

# Acute kidney injury in patients treated with immune checkpoint inhibitors

Shruti Gupta <sup>1</sup>, Samuel A P Short,<sup>2</sup> Meghan E Sise,<sup>3</sup> Jason M Prosek,<sup>4</sup> Sethu M Madhavan,<sup>4</sup> Maria Jose Soler,<sup>5</sup> Marlies Ostermann,<sup>6</sup> Sandra M Herrmann,<sup>7</sup> Ala Abudayyeh,<sup>8</sup> Shuchi Anand,<sup>9</sup> Ilya Glezerman,<sup>10</sup> Shveta S Motwani,<sup>11</sup> Naoka Murakami,<sup>1</sup> Rimda Wanchoo,<sup>12</sup> David I Ortiz-Melo,<sup>13</sup> Arash Rashidi,<sup>14</sup> Ben Sprangers,<sup>15,16</sup> Vikram Aggarwal,<sup>17</sup> A Bilal Malik,<sup>18</sup> Sebastian Loew,<sup>19</sup> Christopher A Carlos,<sup>20</sup> Wei-Ting Chang,<sup>21,22,23</sup> Pazit Beckerman,<sup>24</sup> Zain Mithani,<sup>25</sup> Chintan V Shah,<sup>26</sup> Amanda D Renaghan,<sup>27</sup> Sophie De Seigneux,<sup>28</sup> Luca Campedel,<sup>29</sup> Abhijat Kitchlu <sup>30</sup>, Daniel Sanghoon Shin,<sup>31</sup> Sunil Rangarajan,<sup>32</sup> Priya Deshpande,<sup>33</sup> Gaia Coppock,<sup>34</sup> Mark Eijgelsheim,<sup>35</sup> Harish Seethapathy,<sup>3</sup> Meghan D Lee <sup>3</sup>, Ian A Strohbehm,<sup>3</sup> Dwight H. Owen,<sup>36</sup> Mariam Husain,<sup>36</sup> Clara Garcia-Carro,<sup>5,37</sup> Sheila Bermejo,<sup>5</sup> Nuttha Lumlertgul,<sup>38,39</sup> Nina Seylanova,<sup>38,40</sup> Lucy Flanders,<sup>41</sup> Busra Isik,<sup>7</sup> Omar Mamlouk <sup>8</sup>, Jamie S Lin <sup>8</sup>, Pablo Garcia,<sup>9</sup> Aydin Kaghazchi,<sup>42</sup> Yuriy Khanin,<sup>12</sup> Sheru K Kansal,<sup>14</sup> Els Wauters,<sup>43,44</sup> Sunandana Chandra,<sup>45</sup> Kai M Schmidt-Ott,<sup>19,46</sup> Raymond K Hsu,<sup>20</sup> Maria C Tio,<sup>1</sup> Suraj Sarvode Mothi,<sup>1</sup> Harkarandeep Singh,<sup>1</sup> Deborah Schrag,<sup>47</sup> Kenar D Jhaveri,<sup>12</sup> Kerry L Reynolds,<sup>48</sup> Frank B Cortazar,<sup>49</sup> David E Leaf <sup>1</sup>, ICPi-AKI Consortium Investigators

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For numbered affiliations see end of article.

## Correspondence to

Dr Shruti Gupta;  
sgupta21@bwh.harvard.edu

## ABSTRACT

**Background** Immune checkpoint inhibitor-associated acute kidney injury (ICPi-AKI) has emerged as an important toxicity among patients with cancer.

**Methods** We collected data on 429 patients with ICPi-AKI and 429 control patients who received ICPis contemporaneously but who did not develop ICPi-AKI from 30 sites in 10 countries. Multivariable logistic regression was used to identify predictors of ICPi-AKI and its recovery. A multivariable Cox model was used to estimate the effect of ICPi rechallenge versus no rechallenge on survival following ICPi-AKI.

**Results** ICPi-AKI occurred at a median of 16 weeks (IQR 8–32) following ICPi initiation. Lower baseline estimated glomerular filtration rate, proton pump inhibitor (PPI) use, and extrarenal immune-related adverse events (irAEs) were each associated with a higher risk of ICPi-AKI. Acute tubulointerstitial nephritis was the most common lesion on kidney biopsy (125/151 biopsied patients [82.7%]). Renal recovery occurred in 276 patients (64.3%) at a median of 7 weeks (IQR 3–10) following ICPi-AKI. Treatment with corticosteroids within 14 days following ICPi-AKI diagnosis was associated with higher odds of renal recovery (adjusted OR 2.64; 95% CI 1.58 to 4.41). Among patients treated with corticosteroids, early initiation of corticosteroids (within 3 days of ICPi-AKI) was associated with a higher odds of renal recovery compared with later initiation (more than 3 days following ICPi-AKI) (adjusted OR 2.09; 95% CI 1.16 to 3.79). Of 121 patients rechallenged, 20 (16.5%) developed recurrent ICPi-AKI. There was no difference in survival among patients rechallenged versus those not rechallenged following ICPi-AKI.

**Conclusions** Patients who developed ICPi-AKI were more likely to have impaired renal function at baseline, use a PPI, and have extrarenal irAEs. Two-thirds of patients had renal recovery following ICPi-AKI. Treatment with corticosteroids was associated with improved renal recovery.

## BACKGROUND

Immune checkpoint inhibitors (ICPis) have become some of the most widely prescribed anticancer treatments in current use.<sup>1</sup> Despite their proven efficacy across a wide range of malignancies, ICPis can cause a unique spectrum of autoimmune toxicities known as immune-related adverse events (irAEs). These irAEs can affect virtually any organ in the body, including the kidneys.

Direct renal toxicity from ICPis, referred to here as ICPi-associated acute kidney injury (ICPi-AKI), occurs with an estimated incidence of 3%–5%.<sup>2–9</sup> ICPi-AKI can have serious consequences for patients including dose delay or permanent discontinuation of ICPi therapy, irreversible loss of kidney function (which can impact eligibility to receive other anticancer treatments), and prolonged courses of immunosuppression. Additionally, when a patient undergoing treatment with ICPis develops AKI, there is often uncertainty regarding its etiology, since AKI occurs

commonly in patients with cancer and can be due to a variety of causes.<sup>10 11</sup>

Despite these challenges in diagnosing and treating ICPI-AKI, existing data are largely limited to case reports and small single-center case series.<sup>5 6 12-15</sup> Key questions, therefore, remain unanswered regarding the risk factors, clinical features, histopathological findings, renal outcomes, and overall survival in patients with ICPI-AKI. Further, very limited data are available regarding the safety of rechallenging patients with ICPIs after an episode of ICPI-AKI. To address these and other critical knowledge gaps, we conducted an international multicenter cohort study of patients with ICPI-AKI.

## METHODS

### Study design and oversight

We conducted a multicenter cohort study of adults diagnosed with ICPI-AKI between 2012 and 2020. We contacted nephrologists and oncologists at 40 major academic cancer centers across North America, Europe, and Asia to identify cases of ICPI-AKI, 30 of which provided data on 429 patients with ICPI-AKI (online supplemental table S1 and figure S1). Of the 429 patients with ICPI-AKI in this study, 100 (23.3%) were described previously in 12 publications (online supplemental appendix).

### Data collection

Study personnel at each site collected data by detailed chart review. Data were entered into a standardised case report form using a secure, web-based platform (online supplemental appendix),<sup>16</sup> and included the following: demographics and comorbidities; concomitant treatment with nephrotoxic chemotherapies and medications associated with acute tubulointerstitial nephritis (ATIN), including proton pump inhibitors (PPIs), non-steroidal anti-inflammatory drugs, and antibiotics; prior or concomitant extrarenal irAEs; laboratory data at baseline and at the time of ICPI-AKI; kidney biopsy data; treatments received for ICPI-AKI; and data on renal recovery, ICPI rechallenge, and overall survival.

### Definition of ICPI-AKI

Patients were eligible for inclusion if they had AKI that was directly attributed to the ICPI by the treating provider and if they met either of the following criteria: (1) an increase in serum creatinine (SCr)  $\geq 100\%$  from baseline or treatment with renal replacement therapy (RRT); (2) an increase in SCr  $\geq 50\%$  from baseline and at least one of the following: ATIN on kidney biopsy; ICPI therapy held for at least once cycle due to concern for ICPI-AKI; or treatment with corticosteroids due to concern for ICPI-AKI (online supplemental table S2). Baseline SCr was defined as the closest value prior to ICPI initiation. We excluded kidney transplant patients and those with end stage kidney disease.

