambodian adults for diabetes, hypertension, obesity, and CKD, and initiates treatment for these conditions, paired with diet and lifestyle modification and strict follow-up.\textsuperscript{38} Results from this organization indicate very favorable control of diabetes mellitus, hypertension, and CKD progression.\textsuperscript{39} Similar results have been published for chronic disease outreach programs in the Philippines and the Democratic Republic of Congo.\textsuperscript{40} Such screening and management interventions illustrate the importance and feasibility of chronic disease detection and management to delay progression, even in remote settings.

Within the developed region, although CKD DALY rates have decreased slightly since 1990, mortality rates have not, which could be explained in part by increasing age at death. This is in notable contrast to the almost 45% reduction over that time period in CV disease deaths associated with reduced GFR within the developed world. Success in addressing this CKD burden will likely involve continued efforts at early diagnosis and treatment of hypertension, diabetes mellitus, and early-stage CKD. These measures will also be necessary within developing nations, coupled with infrastructure development for treatment of ESRD.\textsuperscript{31,32}

Within certain regions, unique contributors to ESRD development require focused attention by that nation’s health care infrastructure. This concerns not only diabetes mellitus, hypertension, HIV, and other well known factors leading to kidney damage, but also the CKDu, the latter studied recently mainly in specific Latin American and Southeast Asian countries,\textsuperscript{41–45} with several hypothesized causes, including environmental exposures, toxins, and climate change. To date, there is no clear leading etiological factor.\textsuperscript{36} CKDu often targets young men to a higher degree than CKD caused by known factors,\textsuperscript{41–45} and is now considered endemic in countries such as Sri Lanka, India, El Salvador, and Nicaragua, and is the leading cause of hospital deaths in El Salvador.\textsuperscript{36,37} The challenges surrounding the CKDu epidemic exemplify how methods for detecting and addressing CKD burden may have to extend beyond screening for traditional CKD risk factors within classically defined high-risk portions of the population, in order to prevent ESRD deaths and CKD DALYs.

Making global estimates inherently requires assumptions and has limitations. First, CKD prevalence data were only available for 44 countries; among 1000 surveys, methods were often not optimal; and within most countries, not all time points were included. The Bayesian methods provide the best estimates possible but are limited by the quality of the available data. Furthermore, inconsistency of data collection of time and world region could influence trends and geographic variation. An area for improvement of this analysis would involve incorporating albuminuria into the exposure definition for the population-attributable fractions (PAF) calculation; there is strong evidence in the literature demonstrating the independent association between albuminuria and CV events, in isolation of and in addition to reduced GFR.\textsuperscript{46} Thus, the estimates that we present might possibly underestimate the true burden of CV events associated with CKD. These limitations serve as steps for improvement of future estimations.

Figure 3. Fluctuating ranking of GFR risk factor by world region in 2013. Risk factor ranking for deaths in 2013 per 100,000 among seven super-regions. C, Central; E, East; HI, high-income; Latin Am, Latin America; MENA, Middle East and North Africa; PUFA, polyunsaturated fatty acids; S, South; SE, Southeast; Sub-Sah, sub-Saharan.
As a significant strength of this study involves the relative risk determination for incident CV events by stage of kidney dysfunction in a global consortium. Further strengths involve our ability to estimate fatal and nonfatal burden for specific CV diseases as well as ESRD throughout the world and across time.

Reduced GFR-attributable CV deaths outnumber ESRD deaths at the global level, and the death rate from ESRD is increasing throughout the world and is a leading cause of death in world regions such as Latin America and the Caribbean. Efforts to forestall such rates will involve earlier detection of CKD. Affordable means of detecting early-stage CKD are available, as well as affordable means of treating early-stage CKD to delay progression. In order to evaluate whether such screening methods should be recommended for the general population, further cost-effectiveness analyses will need to be conducted that incorporate the 1.2 million reduced GFR-attributable CV disease deaths. It will be difficult to alter death rates within the CKD population without such studies and screening efforts.

CONCISE METHODS

Study Overview
This analysis follows the risk assessment framework used in the 2013 GBD Study for 79 individual and combined risk factors, where prevalence of the exposure is determined, a theoretical minimum risk is defined, and the relative risk of a causally-related health outcome is quantified (Supplemental Appendix 1, Supplemental Table 3). PAFs
We calculated the CV fatal and nonfatal burden attributable to the categorical exposure of reduced GFR categories using the following equation:

$$PAF = \frac{\sum_{i=1}^{n} P_i (RR_i - 1)}{\sum_{i=1}^{n} P_i (RR_i - 1) + 1}$$

where $RR_i$ is the relative risk for exposure level $i$, $P_i$ is the proportion of the population in that exposure category, and $n$ is the number of exposure categories.

