908 WILEY NEPHROLOGY

There are two reports^{3,5} in children. Both reports were in patients with chronic kidney disease. The first child had received alkaline water as a form of complementary medicine³ and the second child had been prescribed rather high doses of sodium bicarbonate (6 mmol/kg/day).⁴ Our patient had high serum creatinine at presentation, but the renal function had improved to normal.

CONFLICT OF INTEREST

The authors have declared that no Conflict of interest exists.

ETHICS STATEMENT

All procedures performed in studies involving human participants were in accordance with the ethical standards of the institutional and/or national research committee and with the 1964 Helsinki declaration and its later amendments or comparable ethical standards.

> Komatineni Pravallika Sundaramoorthy Mathini Gaddam Srilakshmi Rapur Ram 🕩 Vishnubotla Siva Kumar

(BAPSN

Department of Nephrology, SVIMS, TIrupati, India

Correspondence

Rapur Ram, Department of Nephrology, SVIMS, Tirupati, Email: ram_5_1999@yahoo.com DOI 10.1111/nep.14103

ORCID

Rapur Ram (D) https://orcid.org/0000-0002-3523-3547

REFERENCES

- 1. Kermanshai R, McCarryBE RJ, Summers PS, Weretilnyk EA, Sorger GJ. Benzyl isothiocyanate is the chief or sole anthelmintic in papaya seed extracts. Phytochemistry. 2001;57:427-435.
- 2. Jaiswal N, Pandey VP, Dwivedi UN. Purification of a thermostable alkaline laccase from papaya (Carica papaya) using affinity chromatography. Int J Biol Macromol. 2015:72:326-332.
- 3. Kermond R, Carter S, Quinlan C. A child presents with acute kidney injury, alkalosis and hypercalcaemia-a new-age cause for a historical syndrome: answers. Pediatr Nephrol. 2022;15:1-4.
- 4. Movsés-Neto M. Guimarães FM. Avoub FH. Vieira-Neto OM. Costa JA. Dantas M. Acute renal failure and hypercalcemia. Ren Fail. 2006:28(2):153-159.
- 5. Kari J, El Desoky SM. Milk alkali syndrome in an infant with chronic kidney disease. Pediatric Health Med Ther. 2012;3:19-23.

Minimal change disease with Jack jumper ant stings: A case report

Insect stings from the order Hymenoptera have been associated with nephrotic syndrome, including minimal change disease (MCD) and other glomerular pathologies, since the first reported case following bee sting in 1955.¹ We document the first case of MCD associated with Jack Jumper Ant (JJA) sting. The JJA (Myrmecia pilosula) is endemic to southern and eastern Australia. JJA venom comprises peptides with broad enzymatic activity, histamine-like and eicosanoid releasing factors. The pilosulin peptides are the major components of JJA venom, basic peptides rich in lysine and arginine with cytotoxic effects.²

A 72-year-old female was referred with oliguria and fluid overload 24-h following uneventful Nissen fundoplication. Sixteen days before admission she suffered over a dozen JJA stings. A week before admission she developed swelling of both legs and 10 kg weight gain. She had prior hypertension and a baseline creatinine of 82 µmol/L.

On examination blood pressure was 178/83 mmHg, with elevated jugular venous pressure, anasarca and bibasal crepitations. Urine output was 300 ml/24 hours. Serum creatinine was 140 µmol/L (reference range: 45-90 µmol/L). Urine microscopy revealed no erythrocytes, urine protein: creatinine ratio was 478 mg/mmol (reference range: <3.5 mg/mmol), and albumin: creatinine ratio (uACR) 450 mg/mmol (reference range: <18 mg/mmol). Serum albumin was 23 g/L, total cholesterol 4.8 mmol/L, and international normalized ratio 1.0. Specific immunoglobulin E (IgE) to JJA venom was negative (<0.10 kIU/L).

