A 63-year-old woman was referred to a nephrologist for evaluation of decreased kidney function. The patient had no specific urinary symptoms or history of recurrent urinary tract infections, leg edema or kidney stones. She did not have nausea, vomiting, diarrhea or symptoms of steatorrhea, and had not been using anti-inflammatory agents. Her past medical history included gastresophageal reflux disease, chronic constipation, depression and remote left frontal lobe resection for benign tumour. Her medications included citalopram (20 mg daily), omeprazole (20 mg daily) and risperidone (3 mg at bedtime).

Her physical examination was unremarkable, including normal blood pressure. Laboratory test results showed an elevated serum creatinine level of 172 (normal 35–97) μmol/L and urea level of 11.2 (normal 2–8) mmol/L; eight months earlier, her levels were 71 μmol/L and 5.7 mmol/L, respectively (Appendix 1, available at www.cmaj.ca/lookup/suppl/doi:10.1503/cmaj.151327/-/DC1). Urinalysis was negative for protein, and no active sediment or crystals were reported. Abdominal ultrasonography showed bilateral echogenic kidneys with incidental gallstones. Serum immunoelectrophoresis showed a monoclonal band of IgG kappa measuring 9.3 (normal 6.3–14.9) g/L. These findings and the presence of anemia (hemoglobin 88 [normal 120–160] g/L) led us to consider paraproteinemia, which was ruled out by a normal result for bone marrow aspiration.

The progressive decline in kidney function raised the possibility of light-chain nephropathy, and a kidney biopsy was performed three months after presentation to the nephrophologist that showed acute tubular necrosis and calcium oxalate crystals consistent with oxalate nephropathy (Figure 1).

At that time, the patient’s serum creatinine level was 234 μmol/L, with a urea level of 11.7 mmol/L. Her 24-hour urine calcium level was low at 1.89 (normal range 2.50–7.50) mmol/d, with a high 24-hour urine oxalate level of 1489 (normal range 40–320) μmol/d (Appendix 1).

A preliminary dietary history failed to identify consumption of foods high in oxalate. The patient had a normal pyridoxine levelof
of oxalate content in the patient’s typical diet. The dietitian discovered that the patient had started eating a can (275 g per can) of cashews every two days (about 100–150 g per day or about 1 kg of cashews per week) for their “laxative effect,” about four months before her first visit to the nephrologist, after she learned from a television show that cashews were a “healthy food” and may help relieve constipation. The patient had found the cashews to be very helpful in this regard, and started eating them every day. The patient’s diet was poor in other nutrients, most notably calcium (Box 1).

The patient was advised to stop consuming cashews to limit the intake of oxalates to less than 50 g per day. Calcium carbonate (500 mg twice a day) was added with meals to act as an oxalate binder and to ensure appropriate calcium intake. Two months after following these recommendations, the patient’s urinary oxalate excretion had normalized (88 μmol/d) and her kidney function had improved (serum creatinine 184 μmol/L). Although the patient’s kidney function continued to improve, she was left with residual chronic kidney disease (serum creatinine 130 μmol/L) eight months after stopping the cashews (Appendix 1).

### Discussion

Oxalate, an antinutrient, is a conjugated anion of oxalic acid. It is ubiquitous in nature and is derived from various animal and plant sources. There are two main sources of oxalate: endogenous production in the liver from metabolism of its precursors (hydroxyproline, glycine and serine) and gastrointestinal absorption of oxalates in the diet. The oxalate content of a typical Western diet is approximately 80–120 mg per day, about 10% of which is absorbed via the gastrointestinal tract.1 Oxalate is not metabolized in humans and is excreted unchanged in urine. The normal daily urinary excretion of oxalate ranges between 40 and 320 μmol/d. Urinary excretion of oxalate above this range is classified as hyperoxaluria.2

