Soluble TNF Receptors and Kidney Dysfunction in the Elderly

Axel C. Carlsson,*† Tobias E. Larsson,‡ Johanna Helmersson-Karlqvist,§ Anders Larsson,§ Lars Lind,§ and Johan Ärnlöv†

*Centre for Family Medicine, Department of Neurobiology, Care Sciences and Society, Karolinska Institutet, Huddinge, Sweden; †Department of Medical Sciences, Molecular Epidemiology and Science for Life Laboratory, Uppsala University, Uppsala, Sweden; ‡Department of Clinical Science, Intervention and Technology, Karolinska Institutet, Stockholm, Sweden; §Department of Medical Sciences, Uppsala University Hospital, Uppsala, Sweden; and ||School of Health and Social Studies, Dalarna University, Falun, Sweden

ABSTRACT

The importance of TNF-α and its soluble receptors (sTNFR1 and sTNFR2) in the development of kidney disease is being unraveled. Yet, community-based data regarding the role of sTNFRs are lacking. We assessed serum sTNFRs and aspects of kidney damage cross-sectionally in two independent community-based cohorts of elderly participants: Prospective Investigation of the Vasculature in Uppsala Seniors (n=815; mean age, 75 years; 51% women) and Uppsala Longitudinal Study of Adult Men (n=778; mean age, 78 years). Serum sTNFR1 correlated substantially with different aspects of kidney pathology in the Uppsala Longitudinal Study of Adult Men cohort (R=0.52 for estimated GFR, R=0.22 for urinary albumin-to-creatinine ratio, and R=0.17 for urinary kidney injury molecule-1; P<0.001 for all), with similar correlations in the Prospective Investigation of the Vasculature in Uppsala Seniors cohort. These associations remained significant after adjustment for age, sex, inflammatory markers, and cardiovascular risk factors and were also evident in participants without diabetes. Serum sTNFR2 was associated with all three markers in the Prospective Investigation of the Vasculature in Uppsala Seniors cohort (P<0.001 for all). Our findings from two independent community-based cohorts confirm and extend results of previous studies supporting circulating sTNFRs as relevant biomarkers for kidney damage and dysfunction in elderly individuals, even in the absence of diabetes.


TNF-α is a central player in the human immune system, and inflammatory and stress response pathways are activated as soon as TNF-α binds to TNF receptors (TNFRs).1 However, the effects of TNF-α are also regulated by soluble TNFR (sTNFR) in plasma: One effect is that they block TNF-α from binding its target cell surface receptor, and another is a prolonged and delayed effect of TNF-α. Additional unique properties of sTNFRs that do not involve TNF-α are also present.2 Two sTNFRs—sTNFR1 and sTNFR2—are known, and their importance in the development of kidney diseases is being explored.3

Rat models have suggested a causal role for sTNFRs in diabetic nephropathy.4 Interestingly, two recent studies report that soluble TNFRs predict progression of CKD and development of ESRD in patients with diabetes.5,6 Associations between sTNFRs and progression from microalbuminuria to macroalbuminuria,7 and of decline in eGFR8 in type 1 diabetes, have also recently been reported. However, to date, community-based
data on the association between sTNFRs and kidney damage are scarce.

We hypothesize that soluble TNFRs play a causal role in the development of kidney damage and dysfunction. Herein, we aimed to explore and validate the cross-sectional associations between soluble TNFRs and markers of kidney damage and dysfunction used in clinical practice (eGFR) and microalbuminuria (albumin-to-creatinine ratio [ACR]) in two independent community-based cohorts of elderly persons. As a secondary aim, we wanted to explore the association between sTNFRs and kidney tubular damage (using the specific tubular damage biomarker urinary kidney injury molecule-1 [KIM-1]).

RESULTS

Baseline Characteristics

Baseline characteristics of the study populations are presented in Table 1. The associations between sTNFRs and CKD stages according to the GFR strata in the Kidney Disease Improving Global Outcomes guidelines are shown in Table 2. Higher levels of sTNFRs were seen in individuals with lower GFR.

