Treatment-resistant hypertension in a post-transplant patient with cystic fibrosis: a rare case of pheochromocytoma

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Summary

Pheochromocytoma is a rare catecholamine-producing tumour. We present the case of pheochromocytoma in a young man with a background history of a double-lung transplant for cystic fibrosis (CF). Clinical case: A 25-year-old man, with a background history of CF, CF-related diabetes (CFRD) and a double-lung transplant in 2012 was presented to the emergency department with crampy abdominal pain, nausea and vomiting. He was diagnosed with distal intestinal obstructions syndrome (DIOS). Contrast-enhanced CT imaging of the abdomen and pelvis showed a 3.4 cm right adrenal lesion. This was confirmed by a subsequent MRI of adrenal glands that demonstrated moderate FDG uptake, suggestive of a diagnosis of pheochromocytoma. The patient was noted to be hypertensive with a blood pressure averaging 170/90 mm/Hg despite treatment with three different anti-hypertensive medications – amlodipine, telmisartan and doxazosin. He had hypertension for the last 3 years and had noted increasingly frequent sweating episodes recently, without palpitations or headache. Laboratory analysis showed elevated plasma normetanephrines (NMN) of 3167 pmol/L (182–867) as well as elevated metanephrines (MN) of 793 pmol/L (61–377) and a high 3-MT of 257 pmol/L (<185). Once catecholamine excess was identified biochemically, we proceeded to functional imaging to further investigate. MIBG scan showed a mild increase in the uptake of tracer to the right adrenal gland compared to the left. The case was discussed at a multidisciplinary (MDT) meeting at which the diagnosis of pheochromocytoma was made. Following a challenging period of 4 weeks to control the patient's blood pressure with an alpha-blocker and beta-blocker, the patient had an elective right adrenalectomy, with normalisation of his blood pressure post-surgery. The histopathology of the excised adrenal gland was consistent with a 3 cm pheochromocytoma with no adverse features associated with malignant potential.

Learning points

- Five to ten per cent of patients have a secondary cause for hypertension. Pheochromocytomas are rare tumours, originating in chromaffin cells and they represent 0.1–1.0% of all secondary hypertension cases.
- Secondary causes should be investigated in cases where:
 - Patient is presenting <20 years of age or >50 years of age,
 - There is refractory hypertension, or
 - There is serious end-organ damage present.
- Patients may present with the triad of headache, sweating and palpitations or more vague, non-specific symptoms.

- Patients with suspected pheochromocytoma should have 24-h urinary catecholamines measured and if available, plasma metanephrines measured. Those with abnormal biochemical tests should be further investigated with imaging to locate the tumour.
- Medical treatment involves alpha- and beta-blockade for at least 2 to 3 weeks before surgery as well as rehydration.
- There is a possibility of relapse so high-risk patients require life-long follow-up.

Background

Pheochromocytoma is a rare tumour originating from catecholamine-secreting chromaffin cells that are derived from the ectodermic neural system and mostly situated within the adrenal medulla (1). This can result in catecholamine secretion in sudden bursts, leading to paroxysmal symptoms (2). It has an estimated incidence of less than 0.1% in the global population (3). However, their true incidence may be higher due to a lack of diagnosis until after death; a review of autopsy cases in Australia found that 0.05% had undiagnosed pheochromocytoma (4).

The presentation of pheochromocytomas vary from the classical symptom triad of palpitations, headaches and sweating to a spectrum of non-specific symptomatology, such as flushing, nausea, tiredness, weight loss or even no symptoms at all. All of these symptoms last from only minutes to hours and can occur periodically on different occasions (5). They are secondary to excess catecholamines. Pheochromocytomas differ considerably in rates of catecholamine synthesis, turnover, and release, and in the types of catecholamines and metabolites produced. These differences may explain variations in presenting signs and symptoms; they also can provide useful information about the tumour, including the adrenal or extra-adrenal location, the underlying mutation, tumour size, and the presence of metastatic disease (6).

Regularly described as the 'great mimic' or 'great masquerader', the sometimes non-specific and often intermittent nature of the symptoms caused by pheochromocytomas can result in its detection and diagnosis being missed by clinicians. Despite improved diagnostic techniques, there remains an approximate delay of 3 years between initial symptoms and the final diagnosis (7). This delay can lead to considerable morbidity and indeed mortality for patients.

We describe a case of an adrenal pheochromocytoma in a young man with CF, post lung transplant and multiple other co-morbidities. This case highlights the crucial importance of investigating secondary causes of hypertension.

Case presentation

A 25-year-old man, with a background history of CF, CFRD and a double-lung transplant in 2012 was presented to the emergency department with crampy abdominal pain, nausea and vomiting. He was diagnosed with DIOS, for which he was admitted for rehydration and laxatives. Contrast-enhanced CT imaging of the abdomen and pelvis showed a 3.4 cm right adrenal lesion, which was confirmed by a subsequent MRI of adrenals (Figs 1, 2A, B and 3A, B, C).

