Chronic kidney disease among the Iranian-Azari population; a report from pilot phase of AZAR cohort study

Fariba Mahmoodpoor¹, Mohammad-Reza Ardalan¹ *, Mohammadhossein Somi¹, Elnaz Faramarzi², Sepideh Zununi Vahed¹, Mousa Ghaoyr Nahand¹

¹Kidney Research Center, Tabriz University of Medical Sciences, Tabriz, Iran
²Liver and Gastrointestinal Diseases Research Center, Tabriz University of Medical Sciences, Tabriz, Iran

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**Abstract**

Introduction: Chronic kidney disease (CKD) is a worldwide health problem. Most patients with CKD are asymptomatic and unaware of their kidney disease until reaches its later stages. The worldwide prevalence of CKD is increasing and delayed diagnosis takes from the patients the opportunities for early treatment.

Objectives: Our cohort, named AZAR cohort, is a part of a national cohort program and it is aimed to study the major cardiovascular, pulmonary, renal, diabetes, and cancers diseases risk factors in East-Azerbaijan province in North West of Iran.

Patients and Methods: In this cross-sectional study, all eligible subjects (35-65 years) were recruited. Information about demographic, medical history, some physical and para-clinical were collected. Here, we report the results of pilot phase of this study.

Results: Findings showed that prevalence of CKD (stage 3, eGFR<60 mL/min) among our studied population was 8% (71/898). The studied population was ranged from young adult to pre-elderly (35-65 years). We found a higher proportion of CKD in women (M/F 14/57). The total proportion of diabetics in our study was 126 persons (14%). Distribution of different stages of CKD was as follows: stage 3A (67 persons, M/F; 14/53), stage 3B (4 females), stage 2 (737 persons, M/F; 328/409) and stage 1 (88 persons, M/F; 64/24). There were no cases of CKD stage 4 or 5 in our cohort. Body mass index (BMI), serum triglyceride (TG), and cholesterol levels had a significant correlation with CKD stage 3 (P<0.05). Diabetic patients and female gender were at increased risk of CKD stage 3 (OR: 1.5; 95% CI: 0.857-2.861).

Conclusion: The prevalence of CKD stage 3 in our cohort was compatible with other previous publications. The higher prevalence of CKD in middle aged to early elderly women population could be explained by the high prevalence of obesity among this population.

**Implication for health policy/practice/research/medical education:**
Early diagnosis and implementation of simple preventive measurements are the key elements to fight against the risk of CKD. In this regard awareness about the prevalence of CKD and its probable risk factors are the first steps toward those goals.


Introduction

Chronic kidney disease (CKD) is a worldwide health problem. Most patients with CKD are asymptomatic and unaware of their kidney disease until reaches its later stages (1). The worldwide prevalence of CKD is estimated between 8% to 16% and it is increasing. It seems that this upward trend is more prominent among low and middle-income population with limited access to health care. The proportion of end-stage renal disease (ESRD) population who are impending to start the dialysis vastly underestimates global CKD burden because pre-dialysis CKD exceeds ESRD prevalence by as much as 50 times.
(2,3). Sometimes nonspecific symptoms of CKD direct patients toward different specialties. Delayed diagnosis takes from the patients the opportunities for early treatment. CKD not only progress to the ESRD but also is a major risk factor for cardiovascular disease (CVD). Early detection of kidney diseases and timely referral to nephrology improved the patients’ outcomes (4). The traditional approach recommends screening among patients with diabetes mellitus, hypertension or CVD. For increasing the screening coverage consideration of older than 60 years population and low socioeconomic status (SES) has been proposed. Early stages of CKD can be detected by simple methods such as urine dipstick and serum creatinine measurement (4-6).

**Objectives**

In this study we assessed the prevalence of CKD in Iranian-Azari population.