Correlations and Linear Regression Models

Figure 1, A–F, shows scatterplots of the association between sTNFR1 and eGFR, ACR, and KIM-1 in the Prospective Investigation of the Vasculature in Uppsala Seniors (PIVUS) and Uppsala Longitudinal Study of Adult Men (ULSAM). In these analyses, higher sTNFR1 was significantly associated with lower eGFR, higher ACR, and higher U-KIM-1, with salient similarities in the associations between sTNFR1 and each marker in the two cohorts. The Pearson correlation coefficients and P values are also shown in the figures.

Table 3 shows linear regression models of the association between sTNFR1 and markers of kidney and tubular dysfunction in the ULSAM and PIVUS cohorts. ACR, eGFR, and U-KIM-1 were significantly associated with sTNFR1 in all linear regression models tested. The association was strongest with eGFR, followed by ACR and U-KIM-1.

The linear regression coefficients in a model with all three kidney markers as explanatory variables of sTNFR1 in ULSAM were as follows: eGFR, −0.48 (95% confidence interval [95% CI], −0.54 to −0.42; P<0.001); ACR, 0.068 (95% CI, 0.037 to 0.14; P=0.06); and U-KIM-1, 0.044 (95% CI, −0.026 to 0.11; P=0.21). The corresponding linear regression model with all three kidney markers in PIVUS revealed similar regression coefficients and significant associations between sTNFR1 and the following: eGFR, −0.60 (95% CI, −0.65 to −0.54; P<0.001); ACR, 0.12 (95% CI, 0.062 to 0.17; P<0.001); and U-KIM-1, 0.062 (95% CI, 0.010 to 0.12; P=0.020).

The association between sTNFR1 and eGFR was even more pronounced in participants with eGFR≥60 ml/min per 1.73 m² (significant multiplicative interaction in both ULSAM and PIVUS, P<0.001). The regression coefficients for a 1-SD increment of eGFR in participants with eGFR<60 ml/min per 1.73 m² were −1.13 (95% CI, −1.33 to −0.94; P<0.001) in ULSAM and −1.28 (95% CI, −1.50 to −1.06; P<0.001) in PIVUS. The corresponding regression coefficients in individuals with eGFR≥60 ml/min per 1.73 m² were −0.38 (95% CI, −0.48 to −0.28; P<0.001) in ULSAM and −0.35 (95% CI, −0.42 to −0.28; P<0.001) in PIVUS.

Data on sTNFR2 were available in the PIVUS study only and correlated with TNFR1 (r=0.5937; P<0.001).

Pearson correlations and linear regression models between sTNFR2 and kidney markers are shown in Table 4. ACR and eGFR were associated with sTNFR2 in all linear regression models. KIM-1 was weakly associated with sTNFR2 and was nonsignificant after adjustments for C-reactive protein (CRP) and cardiovascular risk factors. The linear regression coefficients in a model with all three kidney markers as explanatory variables of sTNFR2 were as follows: eGFR, −0.50 (95% CI, −0.55 to −0.44; P<0.001); ACR, 0.064 (95% CI, 0.003 to 0.12; P<0.05); and U-KIM-1, 0.059 (95% CI, −0.018 to 0.12; P=0.06).

The association between sTNFRs and the different aspects of kidney damage and dysfunction were similar after adjustment for level of physical activity (data not shown).

Finally, there were significant differences in medians and their Bonnet-Price CIs in individuals with and without diabetes (P=0.06 for sTNFR1 in ULSAM; P<0.001 for sTNFR1 and P<0.001 for sTNFR2 in PIVUS). The association between the TNFRs and different aspects of kidney damage and dysfunction were also similar in participants with and without diabetes in both cohorts (Supplemental Table 1).