His background medical history was notable for CF for which he underwent a double-lung transplant in 2012 and CFRD. He also had a history of gastroesophageal reflux disease (GORD), for which he underwent a Nissen fundoplication in 2013, nasal polyps, previous episodes of DIOS, and he also had a low body mass density (BMD) for his age.

On review, the patient was noted to be hypertensive with a BP averaging 170/90 despite treatment with

Figure 1

CT of thorax abdomen pelvis (TAP). CT of abdomen and pelvis 28/08/19 (portal venous phase) – right adrenal mass measuring 3 × 2.5 cm. Had been present since 2015 with minimal interval change in size.

Figure 2

(A and B) PET CT. PET/CT 13/08/19 – carried out to assess the lymphadenopathy noted on MRCP. Sagittal view (left): anterior to the upper pole of the right kidney, there is an FDG-avid adrenal mass. Axial view (right).

maximum doses of three different anti-hypertensive medications – amlodipine, telmisartan and doxazosin. In the preceding 3 years, he had developed hypertension, in spite of significant increases in his anti-hypertensive medical therapy. He also complained of frequent sweating episodes over the past few months but denied any palpitations or headache.

Investigation

Biochemical: Our patient had a normal renal function, normal electrolytes, normal HbA1 and thyroid function. His liver function was abnormal with an alkaline phosphatase level of 1005 IU/L (30–130), ALT of 93 IU/L (0–55) and GGT of 684 IU/L (11–67). His albumin was normal at 36 g/L (35–50), as was his bilirubin at 5 μ mol/L (5–24). Further laboratory analysis showed an elevated plasma normetanephrines (NMN) level of 3167 pmol/L (182–867) as well as elevated metanephrines (MN) level of 793 pmol/L (61–377) and high 3-MT of 257 pmol/L (<185). His aldosterone was <138 pmol/L (138–670), his renin was 173.1 mIU/L (9.0–103.5) and his AM cortisol was 69 nmol/L (150–455), in the setting of taking long-term steroids post-transplant in 2012.

Radiological: The CT of abdomen pelvis carried out to work up his GI disturbance on admission showed a right adrenal mass measuring 3 $\times$ 2.5 cm that had been present since 2015 with minimal interval change in size (Fig. 1). This was confirmed by a subsequent MRI of adrenal glands. The PET CT showed that there was an FDG-avid adrenal mass, anterior to the upper pole of the right kidney (Fig. 2A). Finally, the MIBG scan showed only a mild increase in the uptake of tracer to the right adrenal gland compared to the left (Fig. 2B).

Treatment

Our patient's case was discussed at an MDT meeting at which, given the suggestive laboratory and radiologic findings, a diagnosis of pheochromocytoma was made. The consensus decision was for an elective right adrenalectomy to excise the pheochromocytoma.

The optimisation of blood pressure, as well as intravascular volume, is crucial to avoid perioperative adverse haemodynamic events, and we followed the pre-operative guidelines and vitals targets recommended by the Endocrine Society Clinical Guidelines Subcommittee for patients undergoing PCC and PGL resection (8).

Controlling our patient's BP proved challenging. He was initially treated with phenoxybenzamine 10 mg BD PO, however, it was required to up-titrate this to 10 mg TDS PO then 20 mg TDS PO over time due to persistent hypertension. We also started propranolol 10 mg TDS also up-titrating it to 20 mg TDS. Our target was for a BP of 110–100/50–70 with a heart rate of 50–70 bpm. The patient's underlying CF and DIOS may have contributed to malabsorption of these medications rendering pre-operative control of his blood pressure more challenging.

The surgery had to be postponed on one occasion, due to inadequate BP control, further exacerbated by spikes in his BP caused by the patient's abdominal pain. He was treated with regular analgesia and his alpha- and beta-blockade was further up-titrated. His eventual dose of phenoxybenzamine and propranolol was 70 mg QDS and 40 mg TDS, respectively. The peri-operative management included pre-hydration with IV NaCl at 100 mL/h for 24 h, increased oral salt intake, phentolamine IV as well as 100 mg hydrocortisone IV at surgical induction. An elective laparoscopic right adrenalectomy and lymph node biopsy was then carried out. Post-surgery, our patient was treated with hydrocortisone 50 mg IV TDS for 3 days

Figure 3

(A, B and C) MRI of the adrenal gland and MIBG: T2 hASTE sequence: heterogeneous mass in the right kidney. (B) out of phase (OOP). (C) In phase (IP). We compared the OOP to IP. If a lesion is full of fat and water (benign adenoma), it will be bright on IP but lose signal on OOP. This lesion stays the same on both OOP and IP. This can be seen in pheochromocytoma. MIBG: mild increase in the uptake of tracer to the right adrenal gland compared to the left.