### Pathogenesis of hyperoxaluria

Hyperoxaluria² may be primary or secondary. Primary hyperoxaluria results from an inherited defect in oxalate metabolism that results in increased endogenous production of oxalate [Box 2].2,3 Secondary hyperoxaluria may result from either dietary excess, enteric hyperabsorption (Figure 2), or enhanced endogeneous production resulting from exposure to its precursors (ascorbic acid) or from pyridoxine deficiency. The absorption of oxalate in the gastrointestinal tract is influenced by many factors, including gut health and function,4 oxalate intake or oxalate content of food5 (Appendix 2, available at www.cmaj.ca/lookup/suppl/doi:10.1503/cmaj.151327/-/DC1) and, most importantly, the intake of other nutrients such as calcium, magnesium and fat.1 High levels of dietary calcium4 and magnesium6 inhibit oxalate absorption through formation of insoluble oxalate salts in the gastrointestinal tract that are poorly absorbed. In addition, food preparation can affect oxalate absorption, as seen with “juicing.”8 Juices are concentrated forms of fruits and vegetables that may have high amounts of oxalate. Consequently, the increased amount of fluid load in the juices can promote paracellular absorption of oxalate in the intestines.8 The bioavailability of the oxalate in the meal is more important than its amount.

### Box 1: The patient’s typical daily intake of oxalate and calcium* from food

| Food                          | Oxalate content, mg | Calcium, mg* |
|-------------------------------|---------------------|--------------|
| Milk (250 mL)                 | 2                   | 309          |
| Egg (2)                       | 5                   | 87           |
| White bread slice (2)         | 10                  | 84           |
| Oatmeal (~ 75 g cooked)       | 25                  | 63           |
| Mushrooms (125 mL)            | 5                   | 5            |
| Cashews (100–150 g)           | 260–325             | 46           |
| Potatoes, mashed (125 mL)     | 25                  | 4            |
| Ham steak (1)                 | 2                   | 3            |
| Coffee (250 mL)               | 5                   | 5            |
| Total intake per day          | 339–409             | 606          |
| RDA per day                   | No clear recommendations† | 1200‡       |

Note: RDA = Recommended Dietary Allowance.

*Values obtained from food searches in the Canadian Nutrient File (https://food.nutrition.ca/cnf-fce/index-eng.jsp).

†There is no daily RDA for oxalate, because oxalate is an antinutrient. In the United States, oxalate intake is estimated to average between 150 and 200 mg/d.7

The American Academy of Nutrition and Dietetics recommends limiting dietary oxalate intake to less than 50 mg/day in patients with oxalate kidney stones (www.eatrightpro.org).

Dietary Reference Intakes for vitamin D and calcium. Health Canada (http://www.hc-sc.gc.ca/fn-an/nutrition/vitamin/vita-d-eng.php#a7).

### Box 2: Primary hyperoxaluria²,³

Primary hyperoxaluria (PH; autosomal recessive inherited disorders):

- **Type 1 (PH1) (80% of all PH)**
  - Decreased or absent activity of vitamin B₆–dependent hepatic peroxisomal enzyme (alanine-glyoxylate aminotransferase [AGXT]); AGXT gene located on chromosome 2
  - Failure to transaminate glyoxylate that results in increased urinary excretion of oxalate and glycolate

- **Type 2 (PH2) (10% of all PH)**
  - Mutation in the glyoxylate/hydroxyypyruvate reductase gene (GXRPR) that is located on chromosome 10
  - Reduced GXRPR activity leads to increased bioavailability of lactate and hydroxyypyruvate for conversion of oxalate and L-glycerate
  - Characterized by increased urinary excretion of L-glycerate

- **Type 3 (PH3) (about 10% of all PH)**
  - Mutation in the mitochondrial dihydrodipicolinate synthase-like gene (HOGAI)
  - Commonly presents as nephrolithiasis

- **Type 4 (PH4)**
  - Inherited defect in bone metabolism
  - Characterized by osteopenia and cataracts

- **Type 5 (PH5)**
  - Autosomal dominant inheritance
  - Characterized by chronic kidney disease and recurrent urinary tract infections