Table 1. Baseline characteristics of PIVUS and ULSAM

<table>
<thead>
<tr>
<th>Variable</th>
<th>PIVUS</th>
<th>ULSAM</th>
</tr>
</thead>
<tbody>
<tr>
<td>Participants (n)</td>
<td>815</td>
<td>778</td>
</tr>
<tr>
<td>Women, n (%)</td>
<td>414 (51)</td>
<td>0</td>
</tr>
<tr>
<td>Age (yr)</td>
<td>75.3±0.2</td>
<td>77.6±0.8</td>
</tr>
<tr>
<td>CRP (mg/L)</td>
<td>3.9±6.3</td>
<td>3.9±2.7</td>
</tr>
<tr>
<td>IL-6 (pg/ml)</td>
<td>–</td>
<td>3.9±2.7</td>
</tr>
<tr>
<td>sTNFR1 (pg/ml)</td>
<td>2455±1293</td>
<td>2081±865</td>
</tr>
<tr>
<td>sTNFR2 (pg/ml)</td>
<td>6332±2880</td>
<td>–</td>
</tr>
<tr>
<td>Urinary KIM-1–to-creatinine ratio (ng/mmol)</td>
<td>173±1597</td>
<td>118±89</td>
</tr>
<tr>
<td>GFR (ml/min per 1.73 m²)</td>
<td>68±19</td>
<td>73±17</td>
</tr>
<tr>
<td>Urinary ACR (mg/mmol)</td>
<td>6.1±29</td>
<td>4.4±19</td>
</tr>
<tr>
<td>Body mass index (kg/m²)</td>
<td>27±4</td>
<td>26±3</td>
</tr>
<tr>
<td>Systolic BP (mmHg)</td>
<td>149±19</td>
<td>151±21</td>
</tr>
<tr>
<td>Antihypertensive treatment, n (%)</td>
<td>394 (48)</td>
<td>365 (47)</td>
</tr>
<tr>
<td>Serum cholesterol (mmol/L)</td>
<td>5.4±1.1</td>
<td>5.4±1.0</td>
</tr>
<tr>
<td>HDL (mmol/L)</td>
<td>1.49±0.46</td>
<td>1.3±0.3</td>
</tr>
<tr>
<td>Lipid-lowering treatment, n (%)</td>
<td>204 (26)</td>
<td>129 (17)</td>
</tr>
<tr>
<td>Smoking, n (%)</td>
<td>50 (6)</td>
<td>59 (8)</td>
</tr>
<tr>
<td>Diabetes, n (%)</td>
<td>112 (14)</td>
<td>107 (14)</td>
</tr>
</tbody>
</table>

Data are mean±SD for continuous variables and n (%) for categorical variables.
The main finding of this study in two community-based cohorts of elderly persons was that sTNFRs are closely associated with the two most relevant clinical markers defining disease stage and progression risk in CKD: GFR and ACR. The association between sTNFR and eGFR was, however, of a much higher magnitude than that between sTNFRs and ACR, even after adjustment for other inflammatory markers and cardiovascular risk factors. It was also more pronounced in participants with GFR < 60 ml/min per 1.73 m² than in participants with GFR ≥ 60 ml/min per 1.73 m². Of note, these associations were also evident in participants without diabetes. The association between sTNFR1 and KIM-1, a marker of tubular damage, was present in both cohorts.

Comparisons with Previous Studies

Recent studies have shown that higher sTNFRs are associated with deterioration of kidney function or progression of microalbuminuria to macroalbuminuria in patients with type 1 diabetes. In this patient group, sTNFRs have been shown to be superior to many inflammatory markers, including IL-6 and TNF-α, as prognostic markers of eGFR decline. Furthermore, higher sTNFRs predict ESRD in type 2 diabetes. In both cohorts in the present study, circulating levels of sTNFRs were significantly higher in participants with diabetes than in those without diabetes. But more important, the association between sTNFRs and the different aspects of kidney damage and dysfunction were also evident in persons without diabetes. Thus, sTNFRs appear to be a marker for kidney pathology also in the absence of diabetes, a finding that to our knowledge has not been reported before. Our data are in accordance with previous community-based studies that have reported the association between sTNFRs, GFR, and albuminuria.

