Isolated Pheochromocytoma in a 73-Year-Old Man With No Clinical Manifestations of Type 1 Neurofibromatosis Carrying an Unsuspected Deletion of the Entire NF1 Gene

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Pheochromocytomas (PHEOs) are a rare cause of endocrine hypertension that requires genetic counseling since at least 30% of PHEOs are associated with a germline mutation in a susceptibility gene. Neurofibromatosis type 1, NF1 is amongst the 16 known causing genes for pheochromocytomas/paragangliomas. We report a case of a 73-year-old man with PHEO in whom genetic testing revealed a large pathogenic heterozygous deletion of 1.14 Mb encompassing the entire coding sequence of the NF1 gene while the patient showed no signs of clinical NF1. This case illustrates that the diagnosis of NF1 should not be excluded in patients with PHEO in the absence of clinical diagnosis of the disease and support that older patients with PHEO should also be offered genetic counseling.

Keywords: neurofibromatosis, deletion, phenotype, pheochromocytoma, hypertension

INTRODUCTION

Pheochromocytomas (PHEOs) are rare catecholamine-secreting tumors that can cause endocrine hypertension with a prevalence of 0.4–2% in hypertensive patients (1). Germline mutations can be identified in at least 30% of patients with PHEOs and paragangliomas (2). In light of this, genetic testing is now recommended for all patients diagnosed with PHEO (3, 4). Some of these mutations can cause hereditary syndromes such as neurofibromatosis type 1 (NF1), von Hippel-Lindau (VHL), and multiple endocrine neoplasia type 2 (MEN2) (2). NF1 is a neurocutaneous disease caused by a mutation in the NF1 gene that encodes for the neurofibromin protein (5). The NF1 gene is located on the chromosome 17q11.2 (2). Two of the following criteria are needed to establish the diagnosis of NF1: inguinal or axillary freckling, two or more neurofibromas, six or more café-au-lait macules, two or more Lisch nodules, optic gliomas, distinctive bone lesions, or a first degree relative with NF1 (6). We report here the case of a man with pheochromocytoma that was diagnosed at 73 years-old carrying an unsuspected NF1 gene deletion but with no clinical manifestations of NF1.

Abbreviations: PHEO, Pheochromocytoma; NF1, Neurofibromatosis type 1.
CASE REPORT

Following coronary artery bypass grafting, a 73-year-old man presented with symptoms of palpitation, headache, sweating, and blurred vision. His initial systolic blood pressure was over 240 mmHg. The patient's medical history included type 2 diabetes, dyslipidemia, coronary heart disease, prostate cancer and long-standing hypertension that used to be well-controlled with 2 medications; Metoprolol 25 mg twice daily and Valsartan 80 mg daily. However, his hypertension had been harder to control for the past 2 years. His family history was unremarkable except for cases of sudden death for his maternal uncle and aunts. No familial syndromes were noted. Patient was worked up for severe postoperative hypertension with a computed tomography (CT) scan which revealed a small right adrenal mass measured at 31 mm (Figure 1). Biochemical testing included measurements of urinary fractionated metanephrines which revealed high normetanephrines with a value of 880 nmol/d (N < 240). However, metanephrines (148 nmol/d: N < 275), dopamine (936 nmol/d: N < 2570), epinephrines (undetectable: N < 110), and norepinephrines (456 nmol/d: N < 440) were within the normal limits. Plasma free normetanephrines were 2.10 nmol/L (N < 1.20) and free metanephrines 0.4 nmol/L (N < 0.48). The patient had normal calcitonin levels. A meta-iodobenzylguanidine (MIBG) scan showed intense and moderate uptake of a 2 × 2 × 3 cm mass of the right adrenal mass with no distant metastases. The patient underwent adrenalectomy and the diagnosis of PHEO was confirmed by histology post-operatively (Figure 2). PHEO of the Adrenal gland Scaled Score (PASS) was 4, and the ki67 proliferation index was less than one percent.

METHODS

After genetic counseling, the patient gave written consent for genetic analysis. Leucocyte DNA was obtained. A genetic panel of 14 susceptibility genes for PHEOs and paragangliomas was performed. The following genes were evaluated for sequence changes and exonic deletions/duplications (VHL, RET, NF1, SDHD, SDHB, SDHC, SDHA/SDHAF2, TMEM127, MAX, FH, EGLN1, KIF1B, MEN1) (Invitae, San Francisco, CA). Following the identification of a gross deletion, a comparative genomic hybridization (CGH) was performed using Agilent-CGXTM-HD, 4 × 180K oligonucleotide array (CHU Ste Justine, Unité Biologie Médicale: Génétique, Montréal, Quebec) to identify the boundaries of this genetic deletion. A written informed consent was obtained from the patient for the publication of this case report.
FIGURE 3 | Schematic representation of the genomic region at 17q11.2 harboring the NF1 gene and its flanking genes. The deletion of 1.14-Mb is shown with the blue bar and includes 11 OMIM genes (CRLF3, ATAD5, TEFM, ADAP2, RNF135, OMG, EVI2A, EVI2B, NF1, RAB11FIP4, MIR193A) and, 7 other genes (DPRXP4, MIR4733, MIR4724, MIR4725, MIR365B, COPRS, UTP6). Modified from Kehrer-Sawatzki et al. (7).