A kidney biopsy showed normal glomeruli on light microscopy, normal basement membrane on silver staining, and the absence of deposits on immunofluorescence. Electron microscopy confirmed foot process effacement involving >90% of the glomerulus, supporting the clinical diagnosis of MCD. Immunohistochemical staining demonstrated evidence of (i) glomerular and tubular oxidative stress, with positive staining for 4-hydroxynonenal; (ii) glomerular activation of the glucocorticoid-sensitive Angptl4 pathway, without alteration in podocyte marker synaptopodin; and (iii) tubulointerstitial inflammatory CD3⁺ T-cells and CD1c⁺ dendritic cells.

A trial of diuretic therapy failed with fluid gain and elevated creatinine to 181 µmol/L. After a week, prednisolone 75 mg daily was commenced with rapid improvement in clinical status. At 1-month postdischarge, renal biochemistry had improved (creatinine 127 µmol/L, albumin 32 g/L, uACR 173). At 5-months, kidney function and weight returned to baseline (creatinine 87 µmol/L, albumin 34 g/L, uACR 0.5). Prednisolone was tapered off without side-effects. Kidney function remained stable at 10 months.

Only one case of MCD following ant sting has been previously reported,³ and we document the first case as a consequence of JJA sting. JJA stings are associated with high rates of anaphylaxis in endemic areas,² but an allergic mechanism does not appear to be the main driver of pathology in this case where specific IgE to JJA venom was negative. Direct toxicity of JJA venom by interaction with lipid bilayer membranes is reported.⁴ We propose here that anionic podocytes of the glomerular basement membrane (GBM) are susceptible to the toxic effects of cationic *Pilosulin* peptides, resulting in oxidative stress. This case is consistent with previous reports that insect-sting associated nephrotic syndrome is steroid-responsive, with a good prognosis.

AUTHOR CONTRIBUTIONS

Each author has participated sufficiently in the work to take public responsibility for the content.

ACKNOWLEDGEMENTS

Human kidney tissue was obtained with informed patient consent following approval by the Royal Brisbane and Women's Hospital Human Research Ethics Committee (2002/011 and 2006/072). The work was funded by Pathology Queensland and a National Health and Medical Research Council (NHMRC) Project Grant (GNT1161319). Open access publishing facilitated by The University of Queensland, as part of the Wiley - The University of Queensland agreement via the Council of Australian University Librarians.

CONFLICT OF INTEREST

The authors declare that they have no conflict of interest.

Matthew Tunbridge^{1,2} Anca Grivei^{1,3} Andrew J. Kassianos^{1,2,3} Helen Davis¹ Anne Stewart⁴ NEPHROLOGY

WILEY 909

Helen Healy^{1,2,3} Saw Yu Mon^{1,2} George T. John^{1,2}

¹Kidney Health Service, Royal Brisbane and Women's Hospital, Brisbane, Queensland, Australia ²Faculty of Medicine, University of Queensland, Brisbane, Queensland, Australia ³Conjoint Internal Medicine Laboratory, Chemical Pathology, Pathology Queensland, Brisbane, Queensland, Australia ⁴Anatomical Pathology, Pathology Queensland, Brisbane, Queensland, Australia

Correspondence

George T. John, Kidney Health Service, Level 9 Ned Hanlon Building, Royal Brisbane and Women's Hospital, Butterfield St, Herston QLD 4029, Australia.

Email: george.john@health.qld.gov.au DOI 10.1111/nep.14104

ORCID

Andrew J. Kassianos D https://orcid.org/0000-0002-0994-3588

REFERENCES

- Rytand DA. Onset of the nephrotic syndrome during a reaction to bee sting. Stanford Med Bull. 1955;13:224-233.
- Davies NW, Wiese MD, Brown SG. Characterisation of major peptides in'jack jumper'ant venom by mass spectrometry. *Toxicon*. 2004;43: 173-183.
- 3. Swanson GP, Leveque JA. Nephrotic syndrome associated with ant bite. *Tex Med.* 1990;86:39-41.
- 4. Kourie JI, Shorthouse AA. Properties of cytotoxic peptide-formed ion channels. *Am J Physiol Cell Physiol*. 2000;278:C1063-C1087.