Isolated Familial Pheochromocytoma as a Variant of von Hippel-Lindau Disease


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ABSTRACT

Inherited pheochromocytomas are often part of familial syndromes, especially multiple endocrine neoplasia type 2 (MEN 2) and von Hippel-Lindau (vHL) disease. In a family with pheochromocytomas in three generations and with at least seven affected members, we investigated by clinical and genetic analyses the presence or absence of associated conditions. The clinical investigations included ophthalmological and radiological studies for von Hippel-Lindau disease (magnetic resonance imaging of the brain, computed tomography of the abdomen, and direct ophthalmoscopy after mydriasis) and annual calcitonin stimulation tests for C cell disease in five members who agreed to regular follow-up.

Besides the pheochromocytomas (so far, these have been multiple in five of seven individuals) no definite second associated condition was found. Genetic analysis did not identify any MEN 2-specific RET protooncogene point mutations (which are present in 97% of MEN 2 families). However, despite the complete absence of other clinical manifestations of the vHL disease (besides pheochromocytomas), a previously undescribed germline missense mutation in the vHL tumor suppressor gene was found (C775G transversion with a predicted substitution of a leucine by a valine at codon 259 in the putative vHL protein).

We conclude that in this family the sole occurrence of pheochromocytoma is a variant of vHL disease. (J Clin Endocrinol Metab 81: 1035–1037, 1996)

Approximately 10% of pheochromocytomas are believed to be inherited (1). Most of them are part of familial syndromes with an autosomal dominant mode of inheritance and other associated conditions, especially multiple endocrine neoplasia type 2 (MEN 2) and von Hippel-Lindau (vHL) disease. In a family with pheochromocytomas in three generations and with at least seven affected members, we investigated by clinical and genetic analyses the presence or absence of associated conditions. The clinical investigations included ophthalmological and radiological studies for von Hippel-Lindau disease (magnetic resonance imaging of the brain, computed tomography of the abdomen, and direct ophthalmoscopy after mydriasis) and annual calcitonin stimulation tests for C cell disease in five members who agreed to regular follow-up.

Besides the pheochromocytomas (so far, these have been multiple in five of seven individuals) no definite second associated condition was found. Genetic analysis did not identify any MEN 2-specific RET protooncogene point mutations (which are present in 97% of MEN 2 families). However, despite the complete absence of other clinical manifestations of the vHL disease (besides pheochromocytomas), a previously undescribed germline missense mutation in the vHL tumor suppressor gene was found (C775G transversion with a predicted substitution of a leucine by a valine at codon 259 in the putative vHL protein).

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Subjects and Methods

Patients

Since 1988 we have followed a family with pheochromocytomas (the pedigree is given in Fig. 1). The minimal annual examination included a medical history; physical examination; measurement of urinary excretion of vanilmandelic acid, total metanephrines, norepinephrine, and epinephrine; and measurement of serum total calcium concentration and basal and stimulated calcitonin concentrations. With the exceptions stated below, morphological studies were performed in all family members with suspected or proven pheochromocytomas. This included magnetic resonance imaging of the brain, computed tomography of the abdomen, [131]I metaiodobenzylguanidine scintigraphy, direct ophthalmoscopy after mydriasis, and ultrasonography of the testes. These studies were performed using standard techniques and with special attention to the manifestations of MEN 2 and vHL disease. All studies were conducted after informed consent had been obtained.

Biochemical analyses

In 24-h urine collections, epinephrine, norepinephrine, metanephrine, normetanephrine, and vanilmandelic acid were measured, as de-
squares denote males. The right half of each symbol indicates the presence (solid area) or absence (open area) of left-sided pheochromocytoma (and vice versa); the middle portion denotes extraadrenal pheochromocytoma.

Calcitonin stimulation tests were performed by combined stimulation with iv calcium and pentagastrin (6). Measurement of calcitonin was made with an immunoradiometric assay from Medgenix (Fleuris, Belgium). From 1988–1989, the test was considered positive if the stimulated value exceeded 120 pg/mL. In 1990, we established new reference values (7), and stimulated calcitonin values up to 400 pg/mL were considered normal.

Genetic analyses

MEN 2. Genomic DNA was isolated from peripheral leukocytes using an automated DNA extractor, as previously described (8). One hundred nanograms of DNA were PCR amplified for exons 10 (primer CRT 19s and CRT 2C or CRT 19s and CRT 19E), 11 (CRT 19s and 2C or CRT 19B and 2C), and 16 (CRT 5G and CRT 5H or F RET 16 and R RET 16). Twenty-five-microliter samples contained 10 mmol/L Tris-HCl (pH 8.3), 50 mmol/L KCl, 0.75 mmol/L MgCl₂, 0.01% gelatin, 200 pmol/L deoxynucleotides (dNTPs), and 1 U Tag polymerase with 1 µmol/L each of the appropriate PCR primers. DNA was amplified for 40 cycles of 1 min each at 95, 55, and 72°C. PCR products were separated by electrophoresis on 2% low melting temperature agarose gels and eluted using the Magic Prep system (Promega, Southampton, UK). Purified amplification products were sequenced using the delta Taq cycle sequencing kit (U.S. Biochemical Corp., Cleveland, OH). The sequences were resolved on 5% denaturing polyacrylamide gels.

vHL disease. Screening for vHL gene mutations was performed as described previously. Briefly, the cloned coding sequence of the vHL gene was amplified from genomic DNA using six sets of primer sequences, and each fragment was examined for point mutations by single strand conformational polymorphism analysis (9). The nucleotide sequence of PCR products showing band shifts on single strand conformational polymorphism analysis was determined by direct sequencing with nested primers, using a commercially available cycle sequencing kit (Life Technologies, Grand Island, NY) according to the manufacturer’s guidelines.