Exposure
An overview of the modeling method used for determining country-level prevalence of GFR categories <15 ml/min per 1.73 m² (not on dialysis), 15–29 ml/min per 1.73 m², and 30–59 ml/min per 1.73 m² is provided in Supplemental Appendices 2 and 5. We used these country-level prevalence estimates of reduced GFR categories as our exposure categories (Supplemental Tables 4–6).
### Table 3. CV Disease CKD

<table>
<thead>
<tr>
<th>Region</th>
<th>GFR-Attributable Counts among All Ages</th>
<th>Rate per 100,000</th>
<th>% Increase in Rate per 100,000</th>
<th>AS GFR-Attributable Counts among All Ages</th>
<th>Rate per 100,000</th>
<th>% Increase in Rate per 100,000</th>
</tr>
</thead>
<tbody>
<tr>
<td>Global</td>
<td>18,697,954 (16,347,710 to 21,874,776)</td>
<td>304.2 (266.7 to 354.0)</td>
<td>-29.3 (33.6 to 24.2)</td>
<td>33,187,229 (28,460,950 to 37,316,036)</td>
<td>533.0 (426.9 to 556.4)</td>
<td>23.4 (18.0 to 29.2)</td>
</tr>
<tr>
<td>Low-income</td>
<td>2,866,731 (2,392,244 to 3,394,840)</td>
<td>160.0 (134.7 to 189.3)</td>
<td>-52.5 (55.5 to 46.5)</td>
<td>5,940,747 (4,883,654 to 7,080,702)</td>
<td>408.0 (313.1 to 498.9)</td>
<td>18.1 (12.3 to 24.9)</td>
</tr>
<tr>
<td>Central Europe and South Central Asia</td>
<td>871,368,202 (756,131,258 to 994,646,765)</td>
<td>4,207.4 (3,621.8 to 5,113.8)</td>
<td>-7.1 (18.0 to 17.0)</td>
<td>10,643,927 (8,067,850 to 13,215,998)</td>
<td>666.4 (546.7 to 886.1)</td>
<td>-3.1 (11.0 to 7.1)</td>
</tr>
<tr>
<td>Middle East</td>
<td>2,666,731 (2,392,244 to 3,394,840)</td>
<td>160.0 (134.7 to 189.3)</td>
<td>-52.5 (55.5 to 46.5)</td>
<td>5,940,747 (4,883,654 to 7,080,702)</td>
<td>408.0 (313.1 to 498.9)</td>
<td>18.1 (12.3 to 24.9)</td>
</tr>
<tr>
<td>North Africa and Sub-Saharan Africa</td>
<td>2,666,731 (2,392,244 to 3,394,840)</td>
<td>160.0 (134.7 to 189.3)</td>
<td>-52.5 (55.5 to 46.5)</td>
<td>5,940,747 (4,883,654 to 7,080,702)</td>
<td>408.0 (313.1 to 498.9)</td>
<td>18.1 (12.3 to 24.9)</td>
</tr>
<tr>
<td>Latin America and Caribbean</td>
<td>3,512,858 (2,566,177 to 4,457,148)</td>
<td>191.3 (138.9 to 254.9)</td>
<td>-17.4 (25.1 to -7.7)</td>
<td>8,059,824 (6,688,820 to 9,484,590)</td>
<td>598.5 (448.9 to 752.2)</td>
<td>6.7 (11.1 to 2.9)</td>
</tr>
<tr>
<td>South Asia</td>
<td>171,972,315 (147,282,970 to 197,661,440)</td>
<td>1,197.8 (1,081.4 to 1,326.3)</td>
<td>-30.0 (33.4 to -27.3)</td>
<td>2,207,420 (1,758,536 to 2,446,084)</td>
<td>601.1 (446.7 to 756.1)</td>
<td>30.7 (46.5 to 21.8)</td>
</tr>
<tr>
<td>East Asia and Oceania</td>
<td>1,474,518 (1,179,826 to 1,859,920)</td>
<td>1,197.8 (1,081.4 to 1,326.3)</td>
<td>-30.0 (33.4 to -27.3)</td>
<td>2,207,420 (1,758,536 to 2,446,084)</td>
<td>601.1 (446.7 to 756.1)</td>
<td>30.7 (46.5 to 21.8)</td>
</tr>
<tr>
<td>Sub-Saharan Africa</td>
<td>1,867,604 (1,562,751 to 2,234,589)</td>
<td>582.7 (492.2 to 683.4)</td>
<td>-6.1 (13.7 to 1.7)</td>
<td>3,090,221 (2,609,932 to 3,687,591)</td>
<td>568.3 (491.9 to 647.2)</td>
<td>1.0 (16.0 to 13.8)</td>
</tr>
<tr>
<td>North America</td>
<td>3,512,858 (2,566,177 to 4,457,148)</td>
<td>191.3 (138.9 to 254.9)</td>
<td>-17.4 (25.1 to -7.7)</td>
<td>8,059,824 (6,688,820 to 9,484,590)</td>
<td>598.5 (448.9 to 752.2)</td>
<td>6.7 (11.1 to 2.9)</td>
</tr>
<tr>
<td>Central Asia and South Central Asia</td>
<td>2,392,244 (1,902,452 to 2,892,037)</td>
<td>160.0 (134.7 to 189.3)</td>
<td>-52.5 (55.5 to 46.5)</td>
<td>5,940,747 (4,883,654 to 7,080,702)</td>
<td>408.0 (313.1 to 498.9)</td>
<td>18.1 (12.3 to 24.9)</td>
</tr>
<tr>
<td>Sub-Saharan Africa</td>
<td>1,867,604 (1,562,751 to 2,234,589)</td>
<td>582.7 (492.2 to 683.4)</td>
<td>-6.1 (13.7 to 1.7)</td>
<td>3,090,221 (2,609,932 to 3,687,591)</td>
<td>568.3 (491.9 to 647.2)</td>
<td>1.0 (16.0 to 13.8)</td>
</tr>
<tr>
<td>Latin America and Caribbean</td>
<td>3,512,858 (2,566,177 to 4,457,148)</td>
<td>191.3 (138.9 to 254.9)</td>
<td>-17.4 (25.1 to -7.7)</td>
<td>8,059,824 (6,688,820 to 9,484,590)</td>
<td>598.5 (448.9 to 752.2)</td>
<td>6.7 (11.1 to 2.9)</td>
</tr>
</tbody>
</table>