### Definitions of AKI severity, renal recovery, and recurrent ICPI-AKI

AKI severity was based on the maximum SCr achieved in the 4 weeks following ICPI-AKI and staged according

to the Kidney Disease: Improving Global Outcomes criteria (online supplemental table S3).<sup>17</sup> Renal recovery was defined as a nadir SCr  $\leq 1.5$  times the baseline value within 90 days following ICPI-AKI.<sup>18</sup> Recurrent ICPI-AKI after ICPI rechallenge was defined as an increase in SCr  $\geq 50\%$  from the new baseline (at the time of rechallenge) and attributed to the ICPI by the treating provider.

### Patients without ICPI-AKI

To identify risk factors for ICPI-AKI, we also collected data on control patients who received ICPIs contemporaneously but who did not develop ICPI-AKI (defined as absence of an increase in SCr  $\geq 50\%$  from baseline or treatment with RRT that was definitely or probably ICPI related). Each collaborating institution provided data on one control patient for every patient with ICPI-AKI from that site to maintain a 1:1 ratio of cases:controls. Control patients were selected at random to preserve the ability to investigate all characteristics as potential risk factors for ICPI-AKI.

### Statistical analyses

Continuous and categorical data were compared using the Wilcoxon rank-sum and Fisher's exact test, respectively. Multivariable logistic regression was used to identify risk factors for ICPI-AKI, and a sensitivity analysis was performed in patients with stage 2 or 3 ICPI-AKI.

Multivariable logistic regression was also used to identify predictors of renal recovery in patients with ICPI-AKI, including treatment with corticosteroids within 14 days following ICPI-AKI. A sensitivity analysis limited to patients treated with corticosteroids at any time following ICPI-AKI was conducted to assess the effect of early corticosteroid initiation (within 3 days following ICPI-AKI) versus later corticosteroid initiation (anytime after 3 days following ICPI-AKI) on renal recovery. To minimize the potential for confounding due to terminal illness, which could affect the decision to prescribe corticosteroids, each of these analyses were limited to patients who survived at least 14 days following ICPI-AKI.

Kaplan-Meier curves and multivariable Cox regression models were used to assess the association between ICPI-AKI stage, treatment with corticosteroids within 14 days following ICPI-AKI, and other factors with survival, with ICPI-AKI diagnosis serving as time 0. Similar to the analyses above, these analyses were also limited to those who survived at least 14 days following ICPI-AKI.

Kaplan-Meier curves and multivariable Cox regression models were used to estimate the effect of ICPI rechallenge versus no ICPI rechallenge on survival. To eliminate the potential for immortal time bias, we limited this analysis to patients who survived at least 90 days after the initial ICPI-AKI event, and we compared the survival of patients rechallenged in the first 90 days to those not rechallenged in the first 90 days. We repeated this analysis in patients who survived at least 180 days following the initial ICPI-AKI event.

**Table 1** Baseline characteristics

Variable	ICPi-AKI (n=429)	No ICPI-AKI (n=429)	P value
Age at ICPI initiation, years, median (IQR)	68 (59–75)	65 (58–73)	0.02
Male, n (%)	266 (62.0)	251 (58.5)	0.32
Race, n (%)			0.99
White	351 (81.8)	350 (81.6)	
Black	27 (6.3)	24 (5.6)	
Asian	21 (4.9)	21 (4.9)	
Other/unknown	30 (7.0)	34 (7.9)	
Comorbidities, n (%)			
Hypertension	251 (58.5)	229 (53.4)	0.15
Diabetes	77 (17.9)	61 (14.2)	0.16
CHF	17 (4.0)	9 (2.1)	0.16
COPD	45 (10.5)	46 (10.7)	0.99
Cirrhosis	11 (2.6)	10 (2.3)	0.99
Body mass index, median (IQR)	26 (23–30)	26 (22–29)	0.12
Baseline SCr, mg/dL, median (IQR)	0.97 (0.80–1.21)	0.88 (0.73–1.07)	<0.001
Baseline eGFR,* (mL/min per 1.73 m <sup>2</sup> )			
Median (IQR)	73 (57–90)	83 (66–97)	<0.001
eGFR categories, n (%)			<0.001
≥90	111 (25.9)	168 (39.2)	
60–89	192 (44.8)	189 (44.1)	
45–59	72 (16.8)	44 (10.3)	
30–44	43 (10.0)	23 (5.4)	
<30	11 (2.6)	5 (1.2)	
Autoimmune disease, n (%)	42 (9.8)	56 (13.1)	0.16
Extrarenal irAE, † n (%)	201 (46.9)	123 (28.7)	<0.001
Malignancy, n (%)			0.01
Melanoma	104 (24.2)	93 (21.7)	
Lung	126 (29.4)	133 (31.0)	
Genitourinary	100 (23.8)	70 (16.7)	
Other	99 (23.6)	133 (31.7)	
PPI, ‡ n (%)	208 (45.5)	115 (26.8)	<0.001
Concomitant nephrotoxic chemotherapy, § n (%)			
Cisplatin	7 (1.6)	NA	
VEGF/TKI	23 (5.4)	NA	
Other¶	43 (10.0)	NA	
ICPi class, n (%)			
Anti-CTLA-4	103 (24.0)	95 (22.1)	0.57
Anti-PD-1	347 (80.9)	355 (82.8)	0.54
Anti-PD-L1	42 (9.8)	30 (7.0)	0.18
Combo anti-CTLA-4+ anti-PD-1/PD-L1	99 (23.1)	75 (17.5)	0.05

Data are shown as median (IQR) and n (%).

Data on body mass index are missing in one patient with ICPI-AKI and one without ICPI-AKI. Data on PPI use are missing in two patients without ICPI-AKI. All other data are complete.

\*Baseline eGFR calculated based on Chronic Kidney Disease-Epidemiology Collaboration equation.<sup>33</sup>

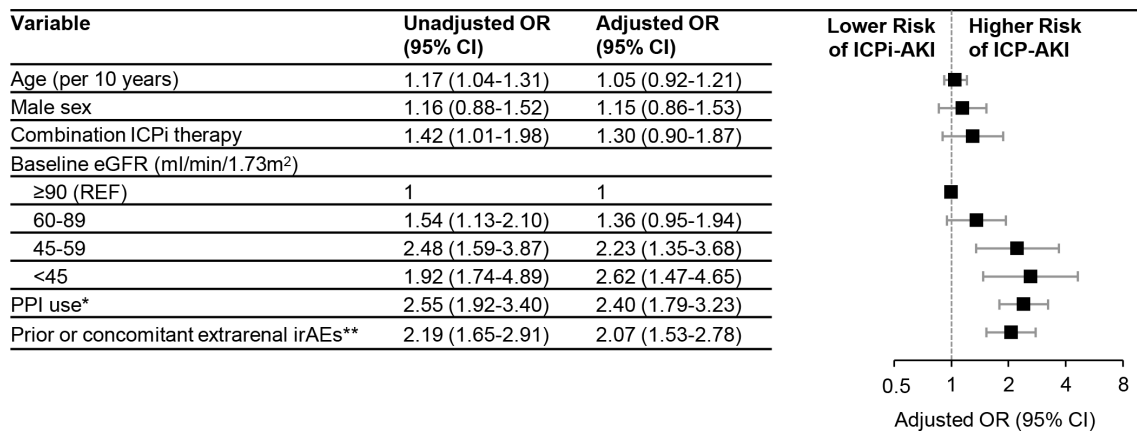
†Extrarenal irAEs were assessed prior to (>14 days) or concomitant (within 14 days before or after) with ICPI-AKI diagnosis among patients with ICPI-AKI, and at any time after ICPI initiation among patients without ICPI-AKI.

‡PPIs were assessed in the 14 days preceding AKI among patients with ICPI-AKI, and were assessed at ICPI initiation in patients without ICPI-AKI.

§Concomitant chemotherapies were assessed in the 30 days preceding ICPI-AKI.

¶Includes pemetrexed (n=28), carboplatin (n=23), BRAF inhibitors (n=2), and paclitaxel (n=1).

