

# Proteinuria and its association with disease progression in IgA nephropathy: Analysis of the UK national RaDaR IgA nephropathy cohort

**David Pitcher<sup>1</sup>, Fiona Braddon<sup>1</sup>, Bruce Hendry<sup>2</sup>, Alex Mercer<sup>3</sup>, Kate Osmaston<sup>1</sup>, Moin A. Saleem<sup>4</sup>, Retha Steenkamp<sup>1</sup>, Neil Turner<sup>5</sup>, Kaijun Wang<sup>2</sup>, Jonathan Barratt<sup>6</sup>, Daniel P. Gale<sup>7</sup>**

<sup>1</sup>UK Kidney Association; <sup>2</sup>Travere Therapeutics, Inc., San Diego, CA; <sup>3</sup>JAMCO Pharma Consulting, Sweden; <sup>4</sup>University of Bristol & Bristol Royal Hospital for Children, UK; <sup>5</sup>University of Edinburgh, UK; <sup>6</sup>University of Leicester & Leicester General Hospital, UK; <sup>7</sup>Department of Renal Medicine, University College London, UK

Contact information:  
David Pitcher, david.pitcher@renalregistry.nhs.uk

Presented at the American Society of Nephrology (ASN) Kidney Week 2022;  
November 03 – November 06, 2022; Orlando, Florida

To obtain a PDF of this poster:



Scan the QR code OR visit  
[www.traverepublications.com/ASN2022/TH-PO494](http://www.traverepublications.com/ASN2022/TH-PO494)

Charges may apply.

No personal information is stored.

**RaDaR**

**RareRenal**  
Information on rare kidney diseases

**UKKA**  
UK Kidney Association  
Rare Renal

**TRAVERE**  
THERAPEUTICS

## Disclosures

- **DP** has nothing to disclose. **FB...** **BH** is an employee and stockholder of Traverre Therapeutics, Inc. **AM** received consultancy fees from Traverre Therapeutics, Inc. **KO...** **MAS** received consultancy fees from Traverre Therapeutics, Inc. and Purespring Therapeutics. **RS** has nothing to disclose. **NT...** **KW** is an employee and stockholder of Traverre Therapeutics, Inc. **JB** received consultancy fees from Traverre Therapeutics, Inc. **DPG** received consultancy fees from Traverre Therapeutics, Inc.

## Acknowledgements

- This study was funded by Traverre Therapeutics, Inc. Writing support was provided by Eve Hunter-Featherstone and David Cork of Genesis Research (Newcastle upon Tyne, UK) which received compensation from Traverre Therapeutics.

- Primary IgA nephropathy (IgAN) is the most common form of glomerulonephritis worldwide and a major cause of kidney failure (KF)<sup>1,2</sup>
- Rate of progression to KF varies widely and can span over decades<sup>3</sup>
- Time-averaged proteinuria (TA-PU) over long-term follow-up is an important predictor of disease progression and KF risk in patients with IgAN<sup>3,4</sup>

## Objective

- To investigate the relationship between proteinuria (PU) measured over follow-up (TA-PU) and rate of kidney function loss and kidney survival in UK IgAN patients within the UK National Registry of Rare Kidney Diseases (RaDaR)

**Abbreviations:** IgAN, IgA nephropathy; KF, kidney failure; PU, proteinuria; TA-PU, time-averaged proteinuria;

**1.** Canney M, et al. *J. Am. Soc. Nephrol.* 2021;32(2):436-447. **2.** McGrogran A, et al. *Nephrol. Dial. Transplant.* 2011;26(2):414-430. **3.** Le W, et al. *Nephrol. Dial. Transplant.* 2012;27(4):1479-1485. **4.** Reich HN, et al. *J Am Soc Nephrol.* 2007;18(12):3177-3183.

## Data Source

- This study uses data from the RaDaR database
- Since 2013, patients with biopsy-proven IgAN and estimated glomerular filtration rate (eGFR)  $<60$  mL/min/1.73 m<sup>2</sup> or PU  $\geq 0.5$  g/day have been enrolled into (RaDaR) IgAN Cohort
- RaDaR contains data on IgAN patients from 87 kidney units across the UK, with automated collection of retrospective and prospective laboratory data

## Definitions and Clinical Measures

- Diagnosis was the earliest of either primary kidney diagnosis date or date of biopsy recorded in RaDaR
- eGFR calculated via the Chronic Kidney Disease Epidemiology Collaboration (CKD-EPI) formula<sup>5</sup> (adults) and the modified Schwartz formula<sup>6</sup> (pediatric)
- KF was defined as the first occurrence of either chronic kidney replacement therapy, a confirmed eGFR  $<15$  mL/min/1.73 m<sup>2</sup>, or KF/CKD stage 5 recorded in RaDaR
- TA-PU was defined as the time-weighted averages for urinary protein-creatinine ratio (PCR), calculated from the area under the curve of serial measurements divided by the length of follow-up

**Abbreviations:** **CKD**, chronic kidney disease; **CKD-EPI**, Chronic Kidney Disease Epidemiology Collaboration; **eGFR**, estimated glomerular filtration rate; **IgAN**, IgA nephropathy; **KF**, kidney failure; **TA-PU**, time-averaged proteinuria; **PCR**, urinary protein-creatinine ratio

**5.** Levey AS, et al. *Ann Intern Med.* 2009;150(9):604-612. **6.** Schwartz GJ, et al. *J Am Soc Nephrol.* 2009;20(3):629-637.