Moreover, we believe we are the first to report an association between sTNFRs and a specific marker of kidney tubular damage, U-KIM-1. In ULSAM, the association between sTNFRs and U-KIM-1 was attenuated and no longer significant after adjustments for eGFR and albuminuria, whereas in PIVUS the association between sTNFRs remained statistically significant after adjustment for eGFR and albuminuria. Thus, whether sTNFRs are independent markers for specific damages in the proximal tubuli remains to be established.

The correlation between sTNFR1 and sTNFR2 in the present PIVUS study was 0.59, confirming high correlations (0.90 and 0.78) seen in a study of patients with type 2 diabetes, 0.90° and a study of patients with type 1 diabetes, respectively. This finding indicates that the strength of association between these two highly correlated markers may depend on the population studied.

Possible Mechanisms for Observed Associations

Several molecular mechanisms may explain our observational findings. Microinflammation driven by interleukins, such as IL-1, IL-6, and IL-18, as well as by TNF-α is directly involved in the pathogenesis and progression of CKD. Our findings, and findings by others, indicate that sTNFRs are closely linked to kidney dysfunction and albuminuria, presumably as direct pathogenic mediators and as markers of high TNF-α activity. The fact that sTNFRs were significant after adjustments for IL-6 and CRP in the present study, as well as in other studies of kidney dysfunction and development of ESRD in individuals with diabetes, further indicates that sTNFRs mirror an independent inflammatory pathway. Specifically, sTNFRs have been shown to be involved in tubulointerstitial fibrosis and thereby contribute to nephropathy. Inflammation identified by sTNFRs may also trigger and promote loss of kidney function due to TNF-driven development of atherosclerosis and malnutrition.

Hyperglycemia has been suggested to affect the levels of oxidative stress and provides one of the main factors explaining the rapid decline in GFR seen in diabetic nephropathy. Oxidative stress also increases TNF-α activity, specifically that of TNF-2. Our study, in contrast to a study of patients with diabetes in which sTNFR1 was more linked to development of ESRD than was TNF-2, showed equally strong associations for both sTNFRs with kidney function and microalbuminuria.

Physical inactivity is accountable for 15/1000 cardiovascular deaths in individuals with CKD and is as important as traditional risk factors, including systolic BP (14/1000 deaths).
Figure 1. Significant correlations between sTNFR1 and the following markers of kidney dysfunction in the elderly: GFR in ULSAM (A), GFR in PIVUS (B), ACR in ULSAM (C), ACR in PIVUS (D), U-KIM-1 in ULSAM (E), and U-KIM-1 in PIVUS (F).
and diabetes (14/1000 deaths). A possible pathway for the increased mortality in sedentary individuals is a higher inflammatory state, which is associated with IL-6 to a greater extent than CRP in patients with CKD. Thus, symptomatic inflammation may decrease ambulation, leading to a vicious circle with more inflammation, progressed CKD, and even less activity. Yet, the present association between sTNFRs and kidney damage and dysfunction remained essentially unaltered in models adjusted for IL-6, CRP, and the level of physical activity, which would argue against this as a major explanation of our findings.

Finally, it is possible that the strong associations between sTNFRs and GFR are partially explained by their impaired renal clearance, but inflammatory mediators are probably the principal cause of their increase in serum that parallels the decline in kidney function.

**CLINICAL IMPLICATIONS**

CKD has a major public impact worldwide, with a global prevalence of 10%. Soluble TNFRs are promising biomarkers of kidney damage and cardiovascular diseases, but more studies are needed to evaluate the clinical value of sTNFR measurements for the detection of kidney damage and for the prediction of GFR decline and the development of cardiovascular disease. The higher correlation between sTNFR and markers of kidney and tubular dysfunction in patients with GFR <60 ml/min per 1.73 m² in the present study indicates that sTNFR measurements may be more clinically relevant in individuals with established CKD. Longitudinal studies of decline of GFR and its association with sTNFR in the community are warranted.