TABLE 1 | Clinical and biochemical descriptions of patients with PHEO and NF1 without clinical NF1 diagnosis described previously and including the current report.

| Sex       | Case 1 | Case 2 | Case 3 | Case 4 |
|-----------|--------|--------|--------|--------|
| Past medical history | Female | Hypertension | Prostatic adenocarcinoma, medullary thyroid cancer | None | Hypertension, prostate cancer |
| Pertinent family history | None | thyroid cancer—type unknown | 22 | None | 73 |
| Age at NF1 diagnosis (years of age) | 53 | 56 | 22 | |
| Age at PHEO diagnosis (years of age) | 39 and 53 | 56 | 22 | 73 |
| PHEO location | Bilateral | Bilateral | Left adrenal | Right adrenal |
| Presenting symptoms | Palpitation, sweating, flushed | Hypertension | Hypertension | Hypertension, palpitation, headache, sweating, and blurred vision |
| Functionality | Metanephrines and normetanephrines | Metanephrines, normetanephrines, epinephrine, norepinephrine, dopamine | Metanephrines, normetanephrines, and methoxytyramine | Normetanephrines |
| NF1 signs | Macrocephaly | On retrospective examination: Café-au-lait spots, axillary and inguinal freckling, Lisch nodules, neurofibromas and macrophagy | Short stature | On retrospective examination:pectus excavatum |
| NF1 Mutation | *c.6084+1G>A—A splice-site mutation | *c.7206_7207del (p.His2402Glnfs*4) | *c.7846C>T, p.Arg2616* | Deletion of 15 oligonucleotides 1.14Mb at 17q11.2.Including the entire coding sequence of NF1 |

*NM_000267.3.

RESULTS

Panel genetic testing revealed a gross heterozygous germline deletion of the genomic region encompassing the full-coding sequence of the NF1 gene (NM_000267.3). In order to better identify the boundaries and the extends of this event CGH array was performed. The analysis showed a deletion of 15 oligonucleotides at 17q11.2 (position 29, 116, 494-30, 260, 501) (Genome Browser UCSC 2009hg19 assembly). The size of this large pathogenic heterozygous deletion is estimated at 1.14 Mb. In addition to the NF1 gene, this deletion includes 10 OMIM genes: CRLF3, ATAD5, TEFM, ADAP2 RNF135, OMG, EVI2A, RAB11FIP4, MIR193A and 7 other genes: DPRXP4, MIR4733, MIR4724, MIR4725, MIR365B, COPRS, UTP6 (Figure 3). Patient was re-examined in light of these findings but showed no signs of clinical neurofibromatosis type 1 except for a pectum excavatum that may be associated with NF1.

DISCUSSION

To our knowledge, we report the oldest patient carrying a large deletion encompassing the entire NF1 gene in the context of an
apparently sporadic pheochromocytoma. Very recently, a case series described three patients with PHEOs carrying unsuspected NF1 mutations (2). In contrast to our case, these patients were younger (22, 39, and 56 years) and carried only point mutations in the NF1 gene rather than the loss of the entire gene. In this report, only one of the three patients had discreet signs of NF1 on a repeat examination (2) (Table 1). This shows that a clinical examination is insufficient to exclude a NF1 diagnosis in a case of PHEO.

In our case, the tumor secreted normetanephrines. NF1 cases of PHEO have been shown to secrete metanephrines or normetanephrines (2). Gross deletions of the NF1 genes have been described previously in individuals with neurofibromatosis type 1 (8, 9). Large deletions of the NF1 gene and its flanking regions are frequently associated with a more severe clinical phenotype of NF1 (dysmorphic features, overgrowth, hypotonia, and cognitive impairment) that was not present in this patient (7).

NF1 is a heterogeneous disorder affecting ~1/3,000 births (10). This disease causes multi-organ involvement, which can include PHEOs in 0.1–7% (2, 10). The incidence might be underestimated due to the lack of formal recommendations to universally screen biochemically NF1 patients for PHEOs, as there are in other genetic syndromes such as Von-Hippel-Lindau or MEN2 (10, 11). Recent data showed that 31% of PHEOs in NF1 are found incidentally and only 7.3% were diagnosed following biochemical testing (12). Moreover, PHEOs tend to be diagnosed at an older age in patients with NF1 when compared to other genetic mutations (48 vs. 30 years old) (10).

Diagnosis of NF1 is important as it is associated with an elevated risk of associated malignancies, including malignant peripheral nerve sheath tumors, breast cancer, gastrointestinal stromal tumors, brain, and central nervous system tumors (13). This entails these patients to a more rigorous screening and active surveillance. Furthermore, recognition of NF1 is important for familial screening. In addition to de novo mutations NF1 may be inherited (14). The patient reported here did not have a family history of NF1. However, NF1 being an autosomal dominant disorder, appropriate genetic counseling for family members is essential (15). Overall, patients with NF1 need a medical follow-up by a multidisciplinary team to assess and manage complications, counsel on genetic screening (6).

Our findings support the recommendation of more recent guidelines that all patients, independently of age and the clinical presentation of PHEO, should be offered genetic counseling (3, 4). This is illustrated in this 73 year-old man carrying an unsuspected major NF1 deletion. Moreover, genetic testing should not only include evaluation for gene sequence changes but also gene deletion/duplication. Finally, the diagnosis of NF1 should not be excluded in patients with PHEO in the absence of clinical diagnosis of the disease.

DATA AVAILABILITY

This manuscript contains previously unpublished data. The name of the repository and accession number(s) are not available.

AUTHOR CONTRIBUTIONS

SP-L collected the data and wrote the manuscript with the support of IB. ND genetic counselor that met the patient. GR endocrinologist that was involved in patients’ care. IB supervised the project and wrote the manuscript with the support of SP-L. ML pathologist that analyzed the specimen and provided the pictures of the tumors' histology. All authors discussed the results and contributed to the final manuscript.

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