Figure 4

(A) Excised pheochromocytoma on the right adrenal gland. The thin rim of adrenal cortex visible in areas. (B) Histological sample of excised pheochromocytoma: circumscribed tumour occupying adrenal medulla with overlying normal adrenal cortical cells.

as well as a supplementary s.c. insulin sliding scale with a target glucose of 6–10 mmol/mol. Our patient recovered well post-op, his blood pressure normalised completely, and he was discharged home 7 days post-surgery without any anti-hypertensive medication, with follow-up at the Endocrine and Transplant clinics.

Outcome and follow-up

Histology: The histopathology of the excised right adrenal gland showed a pheochromocytoma, encapsulated tumour, 30 mm in maximum dimension, composed entirely of nests of large polygonal cells with abundant granular cytoplasm, large nucleoli and prominent nuclear pleomorphism. Vascularity was a prominent feature throughout, but there was no necrosis. No vascular invasion or extension into surrounding tissue was seen (Fig. 4A and B). The PASS score showed a low risk for aggressive behaviour or malignancy with a value of 2/20

(<4/20). The mesenteric lymph nodes showed no evidence of lymphoma or lymphoproliferative disorder.

The patient returned to our Endocrine outpatient clinic in December 'feeling great'. His blood pressure had settled to 123/82 mm/Hg without any anti-hypertensive medication, and he denied any further episodes of sweating/flushing since the surgery. Repetition of biochemical analysis showed normal metanephrines of 72 pmol/L (61–377), normal normetanephrines of 94 pmol/L (182–867) and normal 3-MT of <65 pmol/L (<185).

Discussion

Our case of a young male patient with a history of hypertension who was presented with abdominal pain highlights the importance of screening for secondary aetiologies of hypertension in this population. The differential diagnosis includes renal artery stenosis caused by fibromuscular dysplasia, which is one of the most common secondary aetiologies, particularly in younger patients. Atherosclerotic renal artery stenosis is also a major cause of secondary hypertension in older adults. In middle-aged adults, aldosteronism is the most common secondary cause of hypertension, and the recommended initial diagnostic test is an aldosterone/renin ratio (9). Other differentials include Cushing's syndrome, thyroid dysfunction and obstructive sleep apnoea (10).

Our patient had a history of treatment-resistant hypertension for 3 years, despite being on 3 anti-hypertensive agents. His high blood pressure on admission was attributed to be secondary to the abdominal pain that he was experiencing. However, during the work-up, an abdominal CT showed a mass in the right adrenal gland. Based on these findings, phaeochromocytoma was suspected, which was subsequently confirmed by ancillary tests and pathology. Nausea and abdominal pain are recognised gastrointestinal symptoms of a phaeochromocytoma. Mullen *et al.* found that chronic constipation was noted in 13% of patients with phaeochromocytoma (11). Interestingly, our patient had multiple admissions to hospital for constipation and was diagnosed with DIOS, an issue which may have been exacerbated by both his cystic fibrosis and the catecholamine excess caused by the phaeochromocytoma. Our patient had a high 3-MT level, most commonly associated with extra-adrenal tumours, genetic mutations or malignant tumours. For this reason, we have sent our patient for genetic testing and continue his follow-up for life.

Phaeochromocytomas have been associated with a variety of cardiovascular complications due to overwhelming catecholamine levels. The most common complications include left ventricular hypertrophy, ischaemic heart disease, disturbances in rhythm and conduction, malignant ventricular arrhythmias, and even shock (12). The potentially fatal cardiovascular complications of these tumours are due to the potent effects of catecholamines, especially noradrenaline, the main transmitter released from sympathetic nerve terminals. Therefore, the longer that a diagnosis of phaeochromocytoma is missed, the greater the potential morbidity and mortality for these patients.

We describe a rare case of a right adrenal phaeochromocytoma in a young man with multiple co-morbidities, who completely recovered after tumour resection. This case highlights the crucial importance of investigating secondary causes of hypertension, even in patients with multiple comorbidities. The diagnosis of phaeochromocytoma requires high clinical alertness due to its rare incidence and extremely variable clinical presentation. It also highlights the relevance of considering phaeochromocytoma in the differential diagnosis of unexplained hypertension.

Declaration of interest

The authors declare that there is no conflict of interest that could be perceived as prejudicing the impartiality of the research reported.

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Patient consent

The author confirms that written informed consent has been obtained from the patient for publication of the submitted article and accompanying images.

Author contribution statement

All authors listed contributed equally to this paper and were the primary physicians involved in the care of the patient.

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