TNF-α inhibition reduces albuminuria in rats. Recombinant antibodies against TNF-α and sTNFR have been suggested as potential drugs that may halt decline in kidney function. In fact, studies have demonstrated that progression of CKD can be inhibited by anti-TNF therapy. Additional clinical studies are warranted to elucidate whether anti-TNF therapy can halt GFR decline and microalbuminuria in patients with rapid nephropathy and high circulating levels of sTNFRs.

**STRENGTHS AND LIMITATIONS**

Strengths of our investigation include the validation of our findings in an independent cohort and the detailed characterization of study participants with regard to kidney phenotypes and cardiovascular risk factors. Limitations include the unknown generalizability to other age and ethnic groups. No conclusions regarding causality should be drawn from our cross-sectional observational data; however, the high associations with eGFR and microalbuminuria in the community are of interest because sTNFRs have been shown to be prognostic markers of kidney dysfunction in patients with diabetes.
In conclusion, circulating sTNFRs were associated with different aspects of kidney damage in two independent community-based cohorts of elderly, even in the absence of diabetes. Our findings confirm and extend previous studies in patients with diabetes to the community-based setting, supporting that circulating sTNFRs are relevant biomarkers for kidney damage and dysfunction, and emphasize the importance of microinflammation as a mechanism underpinning kidney damage and nephropathy.

CONCISE METHODS

Description of Study Populations

PIVUS
All 70-year-old men and women living in Uppsala, Sweden, between 2001 and 2004 were eligible for the PIVUS study (described in detail at http://www.medsci.uu.se/pivus/pivus.htm).29 Of 2025 invited individuals, 1016 agreed to participate. In the present study the second examination cycle of PIVUS was used (2006–2009), when participants were 75 years old. Of 964 invited participants, 827 participated (86%); of these, 815 participants had data on sTNFRs.

ULSAM
ULSAM was initiated in 1970. All 50-year-old men, born in 1920–1924 and living in Uppsala, Sweden, were invited to participate in a health survey that focused on identifying cardiovascular risk factors (described in detail at http://www.pubcare.uu.se/ULSAM).30 These analyses are based on the fourth examination cycle, when participants were approximately 77 years of age (1997–2001). Of 1398 invited men, 838 (60%) participated and 778 participants had valid data on sTNFRs.

Ethical Considerations
All participants in both studies gave written informed consent and the Ethics Committee of Uppsala University approved the study protocols. The study was conducted according to the Declaration of Helsinki.

Baseline Investigations
The investigations in PIVUS and ULSAM were performed using similar standardized methods, including anthropometrical measurements; BP; blood sampling; and questionnaires regarding socioeconomic status, medical history, smoking habits, medication, and physical activity level.29–31 Venous blood samples were drawn in the morning after an overnight fast and stored at −70°C until analysis. In the PIVUS cohort, a spot sample of first-morning-void urine was used for analyses. In ULSAM a 24-hour collection of urine was used.

Both sTNFR1 and sTNFR2 were analyzed using a commercially available ELISA kit (DY225 and DY726; R&D Systems, Minneapolis, MN). The assays had a total coefficient of variation of approximately 6%. Soluble TNFR2 was available only in PIVUS.

Cystatin C–based eGFR was calculated as previously described.32,33 Urine albumin was measured by nephelometry (urine albumin; Dade Behring, Deerfield, IL) using a Behring BN ProSpec analyzer (Dade Behring). Urine creatinine was analyzed with a modified kinetic Jaffe reaction on an Architect Ci8200 analyzer (Abbott, Abbott Park, IL), and creatinine-related ACR was calculated. Urinary KIM-1 was analyzed with the commercial sandwich ELISA kit (DY1750; R&D Systems) and adjusted for urinary creatinine (IL test creatinine 181672–00, Monarch 2000 analyzer; Instrumental Laboratories, Lexington, MA).