## Eligibility Criteria

- Patients were included if they had a biopsy date recorded in RaDaR and PU measurements in follow-up (within 2 years from diagnosis and  $\geq 2$  values if follow-up  $> 3$  years)
- Patients were excluded if they had a KF event (CKD stage 5 or kidney replacement therapy [KRT]) or death within 6 months from diagnosis or prior to first PU value

## Statistical Analyses

- TA-PU and rate of eGFR loss (eGFR slope) were calculated over the full duration of follow-up or until KF or death. A linear mixed model was used to estimate each patient's intercept and slope of eGFR
- Kaplan-Meier estimates for kidney survival, from diagnosis to KF/death, were calculated for each TA-PU group. The log-rank test was used for differences between pairwise and all groups
- Association of TA-PU and survival from KF/death evaluated using Cox regression

## Characteristics at diagnosis and clinical outcomes

- The cohort included 923 patients (68% male and 96% adult) with a median age at diagnosis of 41.7 years
- Among patients with available data, the median urinary PCR at diagnosis was 172 mg/mmol (1.5 g/g) and median eGFR was 50 mL/min/1.73 m<sup>2</sup>
- Median duration of follow-up was 4.5 years and 38% of patients progressed to KF/death during follow-up
- Mean eGFR slope was -3.6 mL/min/1.73m<sup>2</sup>/year

**Table 1. Characteristics at diagnosis and clinical outcomes**

	N	%
<b>Age (years)</b>	<b>923</b>	<b>100</b>
Median (IQR)	41.7 (30.3–53.3)	
Pediatric	36	4
<b>Sex</b>	<b>923</b>	<b>100</b>
Female	294	32
Male	629	68
<b>PCR at baseline, n (%)</b>	<b>515</b>	<b>56</b>
mg/mmol Median (IQR)	172 (73–356)	
g/g Median (IQR)	1.5 (0.6–3.1)	
<b>eGFR at baseline, n (%)</b>	<b>565</b>	<b>61</b>
Median, mL/min/1.73 m <sup>2</sup>	49.8	
IQR	33.0–78.2	
<b>Duration of follow-up, n (%)</b>	<b>923</b>	<b>100</b>
Median, years	4.5	
IQR	2.5–6.8	
<b>KF or death event, n (%)</b>	<b>923</b>	<b>100</b>
Yes	355	38
No	568	62
<b>eGFR slope, n (%)</b>	<b>856</b>	<b>93</b>
Mean, mL/min/1.73 m <sup>2</sup> /year	-3.6	
SD	9.4	

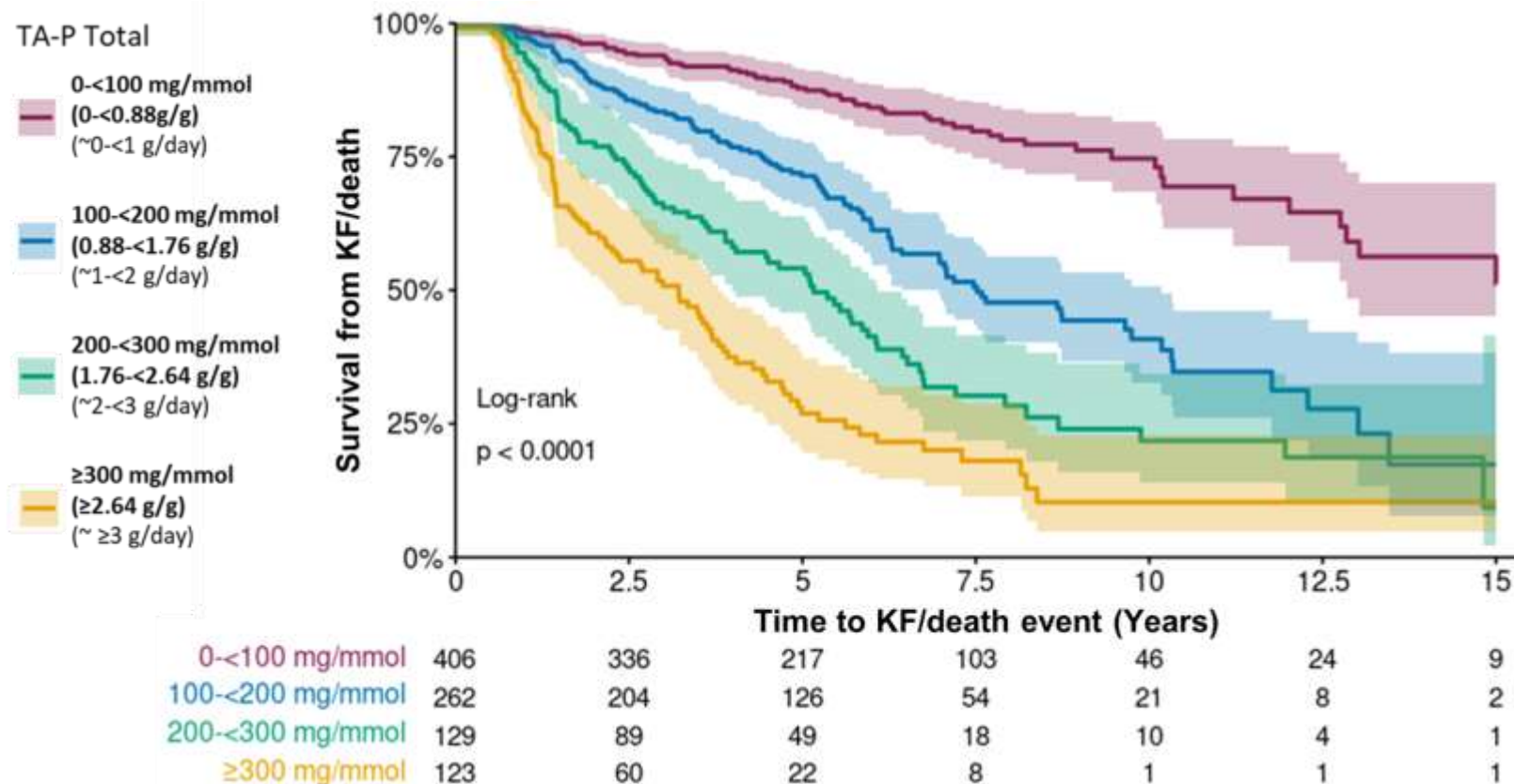
**Abbreviations:** eGFR, estimated glomerular filtration rate; IQR, interquartile range; KF, kidney failure; PCR, protein-creatinine ratio; SD, standard deviation