**Patients and Methods**

**Study design and setting**

Azar cohort is a prospective population-based study and a part of a national screening program named Persian cohort. It is aimed to study the major non-communicable diseases risk factors, including cardiovascular, pulmonary, renal, diabetes and cancers disease. Azar cohort is started from October 2014 and still it is continuing in East-Azarbaijan province in North West of Iran. It is expected up to 15000 persons within the age range of 35-70 years will enter the study. Our study was based on the results of pilot phases of the Azar cohort that was gathered between October 2014 up to January 2015. During the pilot phase all eligible subjects of a small city named ‘Khameneh’ were studied. In our study we entered the population between the ages of 35-65 years. The exclusion criteria were pregnancy and unwillingness to participate.

**Measurements**

In this study all participants underwent general history taking and physical examination. History of presence of diabetes, hypertension, cardiovascular disease, herbal medicine taking and dietary habit were taken. Anthropometric measurements including height, weight, waist, and hip and also wrist circumference were recorded. Blood pressure was measured twice with 10 minutes apart from both upper extremities separately and the mean of measurements was recorded as the blood pressure of each person. From each individual 17 mL of blood were taken after at least 12 hours of fasting, for following measurements; fasting blood glucose (FBS), hemoglobin A1c, lipids profile, liver enzymes, complete cell blood count (CBC), thyroid hormones (T3,T4,TSH), C reactive protein (CRP), ESR. Additionally serum calcium, phosphorus, sodium, potassium and blood urea nitrogen (BUN) was assessed. Serum creatinine was measured by Jaffé reaction. Urinary protein excretion, glucosuria and urinary ketone were evaluated by the dipstick test (Acon –china LCA5721-01). Second screening by urine dipstick was performed in those who had proteinuria and hematuria in their first screening. A 24 hours proteinuric measurement was performed in those who had proteinuria and hematuria. Individuals with persistent hematuria were referred to nephrology clinics for detection of dysmorphic red blood cells. Those with GFR<45 mL/min were also referred to nephrology clinic. Estimated glomerular filtration rate (eGFR) was calculated using the simplified Modification of Diet in Renal Disease (MDRD) study equation (7,8). Body mass index (BMI) was also calculated. Diabetes mellitus was identified as a documented history of diabetes mellitus or elevated blood glucose (FBS ≥126 mg/d).

A person’s blood pressure was calculated as the average of the two measurements in each arm. Subjects with SBP ≥140 and DBP ≥90 or self-reported having hypertension were defined as patients with hypertension.

Our definition of CKD was based on the National Kidney Foundation Kidney Disease Outcomes Quality Initiative working group definition of CKD (7).

CKD stages were defined as follows: stage 1, GFR ≥90 mL/min/1.73 m², stage 2; GFR between 60-89 mL/min/1.73 m²; stage 3, GFR between 30-59 mL/min/1.73 m², stage 3a (46-59), 3b (30-45), stage 4; GFR between 15-29 mL/min/1.73 m²; and stage 5, GFR < 15 mL/min/1.73 m². In our study we only considered persons with CKD stage 3 or higher as our screening target (eGFR below 60 mL/min (7).

**Ethical issues**

This cross-sectional study was approved by Ethics Committee of Tabriz University of medical sciences (tbzmed.rec.1393.205). The research followed the tenets of the Declaration of Helsinki. Written informed consent was obtained from all patients and remained confidential.

**Statistical analysis**

The results of the study are presented as frequencies, percentages, mean ± standard deviation. We used simple statistical methods for descriptive parameters such as age, weight, high and blood pressure measurements. We used conditional logistic regression analysis for estimating crude and adjusted odds ratios (OR) and their respective 95% CIs. The correlation of different stages of CKD with hypertension, BMI, cholesterol was evaluated by Spearman’s correlation coefficient test. The analysis was conducted using the SPSS software (Statistical Package for the Social Sciences, version 11.5) and P value less than 0.05 was considered significant.

