Mercury intoxication presenting with hypertension and tachycardia

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Abstract
An 11 year old girl presented with hypertension and tachycardia. Excess urinary catecholamine excretion suggested phaeochromocytoma but imaging studies failed to demonstrate a tumour. Other symptoms included insomnia and weight loss, and she was found to have a raised concentration of mercury in blood and urine. Mercury intoxication should be considered in the differential diagnosis of hypertension with tachycardia even in patients presenting without the skin lesions typical of mercury intoxication and without a history of exposure.

Case report
A Taiwanese girl was admitted to our institution because of hypertension (160/120 mm Hg) and tachycardia (120 beats/min). She had suffered from painful itching in her extremities but this had resolved before admission. Oscillometric 24 hour blood pressure monitoring revealed severe hypertension without nocturnal dipping (fig 1). Exposure to drugs and toxins was denied, and there was no family history of hypertension or malignant disease. Laboratory values showed normal thyroid and kidney function. Urinary concentrations of vanillylmandelic acid (VMA) were slightly raised (5.6–5.8 nmol/µmol creatinine; normal < 4.7); homovanillic acid (HMA) was within the normal range. Investigations including abdominal and cervical ultrasound, computed tomography scans, and an M-iodobenzylguanidine scan did not reveal a tumour. She was treated with enalapril. After failing to attend for follow up, she presented two months later with insomnia, depression, daytime fatigue, and loss of 12 kg body weight (initial body weight was 36 kg). On examination, she could barely stand because of weakness and ataxia. She was irritable and remained hypertensive despite medication. Nerve conduction velocity of the peroneal and median nerves was reduced and a cerebral magnetic resonance imaging scan showed no abnormality. Again, catecholamine concentrations (noradrenaline, dopamine, and VMA) were slightly raised. A screen for heavy metals was performed despite no history of exposure. The blood and urine of the patient (but not of her parents and younger sister) showed very high mercury concentrations (blood 20 µg/l, normal for adults < 10; urine 217 µg/l, normal for adults < 20). Direct questioning did not reveal any obvious sources of mercury intoxication such as recreational use, presence of a broken thermometer or use of old ointments or Chinese cups during the last year. During treatment with D-penicillamine, mercury concentrations returned to normal. At four month's follow up her heart rate and blood pressure (without medication) were normal. She gained weight, and ataxia resolved slowly over one year.

Discussion
A catecholamine producing tumour is not the only condition that causes hypertension and tachycardia as well as increased catecholamine concentrations. In inorganic mercury poisoning, the metal combines with the sulphydryl moiety of the toxic sulphydryl-containing proteins (reduced glutathione, metallothionein). The metal combines with sulphydryl groups in the free protein and is stored in the liver. The liver maintains a significant pool of mercury, which is inactivated by either storing mercury within toms include insomnia and weight loss, and hypertension with tachycardia even in patients presenting without the skin lesions typical of mercury intoxication and without a history of exposure.

Keywords: hypertension; tachycardia; mercury poisoning; phaeochromocytoma
group of S-adenosylmethionine, which acts as a cofactor for catecholamine-O-methyltransferase (COMT). COMT inhibition leads to accumulation of catecholamines, typically noradrenaline, adrenaline (epinephrine), and dopamine, and to a lesser degree VMA, but not usually HVA. The sympathetic overactivity explains the haemodynamic symptoms of acrodynia. Curiously, in our patient, VMA was the catecholamine most predominantly excreted. It is usual for mercury intoxication to show a much lesser degree of catecholamine excess than would normally be expected in a phaeochromocytoma or neuroblastoma.

Clinical signs of the fully developed form of acrodynia include pronounced mental changes such as insomnia and irritability, pain in the extremities, typical skin lesions (hence the nickname “pink disease”), profuse sweating, and anorexia as well as hypertension and tachycardia. Uncharacteristically, our patient did not have even a transitory rash and was not sweating excessively. However, desquamation and pink palms and soles are unusual in children beyond toddler age because of increased skin thickness.

Since mercury has been excluded from teething powders and ointments, inhalation of mercury from broken thermometers has accounted for most described poisonings in toddlers, and recreational use and inorganic mercury warming have been suggested as further causes of intoxication. In this case, no source could be found even after repeated questioning of family members. We considered exposure at home to be unlikely as the family of our patient live together in a small apartment. Her parents and sister excreted only traces of the heavy metal and we did not test the mercury levels in her home. A Chinese medicine containing a metal substance had been given to the girl, but was found to contain only traces of mercury. The child was, therefore, placed under constant surveillance. Repeated psychological evaluation of both parents and children did not indicate child abuse, but this cannot be excluded.

It is not known how many transient unexplained hypertensive episodes in children might be caused by mercury intoxication. Even though acrodynia is rare, its diagnosis can be made by a single urine analysis and, therefore, acrodynia should be excluded before expensive and invasive procedures are performed to discount phaeochromocytoma as the cause of the symptoms.

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