AKI, acute kidney injury; BRAF, v-raf murine sarcoma viral oncogene homolog B1; CHF, congestive heart failure; Combo, combination therapy; COPD, chronic obstructive pulmonary disease; CTLA-4, cytotoxic T lymphocyte-associated antigen 4; eGFR, estimated glomerular filtration rate; ICPI, immune checkpoint inhibitor; irAEs, immune-related adverse events; NA, not assessed; PD-1, programmed cell death 1; PD-L1, programmed death-ligand 1; PPI, proton pump inhibitor; SCr, serum creatinine; TKI, tyrosine kinase inhibitor; VEGF, vascular endothelial growth factor.



**Figure 1** Risk factors for ICPI-AKI. Total n=856, of whom 429 had ICPI-AKI and 427 did not have ICPI-AKI. All model covariates are shown in the figure. \*Denotes PPI use in the 14 days preceding ICPI-AKI among those with ICPI-AKI, and PPI use at the time of ICPI initiation among patients without ICPI-AKI. \*\*Extrarenal irAEs were assessed prior to (>14 days) or concomitant (within 14 days before or after) with ICPI-AKI diagnosis among patients with ICPI-AKI, and at any time after ICPI initiation among patients without ICPI-AKI. eGFR, estimated glomerular filtration rate; ICPI, immune checkpoint inhibitor; irAEs, immune-related adverse events; PPI, proton pump inhibitor.

For each of the multivariable analyses above, covariate selection was based on univariate associations, biological plausibility, prior knowledge,<sup>3 4</sup> and parsimony. Missing data were not imputed, as less than 1% of data were missing for all variables. Rather, missing data categories were used in multivariable models. Analyses were performed using SAS V.9.5 (SAS Institute), R V.3.6.3 (R Foundation), and GraphPad PRISM V.9.1.0 (GraphPad Software). All comparisons are two tailed, with  $p < 0.05$  considered significant.

## RESULTS

### Baseline characteristics

In total, 429 patients with ICPI-AKI and 429 without ICPI-AKI from 30 institutions were included. Baseline characteristics are shown in table 1. Patients with ICPI-AKI were older, had lower baseline estimated glomerular filtration rate (eGFR), and were more likely to have genitourinary cancer, PPI exposure, and to have received combination ICPI therapy compared with patients without ICPI-AKI (table 1). The temporal distribution of ICPI initiation was similar in patients with and without ICPI-AKI (online supplemental figure S2). Baseline characteristics among biopsied (n=151) and non-biopsied patients (n=278) with ICPI-AKI were largely similar, though biopsied patients were more likely to have more severe AKI and less likely to have extrarenal irAEs (online supplemental table S4).

### Risk factors for ICPI-AKI

Lower baseline eGFR, PPI use, and prior or concomitant extrarenal irAEs were each associated with a higher risk of ICPI-AKI in multivariable models (figure 1). When limited to patients with stage 2 or 3 ICPI-AKI, PPI use and extrarenal irAEs remained associated with ICPI-AKI, whereas baseline eGFR did not (online supplemental figure S3).

### Clinical features of ICPI-AKI

ICPI-AKI developed at a median of 16 weeks (IQR 8–32) after ICPI initiation, with 49 of 429 patients (11.4%) developing ICPI-AKI more than a year after ICPI initiation (figure 2A). Patients developed ICPI-AKI at a median of 3 weeks (IQR 1–4) after the last ICPI cycle (figure 2B). A total of 77 of the 429 patients (17.9%) with ICPI-AKI had stage 1 AKI, 144 (33.6%) had stage 2, and 208 (48.5%) had stage 3, including 33 who received RRT (15.8% of those with stage 3 AKI; 7.7% overall) (figure 2C). SCr at baseline, at ICPI-AKI diagnosis, the peak level within 4 weeks of ICPI-AKI, and the nadir within 90 days following ICPI-AKI are shown in figure 2D. Prior or concomitant extrarenal irAEs occurred in 243 patients (56.6%), with rash and hepatitis being the most common (figure 2E). Among patients who underwent kidney biopsy, ATIN was the most common primary lesion (125/151; 82.7%) (figure 2F).

Additional clinical features of ICPI-AKI stratified by AKI stage are shown in figure 2G–L. At the time of ICPI-AKI, 62% of patients were receiving concomitant medications associated with ATIN, with PPIs being the most common (figure 2G); 39.7% had  $\geq 1+$  blood on urinalysis (figure 2H); 41.3% had  $\geq 1+$  leukocyte esterase on urinalysis (figure 2I); 56.2% had pyuria ( $\geq 5$  white blood cells per high power field) (figure 2J); 58.7% had a urine protein-to-creatinine ratio  $\geq 0.3$  g/g (figure 2K); and 16.5% had  $\geq 500$  eosinophils/ $\mu$ L (figure 2L). Patients with higher stages of AKI were more likely to be receiving concomitant medications associated with ATIN and to have hematuria, proteinuria, and leukocyte esterase on urinalysis.

### Treatment of ICPI-AKI

ICPI therapy was held for at least one cycle in 390 of the 429 patients (90.9%) with ICPI-AKI. Most patients (350/429 [81.6%]) were treated with corticosteroids (figure 3A), including 100 patients treated with intravenous pulse



dose corticosteroids (figure 3B). Patients were initiated on corticosteroids at a median of 4 days (IQR 1–13) following ICPI-AKI diagnosis, and 273 of the 350 patients treated with corticosteroids (78.0%) initiated this treatment in the first 14 days following ICPI-AKI (figure 3C). The median initial corticosteroid dose was 60 mg in prednisone equivalent units (IQR 50–80) (figure 3D). Patients were treated with corticosteroids for a median of 41 days (IQR 26–75) before tapering to  $\leq 10$  mg of prednisone (or the equivalent). A total of 22 patients (5.1%) were treated with additional or alternative immunosuppressive agents, most commonly mycophenolate mofetil (online supplemental table S5).

### Renal recovery

Renal recovery occurred in 276 patients (64.3%) overall, including 90.9% of patients who initially had ICPI-AKI stage 1, 70.8% with stage 2, and 50.0% with stage 3 (figure 4A). Renal recovery occurred at a median of 7 weeks (IQR 3–10) following ICPI-AKI diagnosis (figure 4B). The characteristics of patients with versus without renal recovery are shown in online supplemental table S6). In a multivariable model, higher baseline eGFR, lung cancer, and stage 3 AKI were each associated with a lower odds of renal recovery, whereas concomitant treatment with ATIN-causing medications was associated with higher odds of renal recovery (figure 4C). Importantly,