High-sensitive CRP measurements were performed by latex-enhanced reagent (Siemens) with the use of a BN ProSpec analyzer (Siemens). IL-6 was analyzed in serum using a high-sensitivity IL-6 assay (Quantikine HS ELISA Kit HS600; R&D Systems) according to the instructions of the manufacturer in ULSAM. Diabetes mellitus was diagnosed as fasting plasma glucose level ≥ 7.0 mmol/l (≥126 mg/dl) or use of antidiabetic medication.34

Statistical Analyses
Pearson correlation coefficients were calculated between sTNFR and markers of kidney function (logarithmically transformed variables to promote normal distribution). Linear regression analyses were used to assess cross-sectional associations with log-transformed sTNFR1 and sTNFR2 (PIVUS only) levels as the dependent variable and other parameters as independent variables. The following multivariable models were used:

A: Age and sex (PIVUS).

B: Inflammation (age, sex [PIVUS], and CRP) to test whether sTNFRs add information to models with the clinically most established inflammatory marker, CRP. Additionally, we adjusted for IL-6 in ULSAM.

Table 4. Association between sTNFR2 and kidney markers in the PIVUS cohort (natural logarithm–transformed variables)

<table>
<thead>
<tr>
<th>Kidney marker</th>
<th>Pearson Correlation Coefficients: All</th>
<th>Linear Regression Models, B Coefficients (95% CI)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Model A: All</td>
<td>Model B: All</td>
</tr>
<tr>
<td>GFR</td>
<td>−0.51*</td>
<td>−0.51* (−0.57 to −0.45)</td>
</tr>
<tr>
<td>ACR</td>
<td>0.17*</td>
<td>0.16* (0.088 to 0.22)</td>
</tr>
<tr>
<td>U-KIM-1</td>
<td>0.090*</td>
<td>0.093* (0.024–0.16)</td>
</tr>
</tbody>
</table>

Model A: age and sex (PIVUS). Model B: inflammation (age, sex [PIVUS], CRP, and IL-6 [ULSAM]). Model C: CVD risk factors (age, sex [PIVUS], CRP, IL-6 [ULSAM], body mass index, smoking, systolic BP, HDL, cholesterol, diabetes, and antihypertensive and lipid treatment).

*P<0.001.

bP<0.01.

cP<0.05.
C: Established cardiovascular disease risk factors (age, sex [PIVUS], CRP, IL-6 [ULSAM], systolic BP, cholesterol, HDL, body mass index, lipid-lowering and antihypertensive treatment) to test whether the associations between sTNFRs are independent of CVD risk factors that may promote kidney damage.

Because of non-normal distributions, Spearman correlation coefficients were used to calculate the correlation between sTNFR1 and sTNFR2 in PIVUS (Spearman correlation was also used so that we could compare our data with those in other studies that used Spearman for this analysis).

In secondary models, we also adjusted for leisure time physical activity. We also used a nonparametric metric method to calculate the differences in medians of sTNFRs and between individuals with and without diabetes due to non-normal distributions, by the use of their Bonnett–Price CIs. Stratified multivariable models (model A) in patients with and without diabetes were also calculated. Finally, we also investigated the association between circulating sTNFR2 and markers of kidney damage in PIVUS using the same multivariable linear regression models.

The statistical software package Stata 11.2 (Stata Corp., College Station, TX) was used.

ACKNOWLEDGMENTS

This study was supported by The Swedish Research Council, Swedish Heart-Lung Foundation, Thuréus Foundation, the Marianne and Marcus Wallenberg Foundation, Dalarna University, Upssala-Orebro Regional Research Council, and Uppsala University. The funding sources did not play any role in the design and conduct of the study, collection, management, analysis, and interpretation of the data, and preparation, review, or approval of the manuscript. Dr. Arnlov is the guarantor of this work, had full access to all the data, and takes full responsibility for the integrity of data and the accuracy of data analysis.

DISCLOSURES

None.

REFERENCES


Soluble TNFRs and Kidney Dysfunction

www.jasn.org CLINICAL EPIDEMIOLOGY


This article contains supplemental material online at http://jasn.asnjournals.org/lookup/suppl/doi:10.1681/ASN.2013080860/-/DCSupplemental.