Data in parentheses indicate 95% uncertainty intervals. Super-regions are comprised of the following regions: High-income: Asia Pacific, Australasia, Western Europe, Southern Latin America, North America; Central Europe, Eastern Europe, Central Asia: Central Asia, Central Europe, Eastern Europe; sub-Saharan Africa: Central sub-Saharan Africa, Eastern sub-Saharan Africa, Southern sub-Saharan Africa, Western sub-Saharan Africa; North Africa and Middle East: North Africa, Middle East; South Asia: South Asia; Southeast Asia, East Asia; and Oceania: East Asia, Southeast Asia, Oceania. AS, age-standardized.

---

Outcomes

CV outcomes of ischemic heart disease, stroke, as well as peripheral vascular disease, ESRD deaths, and CKD DALYs at the country, age, sex, and year level were obtained from the 2013 GBD Study (Supplemental Appendix 4).

World Regions

The countries of which regions and super-regions are comprised are listed in Supplemental Table 1.

ACKNOWLEDGMENTS

The authors thank the staff and participants of the Atherosclerosis Risk in Communities (ARIC) Study for their important contributions. The authors thank the other investigators, the staff, and the participants of the Multi-Ethnic Study of Atherosclerosis (MESA) for their valuable contributions. Dade Behring (Marburg, Germany) supplied equipment and reagents for nephelometric measurement of urinary albumin. The authors thank the other investigators, the staff, and the participants of the Reasons for Geographic and Racial Differences in Stroke (REGARDS) Study for their valuable contributions. The authors thank the doctors, nurses, students, volunteers, and organizations involved in country screening programs which made collection of data for the International Society of Nephrology–Kidney Disease Data Center (ISN-KDDC) Study possible.