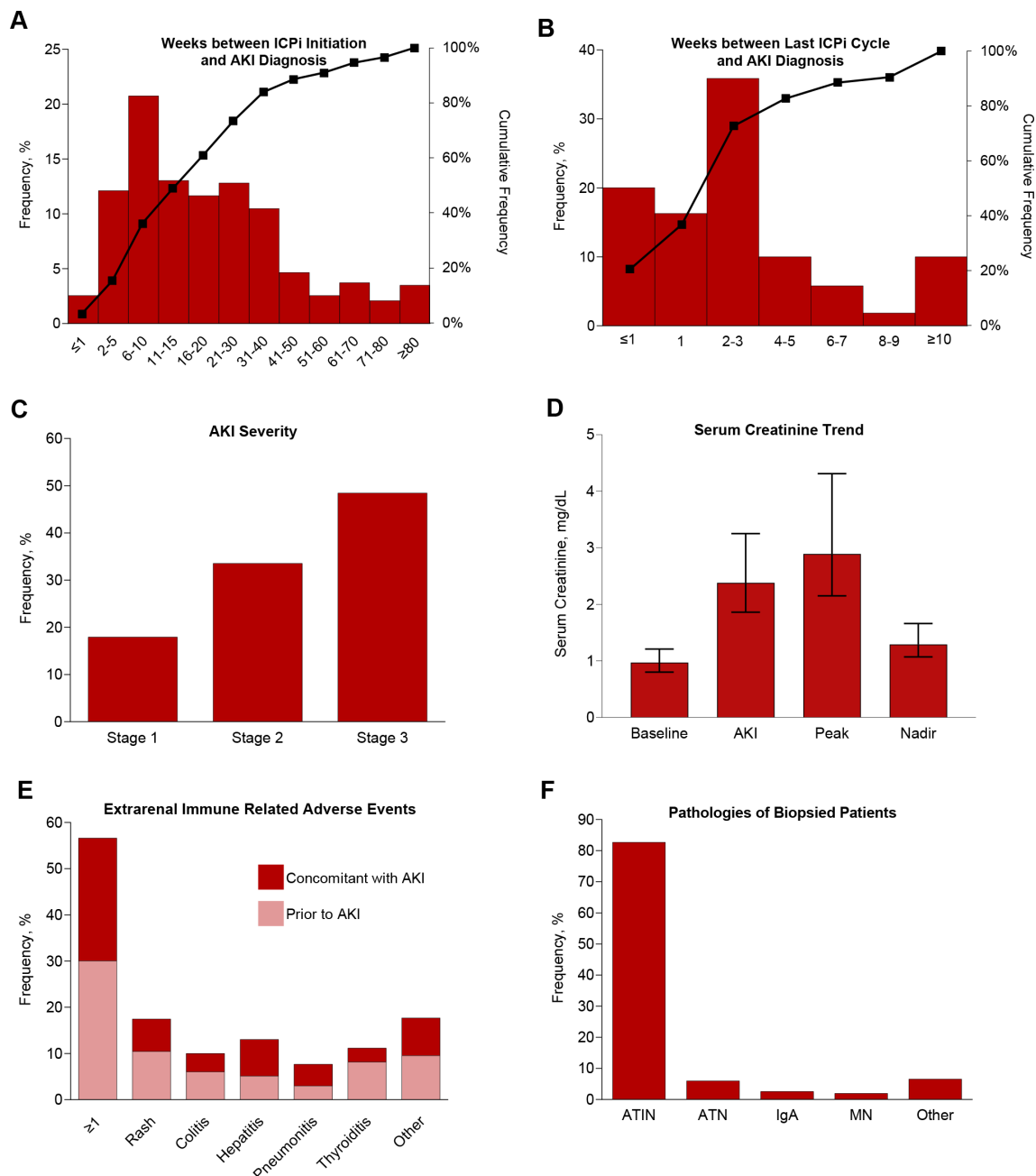
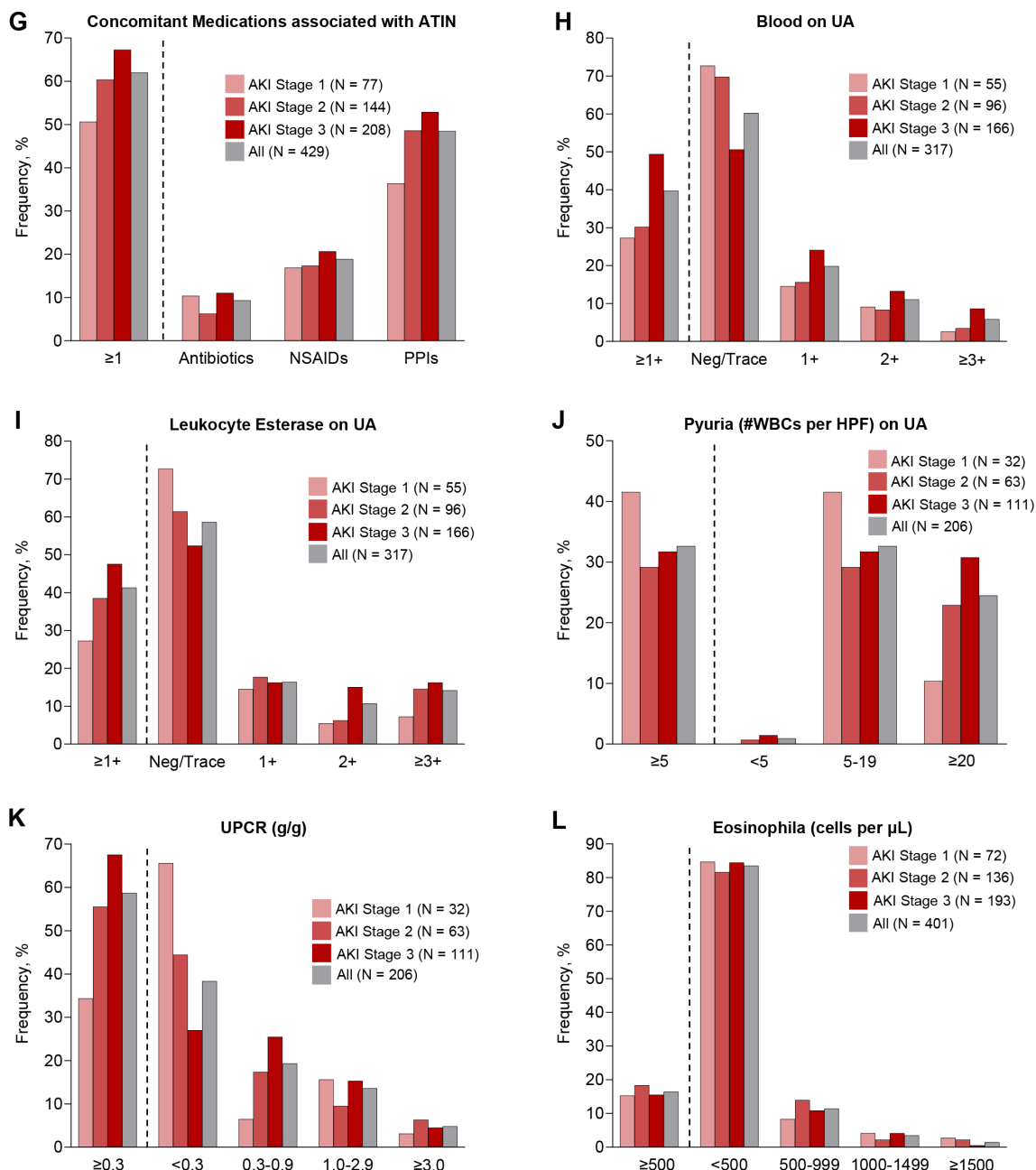
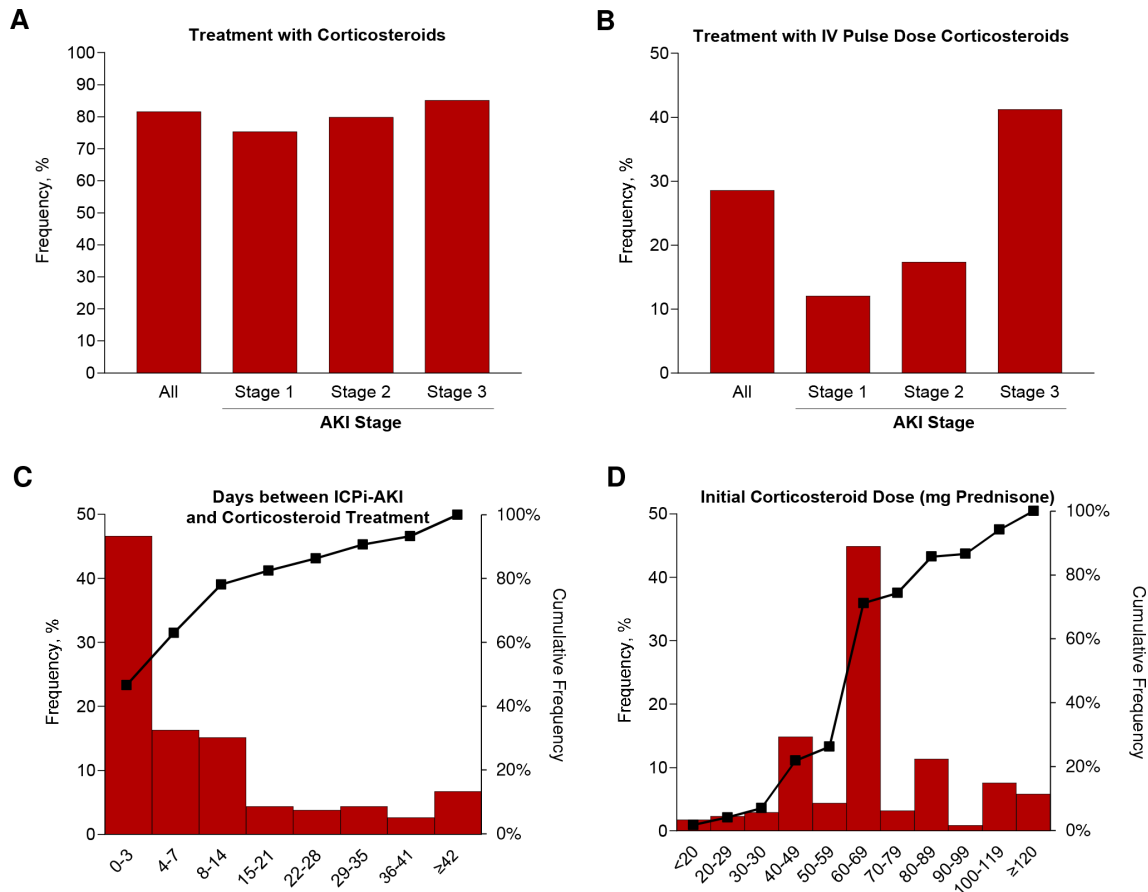