**Results**

Baseline characteristics of the studied population are presented in Table 1. Around 898 (M/F: 406/492 with ages of 35 to 65 years) were included in the study. Proportion
of diabetics in our study was 126 persons (14%) and 28 of them were newly diagnosed in our cohort. The mean of BMI in males and females with stage 3 CKD was as the following (30.12 ±4.32 kg/m² vs 26.54± 3.66 kg/m²; P = 0.004). Around, 71 persons (%8 M/F 14/57) had a reduced eGFR (defined as a value <60 mL/min/1.73 m²). Distribution of different stages of CKD were as the following; stage 3A (67 persons, M/F; 14/53), stage 3B (4 female), stage 2 (737 persons, M/F; 328/409) and stage 1 (88M/F; 64/24) (Table 2). We did not find persons with CKD stage 4 or 5 in our cohort. After one year in second measurements of eGFR only 42 from 71 people who were discovered as CKD stage 3 remained in this group, and 29 persons were returned to CKD stage 2. All individuals with CKD stage 3B or those with persistent proteinuria and hematuria were referred to nephrology clinic for future follow up.

The presence of hypertension tentatively increases the risk of CKD stage 3 two times (OR: 2; 95% CI: 0.90-5.57). BMI, triglyceride (TG), and cholesterol had a meaningful correlation with CKD stage 3 (P<0.05). Diabetic patients were at increased risk of CKD stage 3 (OR: 1.5; 95% CI: 0.85-2.861).

**Discussion**

The result of our study revealed that prevalence of CKD (stage 3, eGFR<60 mL/min) among our studied population (35-65 years) was 8%. The studied population was ranged from young adults to pre-elderlies who are making the most active part of society. We did not investigate pediatrics, teenagers and elderlies (>65 years), hence the result of our study could not extrapolate to the general population. We found a higher proportion of CKD in women. In our study BMI had a positive correlation with CKD prevalence. The higher prevalence of CKD in women could be explained by the high prevalence of obesity among this population. Presumably lower levels of recreational physical activity and regular sport activities could account for the high prevalence of obesity and observed difference in CKD prevalence. The positive correlation between high cholesterol and triglyceride levels and CKD prevalence is also connected to this explanation. As we found in our study, hypertension by itself and as a part of metabolic syndrome is associated within increment of CKD. Importantly, we did not discover any patient with advanced CKD (stage 4 or 5). Age limit of our study (<65) may be an explanation.

In the first 25 years of life the most common causes of CKD are congenital anomalies of the kidneys and urinary tract, steroid-resistant nephrotic syndrome (SRNS), chronic glomerulonephritis and renal cystic diseases and ciliopathies. Monogenic cause of renal disease can be detected in up to 20% of those individuals. Because of our age selection (>35 years) we cannot estimate the above conditions in our study. In another CKD screening program that was performed in Golestan province of Iran the prevalence of CKD stage 3-5 was 5% among the 3591 participants ( >18 years) (9). In a study on >60000 individuals from Rumania, the prevalence of CKD (eGFR <60 mL/min), or positive urinary dipstick test was 6.69%. The mean age of the studied population was 55 years and CKD was more common in elders (10). The approximate prevalence of CKD stage 3 among adults over the age of 65 was 11% in a report from the United States. A selected (>40 years) population study on 6200 individuals in northern Italy showed a prevalence of 6.4% for CKD stage 3 (11). Likewise, in a study from the Netherlands the reported prevalence of CKD was 5.8% (mean age was 49 years) (12). The trend of increased prevalence of CKD with increasing the age was preserved in all above studies and also in a recent cross-sectional study from Spain (13). Serum creatinine as a screening biomarker has several limitations and delays identification of the