100 mg/mmol = 0.88 g/g ≈ 1 g/day

## Elevated time-averaged proteinuria is associated with kidney failure/death

- Time to KF/death was significantly shorter with higher levels of TA-PU
- Approx. 1 in 4 patients with TA-PU <100 mg/mmol progressed to KF/death within 10 years

**Figure 1. Kaplan Meier survival curves for patients categorized by TA-PU**



## Elevated TA-PU is associated with kidney failure/death and more rapid loss of eGFR

- TA-PU 100–<200 mg/mmol was associated with an almost 3-fold increase in risk of KF/death compared with TA-PU <100 mg/mmol
- The risk of KF/death was increased almost 5-fold at TA-PU 200–<300 mg/mmol compared with TA-PU <100 mg/mmol
- TA-PU ≥300 mg/mmol was associated with a 9-fold increase in the risk of KF/death compared with TA-PU <100 mg/mmol
- Higher grades of TA-PU were associated with a higher rate of eGFR loss ( $p < 0.001$ ). The rate of eGFR loss escalated from an eGFR slope of  $-0.35 \text{ mL/min/1.73 m}^2/\text{year}$  for TA-PU <100 mg/mmol to  $-12.41 \text{ mL/min/1.73 m}^2/\text{year}$  with TA-PU ≥300 mg/mmol

**Table 2. Clinical outcomes for patients categorized by TA-PU**

TA-PU	eGFR slope (mL/min/1.73 m <sup>2</sup> /year)			KF/death risk		
	N	Mean	SD	N	HR	95% CI
<100 mg/mmol	385	-0.35	7.15	405	Ref	Ref
100 to <200 mg/mmol	247	-3.32	10.09	264	2.83	2.09-3.82
200 to <300 mg/mmol	113	-6.67	5.73	128	4.82	3.49-6.66
≥300 mg/mmol	111	-12.41	11.28	126	9.00	6.56-12.34



- Adults represent >90% of the cohort with a median baseline age of 41.7 years reflecting a disease with onset at a stage when patients should have a long life-expectancy remaining
- Higher grades of TA-PU are significantly associated with shorter time to KF/death and increased KF/death risk
- Higher grades of TA-PU are also significantly associated with more rapid loss of eGFR
- Proteinuria <1 g/day is commonly perceived as defining patients at low risk, however, in this cohort, approx. 1 in 4 patients progressed to KF/death within 10 years, despite a TA-PU of <100 mg/mmol (approximately <1 g/day)

## Limitations

- The inclusion criteria for RaDaR-IgAN leads to enrollment of patients with progressive disease who represent a high risk IgAN population
- Reporting of proteinuria and eGFR data at disease onset is incomplete and may not be representative of the full cohort, however data are likely to be missing at random with limited bias

- Elevated proteinuria over time is significantly associated with rapid loss of eGFR and greater risk of progression to KF/death in IgAN
- Although TA-PU below 100 mg/mmol is strongly associated with lower risk of KF/death, 25% of patients in this treated, monitored group reached KF/death within 10 years