Figure 2 (Continued)



**Figure 2** Clinical features of ICPI-AKI. (A) The number of weeks between ICPI initiation and ICPI-AKI diagnosis. (B) The number of weeks between the last ICPI cycle and ICPI-AKI diagnosis. (C) Distribution of AKI severity. (D) Serum creatinine trend (median, IQR). (E) Frequency of extrarenal irAEs occurring before (>14 days) or concomitant (within 14 days before or after) with ICPI-AKI diagnosis. Other irAEs include hypophysitis (0.7% prior, 1.4% concomitantly), adrenalitis (0.2% prior, 1.4% concomitantly), type 1 diabetes mellitus (0% prior, 0.5% concomitantly), and myocarditis (1.2% prior, 0.2% concomitantly). (F) Distribution of pathologies among the 151 patients who underwent biopsy. Other includes 2 patients with FSGS and one patient with each of the following: reactive amyloidosis, AA amyloidosis, focal proliferative glomerulonephritis with C3 deposits, immune complex deposition disease not otherwise specified, mesangial proliferative immune complex mediated glomerulonephritis, pauci-immune glomerulonephritis, minimal change disease and thrombotic microangiopathy. (G) Frequency of potential ATIN-causing medications taken within 14 days before ICPI-AKI diagnosis. (H) Frequency of blood on UA at the time of ICPI-AKI. (I) Frequency of leukocyte esterase on UA at the time of ICPI-AKI. (J) Frequency of pyuria on UA at the time of ICPI-AKI. (K) Frequency of proteinuria at the time of ICPI-AKI. (L) Frequency of eosinophilia at the time of ICPI-AKI. AKI, acute kidney injury; ATIN, acute tubulointerstitial nephritis; ATN, acute tubular necrosis; FSGS, focal segmental glomerulosclerosis; HPF, high-power field; ICPI, immune checkpoint inhibitor; MN, membranous nephropathy; UA, urinalysis; UPCR, urine protein-to-creatinine ratio; WBCs, white blood cells.



**Figure 3** Treatment of ICPI-AKI. (A) Frequency of treatment with oral or intravenous corticosteroids by stage of initial ICPI-AKI. (B) Frequency of treatment with intravenous pulse dose corticosteroids by stage of initial ICPI-AKI. (C) Distribution of days between ICPI-AKI diagnosis and initiation of corticosteroids. (D) Distribution of initial corticosteroid dose (in prednisone equivalent units [mg]). AKI, acute kidney injury; ICPI, immune checkpoint inhibitor.

treatment with corticosteroids within 14 days following ICPI-AKI was also associated with a higher odds of renal recovery (OR 2.64; 95% CI 1.58 to 4.41; [figure 4C](#)). In a sensitivity analysis limited to patients treated with corticosteroids at any time following ICPI-AKI, initiation of corticosteroids within 3 days of ICPI-AKI was associated with a higher odds of renal recovery compared with later initiation (OR 2.09; 95% CI 1.16 to 3.79; [online supplemental figure S4](#)).

### Survival in patients with ICPI-AKI

During a median follow-up of 30 weeks (IQR 15–66) from ICPI-AKI diagnosis, 168 of 429 patients (39.2%) died. Survival was lower in patients with ICPI-AKI stage 3 compared with those with stages 1 and 2 in univariate analyses ([figure 5A](#)). In a multivariable model, only lower baseline eGFR was associated with a higher risk of death ([figure 5B](#)).

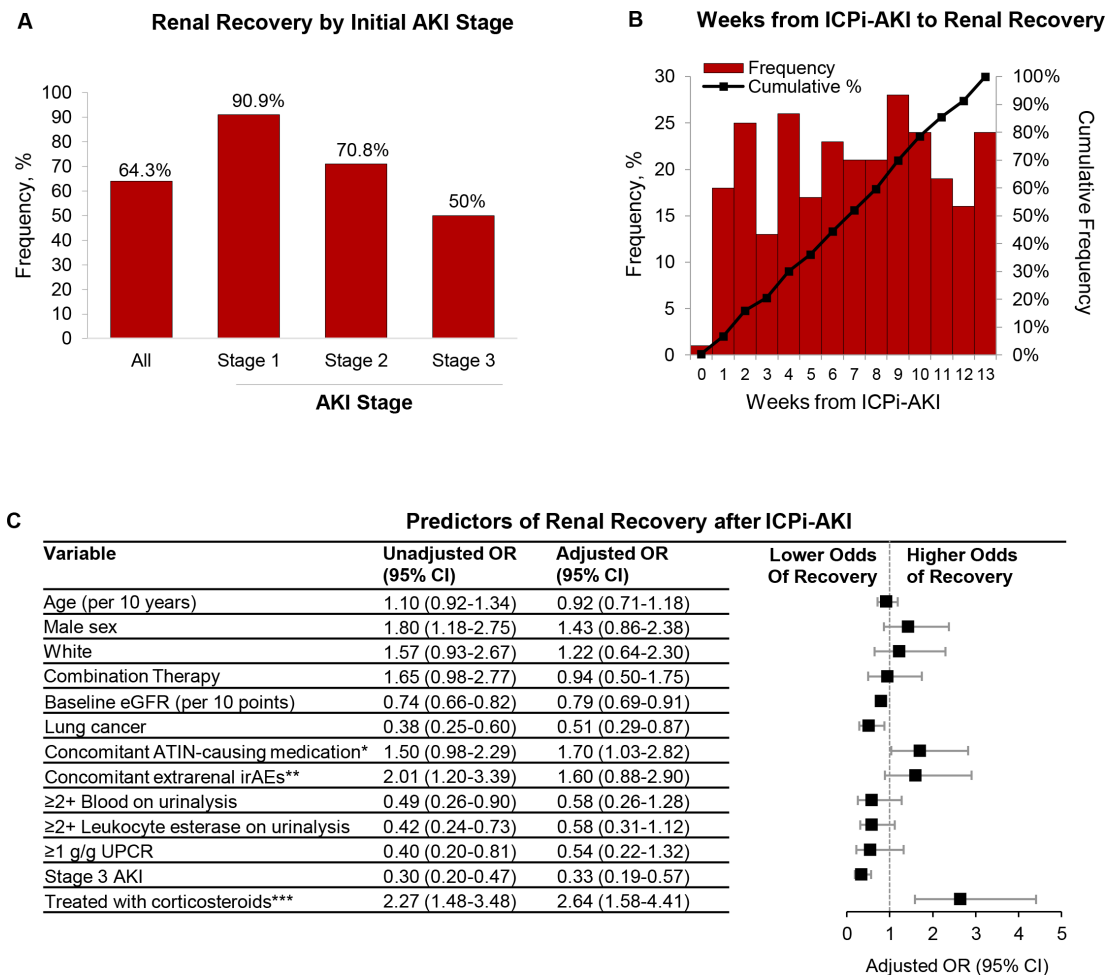
### ICPI rechallenge

A total of 121 of the 429 patients (28.2%) with ICPI-AKI were rechallenged with an ICPI, including 93 (76.9%) who had renal recovery and 28 (23.1%) who did not ([online supplemental figure S5](#)). Rechallenge occurred at a median of 1.9 months (IQR 1.1–4.0) after the initial

ICPI-AKI episode ([online supplemental figure S6](#)). Rechallenged patients were less likely to have had ICPI-AKI stage 3 initially compared with non-rechallenged patients (34.7% vs 53.9%, respectively) and were more likely to have had renal recovery following the initial ICPI-AKI episode (76.8% vs 59.4%, respectively) ([online supplemental table S7](#)). Survival was similar among patients rechallenged versus not rechallenged in the first 90 days ([online supplemental figure S7](#), panels A and B) and in the first 180 days following ICPI-AKI ([online supplemental figure S7](#), panels C and D).

