**“Milky” urine: a case of chyluria**

A 53-YEAR-OLD Asian man with type 2 diabetes mellitus presented to the Emergency Department with acute onset of generalised muscle cramps. He reported a 2-month history of polydipsia, polyuria and passing “milky” urine with blood clots. He had travelled widely throughout subtropical Asia. On examination, he was normotensive, with no lymphadenopathy, abdominal masses or oedema.

Urinalysis showed marked proteinuria, glycosuria and haematuria. The urine protein excretion rate was later confirmed to be 15.57 g/24 h (reference interval [RI], 0.02–0.15 g/24 h). Urine triglyceride measurement and lipoprotein electrophoresis confirmed the appearance of chylomicrons in the urine after an oral fat tolerance test (Box 1).

Biochemical analysis of serum showed the following levels: sodium 121 mmol/L (RI, 134–146 mmol/L), potassium 4.6 mmol/L (RI, 3.4–5.3 mmol/L), creatinine 46 µmol/L (RI, 60–105 µmol/L), glucose 19.7 mmol/L (RI, < 5.5 mmol/L), ferritin 16 mg/L (RI, 30–620 mg/L), 25-hydroxyvitamin D 11 nmol/L (RI, > 50 nmol/L), total cholesterol 5.2 mmol/L (RI, < 5.5 mmol/L), and triglyceride 1.8 mmol/L (RI, < 1.8 mmol/L). The patient had marked hypoproteinaemia and hypoalbuminaemia, with a total protein level of 39 g/L (RI, 63–80 g/L) and albumin level of 21 g/L (RI, 35–50 g/L), respectively. The serum IgE level was also raised (2300 kU/L; RI, < 210 kU/L).

The patient was mildly anaemic (haemoglobin level, 120 g/L [RI, 130–170 g/L]), with a normal erythrocyte sedimentation rate and C-reactive protein level. There was lymphopenia but no eosinophilia. No microfilariae or acid-fast bacilli were detected in thick blood films taken at midnight or in a urine sample. Serological and intradermal tests for filariae were also negative.

A chest x-ray and computed tomography scan of the abdomen and pelvis were normal. A biopsy of the right kidney showed evidence of mild mesangial change, consistent with diabetes mellitus. Lymphoscintigraphy showed delayed lymphatic transport, particularly on the left side, but no physical obstruction to lymphatic drainage. Contrast lymphangiography demonstrated a grossly abnormal lymphatic system in the pelvis and groin with a unilateral left-sided lymphorenal communication (Box 2).

A presumptive diagnosis of filariasis was made and the communication between the left kidney and the lymphatics was surgically disconnected. The chyluria recurred after surgery, but within 4 weeks of a therapeutic course of the antifilarial diethylcarbamazine there was complete resolution. Furthermore, the biochemical abnormalities reversed, consistent with urinary loss as the mechanism.

At 3-year follow-up, the patient remained symptom-free and without chyluria.

**Discussion**

Chyluria is rare in Australia but common in many parts of the world, particularly where *Wuchereria bancrofti*, the main agent of filariasis, is endemic. It occurs, on average, 5–10 years after the worm has died, and so there may be no evidence of active filariasis. However, a therapeutic trial of diethylcarbamazine should be considered before undertaking surgery. As spontaneous remissions of chyluria have been reported, we can not be certain whether the antifilarial therapy was responsible for resolution of the chyluria in this case.

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**1: Oral fat tolerance test**

Urine of patient before a 75 g oral fat tolerance test (0 h) and at serial time points (0.5, 1, 2, 3, 4, and 5 h) after the test.

**2: Contrast lymphangiography**

Contrast lymphangiogram, showing a grossly abnormal lymphatic system with a unilateral left-sided lymphorenal communication (arrow).