**Table 1.** Demographic characteristics of participants (n = 898)

<table>
<thead>
<tr>
<th>Age (y)</th>
<th>Male</th>
<th>Female</th>
<th>Total</th>
<th></th>
</tr>
</thead>
<tbody>
<tr>
<td>Weight (kg)</td>
<td>49.9±8.026</td>
<td>48±7.67</td>
<td>48.9±7.9</td>
<td></td>
</tr>
<tr>
<td>Height (cm)</td>
<td>78.5±13.7</td>
<td>72.6±13.3</td>
<td>75.2±13.7</td>
<td></td>
</tr>
<tr>
<td>BMI (kg/m²)</td>
<td>26.5±4.22</td>
<td>29.3±5.2</td>
<td>284±8.4</td>
<td></td>
</tr>
<tr>
<td>Cholesterol (mg/dL)</td>
<td>179.89±37.2</td>
<td>191.21±38.6</td>
<td>186.1±38.3</td>
<td></td>
</tr>
<tr>
<td>HDL (mg/dL)</td>
<td>43.9±9.6</td>
<td>51.4±10.5</td>
<td>48±10.8</td>
<td></td>
</tr>
<tr>
<td>Triglyceride (mg/dL)</td>
<td>160.7±88.1</td>
<td>155.6±83.7</td>
<td>157.9±86.1</td>
<td></td>
</tr>
<tr>
<td>Blood pressure</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Systolic (mm Hg)</td>
<td>109±15</td>
<td>106.5±15.4</td>
<td>107.6±15.3</td>
<td></td>
</tr>
<tr>
<td>Diastolic (mm Hg)</td>
<td>69.6±18.5</td>
<td>68.3±8.2</td>
<td>68.9±8.4</td>
<td></td>
</tr>
<tr>
<td>Fasting blood glucose (mg/dL)</td>
<td>102.4±32.2</td>
<td>103.9±37.5</td>
<td>103.2±35.2</td>
<td></td>
</tr>
</tbody>
</table>

**Table 2.** The prevalence of CKD stratified by age and gender

<table>
<thead>
<tr>
<th>CKD stages</th>
<th>Female, No. (%)</th>
<th>Male, No. (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>35-44</td>
<td>45-54</td>
<td>55-65</td>
</tr>
<tr>
<td>Stage 1 (GFR ≥90 ml/min)</td>
<td>14 (8.3)</td>
<td>8 (3.6)</td>
</tr>
<tr>
<td>Stage 2 (GFR=60-89 ml/min)</td>
<td>148 (88)</td>
<td>183 (82)</td>
</tr>
<tr>
<td>Stage 3a (GFR=46-59 ml/min)</td>
<td>6 (3.8)</td>
<td>31 (13.9)</td>
</tr>
<tr>
<td>Stage 3b (GFR=30-45 ml/min)</td>
<td>0</td>
<td>1 (0.45)</td>
</tr>
<tr>
<td>Stage 4 (GFR=15-29 ml/min)</td>
<td>0</td>
<td>0</td>
</tr>
<tr>
<td>Stage 5 (GFR&lt;15 ml/min)</td>
<td>0</td>
<td>0</td>
</tr>
</tbody>
</table>
disease process. For example, in AKI, creatinine does not elevate until the injury is well established (14). In this study, serum creatinine as a marker of renal function was used. In a cross-sectional study from Japan the prevalence of proteinuria (dipstick test >30 mg/dL) and hematuria (dipstick) in general population was 5.3% and 9.0% respectively (15). Japanese investigators have reported a change from glomerulonephritis to diabetic nephropathy and hypertensive nephrosclerosis was the most common causes of CKD (14,16). We believe such a deviation has happened in our population too. Some studies did not recommend the community based CKD screening in asymptomatic adults (17), and rather considered it for diabetics or hypertensive population, or those with a family history of diabetes, hypertension, or CKD (6,18).

Conclusion
The prevalence of kidney disease in our report was in concordance with other national and international results and shows the influence of known CKD risk factors in our society. Screening for kidney disease and education of health givers is very important. Implementation of simple preventive measurements and the early diagnosis are the key elements to combat this rising threat.

Limitations of the study
The most important limitation of the study was its small sample size due to the pilot study of cohort. We suggest larger studies on this topic.

Authors’ contribution
Study concept, design, and supervision: MRA and MS. Acquisition of data: EF and FM. Drafting of the manuscript: MRA, EF, and FM. Critical revision of the manuscript for important intellectual content: MRA, SZV. Statistical analysis: FM and EF. Administrative, technical and material support by MS and MGN.

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Conflicts of interest
The authors declared no potential conflicts of interest with respect to the research, authorship, and/or publication of this article.

Ethical considerations
Ethical issues (including plagiarism, data fabrication, double publication) have been completely observed by the authors.

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References