### Recurrent ICPI-AKI after rechallenge

Of the 121 patients rechallenged, 20 (16.5%) developed recurrent ICPI-AKI, including 4 patients (20%) who developed AKI stage 1, 8 (40%) who developed AKI stage 2, and 8 (40%) who developed AKI stage 3 ([online supplemental figure S8](#)). Recurrent ICPI-AKI occurred at a median of 10 weeks (IQR 3–17) following rechallenge. There were no significant differences in the characteristics of rechallenged patients who developed recurrent ICPI-AKI versus those who did not, including receipt of corticosteroids at the time of rechallenge ([online supplemental table S8](#)).



**Figure 4** Characteristics of renal recovery among patients with ICPI-AKI. (A) Renal recovery overall and according to initial ICPI-AKI stage. (B) Time (in weeks) from ICPI-AKI diagnosis to renal recovery. (C) Predictors of renal recovery (total  $n=405$ , of whom 270 (66.7%) had renal recovery and 135 (33.3%) did not). Renal recovery was defined as a return of serum creatinine to  $\leq 50\%$  of the baseline value within 90 days of ICPI-AKI. Patients who died within 14 days of ICPI-AKI ( $n=24$ ) were excluded. All model covariates are shown in the figure. \*Denotes receipt of NSAIDs, PPIs, or antibiotics in the 14 days preceding ICPI-AKI. \*\*Extrarenal irAEs were assessed concomitantly (within 14 days before or after) with ICPI-AKI diagnosis. \*\*\*Refers to oral or intravenous corticosteroids initiated within 14 days following ICPI-AKI. AKI, acute kidney injury; ATIN, acute tubulointerstitial nephritis; eGFR, estimated glomerular filtration rate; ICPI, immune checkpoint inhibitor; irAEs, immune-related adverse events; NSAIDs, non-steroidal anti-inflammatory drugs; PPI, proton pump inhibitor; UPCr, urine protein-to-creatinine ratio.

All patients with recurrent ICPI-AKI had their ICPI therapy held, and most (14/20 [70%]) were treated with corticosteroids (online supplemental table S8). A total of 12 of the 20 patients (60%) with recurrent ICPI-AKI had renal recovery (online supplemental figure S5), which occurred at a median of 34 days (IQR 27–38) following diagnosis of recurrent ICPI-AKI. Patients who developed recurrent ICPI-AKI had higher mortality compared with those who did not (12/20 [60%] vs 32/101 [31.7%];  $p=0.02$ ).

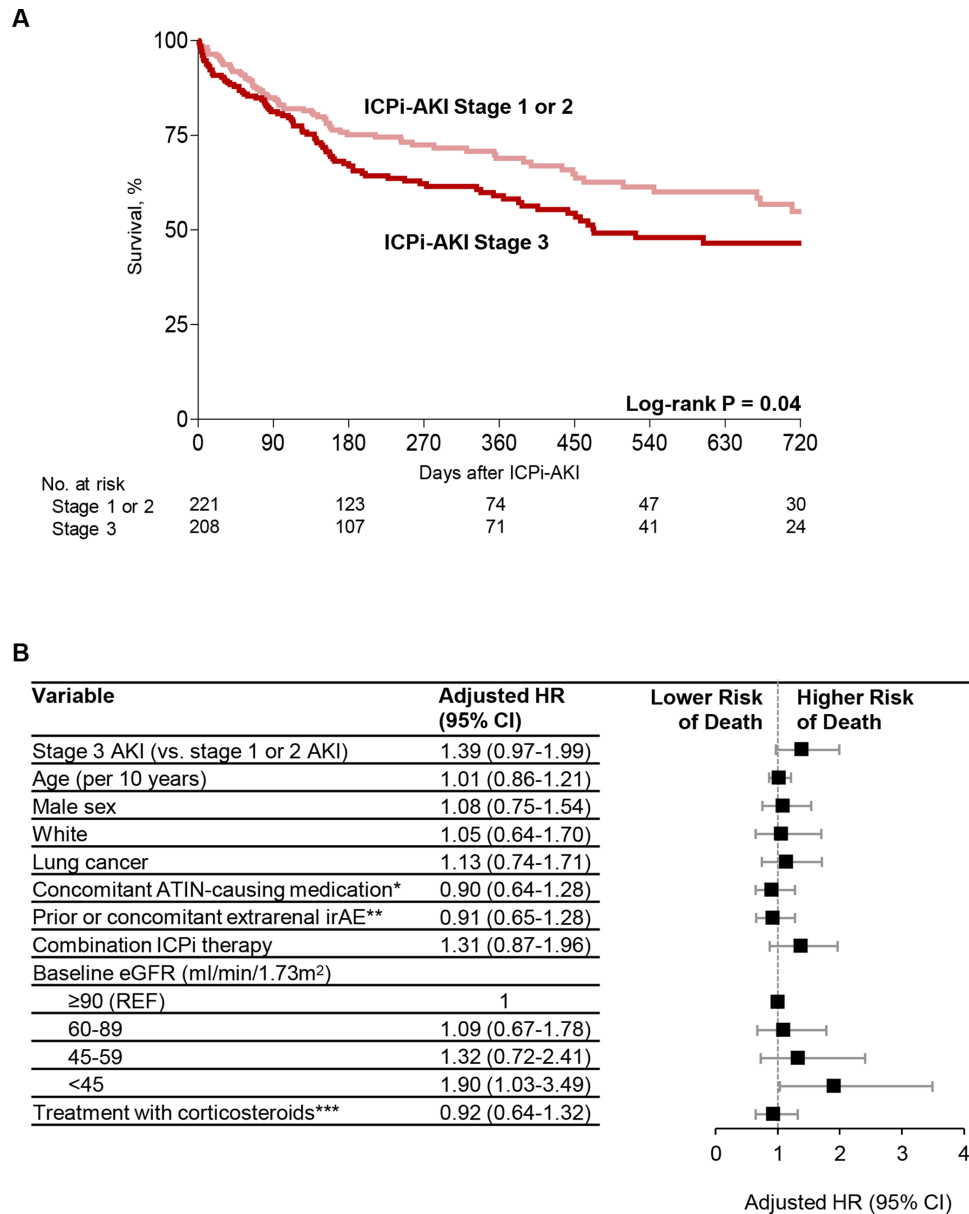
## DISCUSSION

In this international multicenter cohort study of over 400 patients with ICPI-AKI treated at 30 sites, we identified key risk factors, clinical features, and outcomes associated with ICPI-AKI. First, we found that lower baseline eGFR, PPI use, and extrarenal irAEs are each independently

associated with a higher risk of ICPI-AKI. Second, we expand on initial observations by our group<sup>3,19</sup> and others<sup>5–7,12–15,20</sup> regarding the clinicopathological features of ICPI-AKI, including the variable and often prolonged delay between ICPI initiation and ICPI-AKI, the high frequency of extrarenal irAEs in patients with ICPI-AKI, and ATIN as the most common histopathological lesion. Third, we found that renal recovery occurs in approximately two thirds of patients, and that early initiation of corticosteroids is associated with a higher likelihood of renal recovery. Fourth, among patients with ICPI-AKI who are rechallenged, fewer than one in five develop recurrent ICPI-AKI, half of whom subsequently have renal recovery.

