these characteristics may respond adequately to less intense immunosuppression.

In summary, patients with lupus and evidence of an active urine sediment but low levels of proteinuria (<0.5 g/d) may have important kidney pathology that warrants aggressive treatment. Despite current guidelines that recommend a proteinuria threshold of >0.5 to 1 g/d, a diagnostic kidney biopsy in SLE patients with low-grade proteinuria should be considered.

DISCLOSURE
All the authors declared no competing interests.

SUPPLEMENTARY MATERIAL
Supplementary File (PDF)
Supplementary Methods.
Figure S1. Study design and cohort flowchart.

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Time to Acute Kidney Injury in β-Lactam—Induced Acute Interstitial Nephritis

Benjamin Lazarus¹,², Matthew R.P. Davies², Jason A. Trubiano³,⁴ and Rebecca Pellicano¹,⁵

¹Department of Nephrology, Monash Health, Victoria, Australia; ²Department of Nephrology, Austin Health, Victoria, Australia; ³Department of Infectious Diseases, Austin Health, Victoria, Australia; ⁴Department of Medicine (Austin Health), University of Melbourne, Victoria, Australia; and ⁵Department of Medicine, Monash University, Victoria, Australia

Correspondence: Benjamin Lazarus, Monash Medical Centre, 246 Clayton Road, Clayton, Victoria 3168, Australia. E-mail: Benjamin.Lazarus@monashhealth.org

Received 30 January 2020; revised 7 April 2020; accepted 8 April 2020; published online 20 April 2020

Kidney Int Rep (2020) 5, 1061–1089; https://doi.org/10.1016/j.ekir.2020.04.008
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Acute interstitial nephritis (AIN) is a leading cause of acute kidney injury (AKI) in hospitalized patients, and β-lactam antibiotics are a common cause.¹ Early diagnosis of β-lactam—induced AIN is essential to direct changes to therapy, but it is often difficult to distinguish AIN from other causes of AKI, such as acute tubular necrosis. A kidney biopsy is the gold-standard diagnostic test, but carries significant risks and is often delayed. Mislabeling patients with a β-lactam allergy, in the absence of a biopsy, also restricts future antibiotic choice and drives inappropriate prescribing.² Clinical features alone cannot reliably differentiate AIN from other causes of AKI, and the temporal relationship between drug exposure and onset of AKI remains poorly understood. Some reports suggest that β-lactam—induced AIN typically occurs
more than 8 days after exposure; however, data are limited, and existing studies use nonstandardized definitions for AKI. We aimed to characterize the time between β-lactam exposure and AKI, defined by doubling of baseline serum creatinine, in a series of hospitalized patients with biopsy-proven AIN across 2 health services in Melbourne, Australia.

RESULTS
A total of 72 cases of biopsy-proven AIN were identified, of which 49 were attributed to drugs and 14 to β-lactams specifically (Figure 1). I.v. flucloxacillin, piperacillin–tazobactam, and benzylpenicillin were used in 8 cases, 5 cases, and 1 case, respectively. Among the 14 patients with β-lactam–induced AIN, the median age was 57 years (interquartile range [IQR], 40–68 years); 6 patients were female; mean baseline creatinine was 78 µmol/l; and median duration of β-lactam use was 8 days (IQR, 3–12 days). The indications for β-lactams included cellulitis (n = 5), methicillin-susceptible Staphylococcus aureus (MSSA) osteomyelitis (n = 2), MSSA bacteremia (n = 2), fever of unknown origin (n = 2), dog bite, pneumonia, and cholangitis. In 7 patients (50%) there were concurrent histopathological features on the kidney biopsy specimen that could contribute to AKI (i.e., acute renal pathology), 6 with acute tubular necrosis and 1 with IgA co-dominant postinfectious glomerulonephritis (PIGN). Three had features of background mild chronic damage. The median time to AKI was 2.5 days (IQR, 2–8 days). There was no appreciable difference in the time to AKI between centers (Figure 2). The mean time to AKI among 7 patients with concurrent acute renal pathology was 3.6 days (range, 1–13 days), whereas among 7 patients without concurrent acute renal pathology it was 6.1 days (range, 1–11 days); however, this difference was not statistically significant (P = 0.27, 2-sided t test). Two patients, 1 patient with and 1 patient without concurrent acute renal pathology, had a known pre-existing allergy to penicillin. One patient reported an unknown reaction, and the other reported prior kidney injury following penicillin exposure.

DISCUSSION
In our retrospective multicenter case series of 14 patients with β-lactam–induced biopsy-proven acute interstitial nephritis, the median time to doubling of serum creatinine was 2.5 days, which is earlier than previous estimates and shorter than typically expected in primary T-cell–mediated hypersensitivities. Our finding likely reflects the realities of clinical practice, in which a combination of other factors may contribute to earlier onset of AKI, and selection bias toward more severe cases that warrant biopsy.
Our study is consistent with previous case series of drug-induced AIN, affirming that β-lactams, particularly flucloxacinil and piperacillin–tazobactam, nonsteroidal anti-inflammatory drugs, and proton pump inhibitors are the leading culprit agents in Australia.6 Cephalosporin and carbapenem antibiotics were not identified as culprit agents in either study. Recovery of renal function was observed in all cases, despite 6 of the 14 patients requiring hemodialysis. Serum creatinine was measured on a daily basis during inpatient stay. Flucloxacillin was the most common antibiotic, and to further characterize the contribution of serum creatinine were not prospectively measured, and thus not standardized, but in most cases serum creatinine was measured on a daily basis during the inpatient stay. Flucloxacinil was the most common culprit agent identified in our case series; however, the frequency with which each of the β-lactams was used is unknown, and prospective studies are required to determine the relative incidence of AIN induced by different β-lactams.

In conclusion, a diagnosis of β-lactam—induced AIN should be considered in hospitalized patients who sustain an AKI following β-lactam use, even if doubling of serum creatinine occurs within 2 or 3 days of exposure. This time frame may reflect more severe or atypical cases, because only cases in which patients underwent biopsy were available for our analysis. Larger studies are required to determine risk factors for early-onset AKI in β-lactam—induced AIN. A case-control study would be a logical next step to investigate this issue. Furthermore, we believe that the standard practice of avoiding all β-lactams following an episode of penicillin-induced AIN should be revisited, because re-exposure to alternative β-lactam antibiotics was not associated with recurrence of AKI in our case series.

DISCLOSURE
All the authors declared no competing interests.

SUPPLEMENTARY MATERIAL
Supplementary File (PDF)
Supplementary Methods.

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Author/s:
Lazarus, B; Davies, MRP; Trubiano, JA; Pellicano, R

Title:
Time to Acute Kidney Injury in beta-LactamInduced Acute Interstitial

Date:
2020-07-01

Citation:
Lazarus, B., Davies, M. R. P., Trubiano, J. A. & Pellicano, R. (2020). Time to Acute Kidney Injury in beta-LactamInduced Acute Interstitial. KIDNEY INTERNATIONAL REPORTS, 5 (7), pp.1068-1070. https://doi.org/10.1016/j.ekir.2020.04.008.

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