- **Primary hyperoxaluria (PH1, PH2, PH3, PH4, PH5)**
  - Also classified as hyperoxaluria²

- **Secondary hyperoxaluria**
  - Resulting from increased dietary intake of oxalate
  - Increased enteric hyperabsorption
  - Increased endogeneous production

High levels of dietary calcium4 and magnesium6 inhibit oxalate absorption through formation of insoluble oxalate salts in the gastrointestinal tract that are poorly absorbed. In addition, food preparation can affect oxalate absorption, as seen with “juicing.”8 Juices are concentrated forms of fruits and vegetables that may have high amounts of oxalate. Consequently, the increased amount of fluid load in the juices can promote paracellular absorption of oxalate in the intestines.8 The bioavailability of the oxalate in the meal is more important than its amount.
Consequences of hyperoxaluria

Hyperoxaluria, whether primary or secondary, can cause nephrolithiasis. However, it can also cause crystalline nephropathy through deposition of calcium oxalate crystals in the renal parenchyma that cause tubular damage, interstitial inflammation and fibrosis, and may present as acute kidney injury or chronic kidney disease. In patients with chronic diarrhea, the associated volume depletion and metabolic acidosis results in low urinary pH and hypocitraturia that, in conjunction with hyperoxaluria, potentiate calcium oxalate precipitation in the kidney and promote kidney injury. Oxalate nephropathy has been reported with excessive consumption of different oxalate-rich foods, including nuts and iced tea.

Cashew nuts (actually seeds) are attached to the bottom of the cashew apple (Figure 3), the fruit of the cashew tree (*Anacardium occidentale*). Cashews are considered a “healthy” food because of their monounsaturated fat content that is similar to olive oil. Nuts, despite their overall healthy nutrient profile, have substantial amounts of oxalate.

The oxalate content of cashews is moderate compared with other nuts; however, by consuming large amounts daily, the patient increased her total daily intake of oxalate to a level much higher than average (Box 1). In addition, her lower calcium intake promoted hyperabsorption of oxalate, because calcium was not available to bind oxalate in the gut. Furthermore, the higher fat content of cashews (about 45%), although healthy for the heart, perpetuated oxalate absorption by further reducing the calcium available to bind oxalate in the gut. All of these factors and possibly the higher concentration of intestinal soluble oxalate promoted hyperabsorption of oxalate, which resulted in increased urinary oxalate concentration, with precipitation of calcium oxalate crystals in the renal tubules causing kidney injury.

### Nephrolithiasis versus nephropathy

The exact mechanism of why kidney stones develop in some patients with hyperoxaluria and kidney injury in others is unknown and possibly multifactorial. However, recent evidence suggests that supersaturation of soluble oxalate causes increased expression of nucleotide-binding oligomerization domain–like receptor, pyrin domain 3 (NLRP3) inflammasomes in the kidney that trigger inflammation and potential kidney injury that may be acute or chronic rather than stone formation (nephrolithiasis).

### Management

Prevention is key, and foods rich in oxalate content should be consumed with caution. Even foods that are moderate in oxalate content may cause kidney injury, especially if dietary calcium intake is poor; oxalate bioavailability plays an important role.

Therapeutic options are extrapolated from studies in patients with primary hyperoxaluria and oxalate stones. Based on expert opinion, conservative measures should be implemented soon after diagnosis and include a diet low in oxalate and relatively high in calcium, maintenance of high fluid intake (at least 1.5 L per 1.73 m² per day) and consideration of potassium citrate to alkalize urine to increase solubility of oxalate crystals. The role of probiotics is controversial, with studies showing conflicting results.

### Conclusion

Crystalline nephropathies should be considered in the differential diagnosis of patients with unexplained renal dysfunction. A detailed dietary history may provide a clue to the cause, as in this patient’s case, and measures to control reversible factors should be implemented at the earliest opportunity to prevent further progression.
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