Our findings regarding risk factors for development of ICPI-AKI, including lower baseline eGFR, PPI use, and extrarenal irAEs, are consistent with and expand





**Figure 5** Risk factors for death in patients with ICPI-AKI. (A) Survival among patients with stages 1 and 2 ICPI-AKI versus stage 3. (B) Multivariable Cox regression model showing predictors of death among patients with ICPI-AKI (total n=405, of whom 144 (35.6%) died). Patients who died within 14 days of ICPI-AKI (n=24) were excluded. All model covariates are shown in the figure. \*Denotes receipt of NSAIDs, PPIs, or antibiotics in the 14 days preceding ICPI-AKI. \*\*Extrarenal irAEs were assessed prior to (>14 days) or concomitant (within 14 days before or after) with ICPI-AKI. \*\*\*Refers to oral or intravenous corticosteroids initiated within 14 days following ICPI-AKI. AKI, acute kidney injury; ATIN, acute tubulointerstitial nephritis; eGFR, estimated glomerular filtration rate; ICPI, immune checkpoint inhibitor; irAEs, immune-related adverse events; NSAIDs, non-steroidal anti-inflammatory drugs; PPI, proton pump inhibitor.

on prior studies.<sup>3-5</sup> We previously reported an association between lower baseline eGFR and higher risk of ICPI-AKI.<sup>3</sup> However, it is unclear whether the risk of immunological injury to the kidneys from ICPIs is truly increased in patients with underlying chronic kidney disease, or if lower renal reserve simply facilitates the crossing of a threshold increase in SCr (eg, ≥50%) considered to be ‘AKI’ in response to an insult such as ATIN. Our finding that baseline eGFR was no longer associated with ICPI-AKI when limited to patients with moderate-to-severe kidney injury suggests the latter. PPIs have

been recognised as an important cause of drug-induced ATIN in both the general population<sup>21-23</sup> and in patients receiving ICPIs.<sup>3,4</sup> PPIs and other medications associated with ATIN may predispose to ICPI-AKI through loss of tolerance via activation or reactivation of drug-specific T cells. Thus, PPIs should be used with caution in patients receiving ICPI therapy. Finally, prior or concomitant extrarenal irAEs were present in over half of the patients with ICPI-AKI in our cohort. Thus, the presence of extrarenal irAEs should raise clinical suspicion for ICPI-AKI in patients receiving ICPI therapy who develop AKI.<sup>24</sup>

We did not identify any clinical features that were reliably present or absent in patients with ICPI-AKI. The latency period between ICPI initiation and ICPI-AKI was highly variable, with some patients developing ICPI-AKI more than a year after ICPI initiation, suggesting that clinicians should remain vigilant regarding ICPI-AKI even when it occurs late. The finding that ICPI-AKI occurred at a median of 16 weeks suggests there is a longer latency period with ICPI-AKI than extrarenal irAEs.<sup>25–27</sup> Urinary findings such as pyuria, hematuria, and proteinuria were present in only 30%–60% of the patients in our cohort, and thus cannot be used in isolation to differentiate ICPI-AKI from other causes of AKI. Among the patients biopsied, ATIN was by far the most common lesion (occurring in 83% of patients), as has been described in previous reports.<sup>3 4 12 19</sup> However, one in six patients had an alternative primary lesion, including glomerulonephritis and other lesions that have been described previously.<sup>14 28</sup> The lack of clinical features that distinguish ICPI-AKI from other causes of AKI, as well as the wide spectrum of histopathological findings in patients with ICPI-AKI, underscores the importance of performing renal biopsy in patients without contraindications, particularly in patients with atypical features (eg, nephrotic range proteinuria) or those who do not respond to corticosteroids.<sup>24</sup>

We found that approximately two-thirds of patients with ICPI-AKI have renal recovery, occurring at a median of 7 weeks. Patients taking concomitant ATIN-causing medications had a higher rate of renal recovery, consistent with our previous findings.<sup>3</sup> This may be a reflection of T-cell reactivity to the exogenous drug, whereby cessation of the causative agent in conjunction with corticosteroids can rapidly attenuate the immune response. We paradoxically found that higher baseline eGFR was associated with a lower likelihood of renal recovery. One possible explanation for this finding is that patients with more renal reserve would need to sustain a greater renal insult in order to achieve the same relative increase in SCr as a patient with less renal reserve.<sup>29</sup> We also found that patients with higher stages of ICPI-AKI were less likely to have renal recovery, highlighting the importance of early recognition and treatment of ICPI-AKI. This is further supported by the finding that early initiation of corticosteroids (within 3 days of ICPI-AKI diagnosis) was associated with a higher rate of renal recovery. Similar findings have been observed with extrarenal irAEs, including myocarditis,<sup>30</sup> as well as with other forms of drug-induced AKI.<sup>31</sup>

Data on renal and overall outcomes in patients with ICPI-AKI who are rechallenged are sparse.<sup>4 15 32</sup> A total of 121 patients in the current study were rechallenged, and fewer than one in five developed recurrent ICPI-AKI. We did not detect a survival difference in patients rechallenged versus those not rechallenged; however, patients with more aggressive malignancies may have been preferentially selected for rechallenge, which could have obscured our ability to identify a survival benefit. Given the low incidence of recurrent ICPI-AKI, it seems

reasonable to consider rechallenge in patients for whom ICPIs are the optimal therapy.

Although this is the largest study of ICPI-AKI to date, we acknowledge several limitations. First, not all patients had a kidney biopsy to confirm the diagnosis, which reflects clinical practice in which patients are often treated empirically. Second, our outcomes analyses were focused on renal recovery and overall survival, and we did not collect data on tumor response to ICPI therapy. We, therefore, do not have data on cancer status at the time of ICPI-AKI or rechallenge. Third, patients in our cohort were disproportionately treated at sites in the USA, which may affect the generalizability of our findings.

In this international multicenter cohort study, we identified risk factors, clinical features, and histopathological findings associated with ICPI-AKI, predictors, rates, and timing of renal recovery following ICPI-AKI, and the incidence of recurrent ICPI-AKI after rechallenge. Future studies with longitudinal biospecimen collection are needed to provide additional insight into the mechanisms of ICPI-AKI, and to aid clinicians in differentiating it from other causes of AKI. Further, randomized clinical trials are needed to evaluate the efficacy and safety of different corticosteroid dosing regimens and other forms of immunosuppression for optimal management of ICPI-AKI.

#### Author affiliations

<sup>1</sup>Division of Renal Medicine, Brigham and Women's Hospital, Boston, Massachusetts, USA

<sup>2</sup>University of Vermont Larner College of Medicine, Burlington, Vermont, USA

<sup>3</sup>Division of Nephrology, Massachusetts General Hospital, Boston, Massachusetts, USA

<sup>4</sup>Division of Nephrology, Department of Internal Medicine, The Ohio State University Medical Center, Columbus, Ohio, USA

<sup>5</sup>Nephrology Department, Vall d'Hebron University Hospital, Vall d'Hebron Institute of Research, Barcelona, Spain

<sup>6</sup>Department of Critical Care & Nephrology, King's College London, Guy's and St Thomas' Hospital, London, UK

<sup>7</sup>Division of Nephrology & Hypertension, Mayo Clinic, Rochester, Minnesota, USA

<sup>8</sup>Division of Internal Medicine, Section of Nephrology, University of Texas MD Anderson Cancer Center, Houston, Texas, USA

<sup>9</sup>Division of Nephrology, Stanford University School of Medicine, Palo Alto, California, USA

<sup>10</sup>Renal Service, Memorial Sloan Kettering Cancer Center and Weill Cornell Medical College, New York, New York, USA

<sup>11</sup>Dana-Farber Cancer Institute Survivorship Program, Boston, Massachusetts, USA

<sup>12</sup>Division of Kidney Diseases and Hypertension, Donald and Barbara Zucker School of Medicine at Hofstra/Northwell, Northwell Health, Hempstead, New York, USA

<sup>13</sup>Division of Nephrology, Duke University School of Medicine, Durham, North Carolina, USA

<sup>14</sup>Division of Nephrology and Hypertension, University Hospital Cleveland Medical Center, Cleveland, Ohio, USA

<sup>15</sup>Department of Microbiology, Immunology and Transplantation, Laboratory of Molecular Immunology (Rega Institute for Medical Research), KU Leuven, Leuven, Belgium

<sup>16</sup>Division of Nephrology, University Hospitals Leuven, Leuven, Belgium

<sup>17</sup>Department of Nephrology and Hypertension, Northwestern University Feinberg School of Medicine, Chicago, Illinois, USA

<sup>18</sup>Division of Nephrology, University of Washington, Seattle, Washington, USA

<sup>19</sup>Department of Nephrology and Medical Intensive Care, Charité – Universitätsmedizin Berlin, Berlin, Germany

<sup>20</sup>Division of Nephrology, Department of Medicine, University of California San Francisco, San Francisco, California, USA