The ARIC study is carried out as a collaborative study supported by the National Heart, Lung, and Blood Institute (NHLBI; contracts HHSN268201100005C, HHSN268201100006C, HHSN268201100007C, HHSN268201100008C, HHSN268201100009C, HHSN268201100010C, HHSN268201100011C, and HHSN268201100012C). The Australian Diabetes, Obesity and Lifestyle Study is supported by the Baker IDI Heart and Diabetes Institute, Melbourne, Australia, their sponsors, and the National Health and Medical Research Council of Australia (grant 233200), Amgen Australia, Kidney Health Australia, and The Royal Prince Alfred Hospital, Sydney, Australia. Beaver Dam is supported by the National Institutes of Health (NIH) and National Institute of Diabetes and Digestive and Kidney Diseases (NIDDK) grants NIH/NIDDK DK7321 and NIH/NEI EY F06594, and an unrestricted grant from Research to Prevent Blindness. The Epidemiological investigations on chances of preventing, recognizing early and optimally treating chronic diseases in an elderly population study is supported by the Ministry of Research, Science and the Arts Baden-Württemberg (Stuttgart, Germany), Federal Ministry of Education and Research (Berlin, Germany), Federal Ministry of Family Affairs, Senior Citizens, and...
Women and Youth (Berlin, Germany), and the European Commission FP7 framework programme of DG-Research (Consortium on Health and Ageing: Network of Cohorts in Europe and the United States Project). Measurement of urinary albumin was funded by Dade-Behring, Marburg, Germany. Boris Bikbov has received funding from the European Union Horizon 2020 research and innovation programme under the Marie Skłodowska-Curie grant agreement no. 703226. The NHLBI Framingham Heart Study is supported by grant N01-HC-25195. The Global Burden of Disease Study is supported by the Bill and Melinda Gates Foundation. The MESAs study is supported by contracts N01-HC-95159 to N01-HC-95169 from the NHLBI and by grants UL1-RR-00040 and UL1-RR-025005 from National Center for Research Resources. The Ohasama Study is supported by grants-in-aid (H20-22)Junkankitou [Seishuu]-Ippan-009, 013 and H23-Junkankitou [Senshhu]-Ippan-005) from the Ministry of Health, Labor and Welfare, Health and Labor Sciences Research Grants, Japan, and the Japan Atherosclerosis Prevention Fund. The Uppsala Longitudinal Study of Adult Men is supported by the Swedish Research Council, the Swedish Heart-Lung Foundation, the Marianne and Marcus Wallenberg Foundation, Dalarö University, and Uppsala University. The Prevention of Renal and Vascular End-stage Disease Study is supported by several grants from the Dutch Kidney Foundation, and grants from the Dutch Heart Foundation, the Dutch Government, the NIH, and the University Medical Center Groningen, The Netherlands. The REGARDS research project is supported by a cooperative agreement (U01 NS041588) from the National Institute of Neurological Disorders and Stroke, NIH, Department of Health and Human Service. Additional funding was provided by an investigator-initiated grant-in-aid from Amgen and an investigator-initiated NHLBI grant (R01 HL080477). The Severance Study is supported by the Seoul City R&D program (10526), Korea, the National R&D Program for Cancer Control, Ministry for Health, Welfare and Family Affairs, Republic of Korea (1220180), and the National Research Foundation of Korea grant funded by the Korean Government (2011-0029348). The Gubbio Study is supported by the Municipal and Health Authorities of Gubbio, Italy; the Federico II University, Naples, Italy; University of Milan, Milan, Italy; Istituto Superiore di Sanità, Rome, Italy; Northwestern University, Chicago; University of Salerno, Italy; and Merck Sharp & Dohme, Italy. The ISN-KDDC Study is supported by the International Society of Nephrology, Research and Prevention Committee, Brussels, Belgium and the Istituto di Ricerche Farmacologiche Mario Negri, Clinical Research Center for Rare Diseases Aldo e Cele Daccò, Bergamo, Italy. The Rancho Bernardo Study is supported by grants National Institute on Aging AG07181 and AG028507 NIDDK DK31801.

These results do not reflect the views of The Bill and Melinda Gates Foundation. The content is solely the responsibility of the authors and does not necessarily represent the official views of the National Institute of Neurological Disorders and Stroke or the NIH. Representatives of the National Institute of Neurological Disorders and Stroke, NIH, Department of Health and Human Service have been involved in the review of the manuscript but not directly involved in the collection, management, analysis, or interpretation of the data. Representatives from Amgen and NHLBI did not have any role in the design and conduct of the
study, the collection, management, analysis, and interpretation of the data, or the preparation or approval of the manuscript.

A full list of participating MESA investigators and institutions can be found at http://www.mesa-nhlbi.org. A full list of participating REGARDS investigators and institutions can be found at http://www.regardsstudy.org.

DISCLOSURES
None.

REFERENCES


37. Rodríguez MI: Sounding the alarm on chronic kidney disease in farming communities: María Isabel Rodríguez MD. Minister of health, El Salvador. Interview by Conner Gorry. MEDICC Rev 15: 8–10, 2013


This article contains supplemental material online at http://jasn.asnjournals.org/lookup/suppl/doi:10.1681/ASN.2016050562/-/DCSupplemental.
Huntington, WV; Institute of Health Care and Social Sciences, Hochschule für Oekonomie & Management University, Essen, Germany; St. Louis, MO; Federal University of Santa Catarina, Florianopolis, Brazil; Health Care Center of Anjo Kosei Hospital, Anjo City, Japan; Department of Internal Medicine, Federal Teaching Hospital, Abakaliki, Nigeria; Ibaraki Prefectural University of Health Sciences, Ibaraki, Japan; Center for Disease Burden, Norwegian Institute of Public Health, Bergen, Norway; Department of Global Public Health and Primary Care, University of Bergen, Bergen, Norway; Competence Center Mortality-Follow-Up of the German National Cohort, Federal Institute for Population Research, Wiesbaden, Germany; Department of Public Health Medicine, University of Tsukuba, Tsukuba, Japan; Department of Preventive Medicine, Northwestern University, Chicago, IL; and Department of Biostatistics, School of Public Health, Kyoto University, Kyoto, Japan