- <sup>21</sup>Institute of Clinical Medicine, College of Medicine, National Cheng Kung University, Tainan, Taiwan
- <sup>22</sup>Division of Cardiovascular Medicine, Chi-Mei Medical Center, Tainan, Taiwan
- <sup>23</sup>Department of Biotechnology, Southern Taiwan University of Science and Technology, Tainan, Taiwan
- <sup>24</sup>Institute of Nephrology and Hypertension, Sheba Medical Center and Sackler School of Medicine, Tel Aviv University, Tel Hashomer, Tel Aviv, Israel
- <sup>25</sup>Katz Family Division of Nephrology and Hypertension, Department of Medicine, University of Miami Miller School of Medicine, Miami, Florida, USA
- <sup>26</sup>Division of Nephrology, Hypertension, and Renal Transplant, Department of Medicine, University of Florida, Gainesville, Florida, USA
- <sup>27</sup>Division of Nephrology, University of Virginia Health System, Charlottesville, Virginia, USA
- <sup>28</sup>Service of Nephrology, Department of Medicine, University Hospital of Geneva, Geneva, Switzerland
- <sup>29</sup>Department of Medical Oncology, Assistance Publique - Hôpitaux de Paris, Paris, France
- <sup>30</sup>Division of Nephrology, University Health Network, University of Toronto, Toronto, Ontario, Canada
- <sup>31</sup>Division of Hematology-Oncology, VAGLAHS, Department of Medicine, David Geffen School of Medicine at University of California-Los Angeles, Los Angeles, California, USA
- <sup>32</sup>Division of Hematology/Oncology and Division of Nephrology, The University of Alabama School of Medicine, Birmingham, Alabama, USA
- <sup>33</sup>Division of Nephrology, Department of Medicine, Icahn School of Medicine at the Mount Sinai Hospital, New York, New York, USA
- <sup>34</sup>Renal-Electrolyte and Hypertension Division, Perelman School of Medicine, University of Pennsylvania, Philadelphia, Pennsylvania, USA
- <sup>35</sup>Department of Internal Medicine, Division of Nephrology, University Medical Center Groningen, University of Groningen, Groningen, The Netherlands
- <sup>36</sup>Division of Medical Oncology, Department of Internal Medicine, The Ohio State University, Columbus, Ohio, USA
- <sup>37</sup>Nephrology Department, San Carlos Clinical University Hospital, Madrid, Spain
- <sup>38</sup>Department of Critical Care & Nephrology, Guy's and St Thomas Hospital, London, UK
- <sup>39</sup>Division of Nephrology, King Chulalongkorn Memorial Hospital, Bangkok, Thailand
- <sup>40</sup>Sechenov Biomedical Science and Technology Park, Sechenov First Moscow State Medical University, Moscow, Russian Federation
- <sup>41</sup>Department of Oncology, Guy's & St Thomas Hospital, London, UK
- <sup>42</sup>Division of Epidemiology and Population Health, Stanford University, Palo Alto, California, USA
- <sup>43</sup>Laboratory of Respiratory Diseases and Thoracic Surgery (BREATHE), Department of Chronic Diseases and Metabolism, KU Leuven, Leuven, Belgium
- <sup>44</sup>Department of Pneumology, University Hospitals Leuven, Leuven, Belgium
- <sup>45</sup>Robert H. Lurie Comprehensive Cancer Center, Northwestern University, Chicago, Illinois, USA
- <sup>46</sup>Max Delbrück Center for Molecular Medicine, Helmholtz Association, Berlin, Germany
- <sup>47</sup>Division of Population Sciences, Medical Oncology, Dana-Farber Cancer Institute, Boston, MA, USA
- <sup>48</sup>Division of Oncology, Massachusetts General Hospital, Boston, MA, USA
- <sup>49</sup>New York Nephrology Vasculitis and Glomerular Center, Albany, New York, USA

**Twitter** Shruti Gupta @ShrutiGKidney, Sandra M Herrmann @HerrmannMd and David E Leaf @DavidLeaf9

**Collaborators** ICPi-AKI Consortium Investigators: Assistance Publique-Hôpitaux de Paris (AP-HP), Sorbonne Université, Hôpital Pitié-Salpêtrière: Luca Campedel, Joe-Elie Salem, Corinne Isnard Bagnis. Brigham and Women's Hospital/Dana-Farber Cancer Institute: Shruti Gupta, David E. Leaf, Harkarandeep Singh, Shveta S. Motwani, Naoka Murakami, Maria C. Tio, Osama E. Rahma, Suraj S. Mothi, Umud Selamet, Deborah Schrag. Charité – Universitätsmedizin Berlin: Sebastian Loew, Kai M. Schmidt-Ott. Chi-Mei Medical Center: Weiting Chang. Donald and Barbara Zucker School of Medicine: Kenar D. Jhaveri, Rimda Wanchoo, Yuriy Khanin, Jamie S. Hirsch, Vipulbhai Sakhiya, Daniel Stalbow, Sylvia Wu. Duke University Medical Center: David I. Ortiz-Melo. Guy's and St. Thomas NHS Hospital: Mariëes Ostermann, Nuttha Lumlertgul, Nina Seylanova, Lucy Flanders, Armando Cennamo, Sophie Papa, Anne Rigg, Nisha Shaunak. Harvard Medical School: Zoe A. Kibbelaar. Heidelberg University Hospital: Karolina Benesova. Icahn School of Medicine at Mount Sinai Hospital: Priya Deshpande. Massachusetts General Hospital: Meghan

E. Sise, Kerry L. Reynolds, Harish S. Seethapathy, Meghan Lee, Ian A. Strohben. Mayo Clinic: Sandra M. Herrmann, Busra Isik. Memorial Sloan Kettering Cancer Center: Ilya G. Glezerman. New York Nephrology Vasculitis and Glomerular Center: Frank B. Cortazar. Northwestern University: Vikram Aggarwal, Sunandana Chandra. Ohio State University: Jason M. Prosek, Sethu M. Madhavan, Dwight H. Owen, Marium Husain. Sheba Medical Center: Pazit Beckerman, Sharon Mini. Stanford University School of Medicine: Shuchi Anand, Pablo Garcia, Aydin Kaghazchi. University of Alabama at Birmingham: Sunil Rangarajan. University of California-Los Angeles: Daniel Sanghoon Shin, Grace Cherry. University of California-San Francisco: Christopher A. Carlos, Raymond K. Hsu, Andrey Kisel. University Hospitals Cleveland Medical Center: Arash Rashidi, Sheru K. Kansal, Nicole Albert, Katherine Carter, Vicki Donley, Tricia Young, Heather Cigoi. University Hospital of Geneva: Sophie De Seigneux, Thibaud Koessler. University Hospitals Leuven: Ben Sprangers, Els Wauters. University of Florida: Chintan V. Shah. University Medical Center Groningen: Mark Eijgelsheim. University of Miami Miller School of Medicine: Zain Mithani, Javier A. Pagan. University of Pennsylvania Health System: Gaia Coppock, Jonathan J. Hogan. University of Texas MD Anderson Cancer Center: Ala Abudayyeh, Omar Mamlouk, Jamie S. Lin, Valda Page. University of Toronto: Abhijat Kitchlu. University of Vermont Larner College of Medicine: Samuel AP Short. University of Virginia Health System: Amanda D. Renaghan, Elizabeth M. Gaughan. University of Washington: A. Bilal Malik. Vall d'Hebron University Hospital: Maria Jose Soler, Clara Garcia-Carro, Sheila Bermejo, Enriqueta Felip, Eva Muñoz-Couselo, Maria Josep Carreras.

**Contributors** Concept and design: all authors. Acquisition, analysis, or interpretation of data: all authors. Drafting of the manuscript: SG and DEL. Critical revision of the manuscript for important intellectual content: all authors. Statistical analysis: SG, SAPS, and DEL. Preparation of figures and tables: SG, SAPS, and DEL. All authors approved the final version of the manuscript, and accept responsibility to submit for publication. SG and DEL have verified the underlying data and are accountable for all aspects of the submitted work.

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#### ORCID iDs

Shruti Gupta <http://orcid.org/0000-0002-5747-2151>

Abhijat Kitchlu <http://orcid.org/0000-0002-4340-5046>

Meghan D Lee <http://orcid.org/0000-0002-6895-5968>

Omar Mamlouk <http://orcid.org/0000-0002-1242-2740>

Jamie S Lin <http://orcid.org/0000-0002-7073-9658>

David E Leaf <http://orcid.org/0000-0001-7875-090X>

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