Results

Patients

With respect to the pedigree shown in Fig. 1, patient I,1 refused clinical investigations, with the exception of an ultrasonography that demonstrated bilateral adrenal tumors but normal renal and thyroid morphology. She also agreed to determination of the basal calcitonin level, which was normal, and a genetic analysis.

Patient I,1 died in childbirth with the symptoms and signs of a hypertensive crisis. Whole body autopsy showed a left adrenal and an extraadrenal pheochromocytoma, but no other features of MEN 2 or vHL disease. The child died 1 day after birth from a cerebral hemorrhage.

Patient I,3 claimed that after performing an ultrasonography her family doctor had told her that she was not affected by the family disease. She agreed only to the genetic analysis.

All other patients (II,2; II,3; II,4; II,5; and III,1) were screened for MEN 2 and vHL disease. With one exception all results were normal. In patient II,2, two calcitonin stimulation tests were performed, which demonstrated increases from a normal basal value (37 and 39 pg/mL) to 141 and 295 pg/mL. At that time our upper normal value for calcitonin during the stimulation test was 120 pg/mL, and we considered this increase to be indicative of C cell hyperplasia. After extensive discussion with the patient, thyroidectomy was performed despite normal ultrasonography of the thyroid. Only a focal presence of C cells, but no medullary thyroid carcinoma, was found. Postoperatively, calcitonin was no longer detectable (<10 pg/mL basally and after stimulation).

Genetic analyses

We were not able to identify any MEN 2A-specific RET point mutations in genomic DNA of the family members tested at codons 609, 611, 618, 620, and 634 in exons 10 and 11, respectively. Furthermore, no mutations were found in exon 16 (codon 918).

Examination of the vHL gene identified a C to G transversion at nucleotide 775 of the published coding sequence (10). This sequence change was predicted to cause a substitution of a leucine by a valine at codon 250 in the putative vHL protein. The C775G change segregated with the disease and has not been detected in analysis of more than 400 normal vHL alleles.

Discussion

We describe a family suffering from familial pheochromocytoma with seven affected subjects in three generations. In the absence of cutaneous manifestations and, therefore, exclusion of neurofibromatosis type 1, we had to differentiate among the diagnoses of MEN 2, retinal cerebellar hemangioblastomatosis, and isolated familial pheochromocytoma.

The diagnosis of MEN 2 appeared unlikely because none of the affected family members (at 14, 20, 29, 42, 45, 59, and 71 yr of age) suffered from overt thyroid disease, and an adequately and repeatedly performed screening for medullary thyroid carcinoma gave normal results in 4 of 5 patients. In 25 patients, members of 5 MEN 2 families, we have found no patient with a pheochromocytoma or primary hyperparathyroidism without C cell disease (unpublished data). However, Gagel et al. (11) described adrenal disease preceding by 5 and 8 yr the detection of C cell abnormalities in 2 of 19 MEN 2 patients with pheochromocytoma. Thus, the detection of adrenal disease as the first manifestation in MEN 2 patients is occasionally possible; however, the absence of thyroid disease 24 yr (patient II,4) and 12 yr (patient II,5) after detecting pheochromocytoma would have been unprecedented.

One patient (II,2) had a calcitonin stimulation test that was considered abnormal. Consequently, thyroidectomy was performed. At that time, we used an upper normal limit for calcitonin during the stimulation test of 120 pg/mL. Subsequently, based on tests in 20 healthy volunteers, we established an upper normal limit for stimulated calcitonin of 400 pg/mL (7). The histological evaluation in this patient showed only focal C cell hyperplasia, but no medullary thyroid carcinoma. Focal C cell
hyperplasia can represent a normal finding in the thyroids of children less than 6 yr of age and in elderly patients (12). Recently, positive stimulation tests and C cell hyperplasia in the operated thyroid were described as a normal finding in unaffected members of MEN 2 families (13). Thus, we now judge the calcitonin stimulation test performed preoperatively on patient II.2 as normal and the histological findings as not proven for MEN 2 associated C cell disease.

Mutations in the RET protooncogene were identified as the genetic defect in MEN 2 (14, 15). Based on this experience, genetic analysis of the family was carried out. RET protooncogene mutations in exons 10 and 11 are present in 97% of MEN 2A families and in 86% of the patients with familial medullary thyroid carcinoma (16). In almost all MEN 2B patients tested recently, mutations in exon 16 were detectable (17).

We were not able to detect any MEN 2-specific RET protooncogene mutations in exons 10, 11, and 16 and concluded that the diagnosis of MEN 2 was extremely unlikely. Therefore, calcitonin stimulation tests were no longer needed in the care of this family.

At about the time of the discovery of the MEN 2 gene, rearrangements in a gene named vHL disease tumor suppressor gene were identified as the causal defect for vHL disease (10). Therefore, a search for a defect in this gene in the described family was performed. Despite the complete absence of other clinical manifestations, a mutation in the vHL tumor suppressor gene was found.

The positive finding of a mutation in this gene obviates the need for further testing in the RET protooncogene, as additional mutations in 3% of MEN 2a families were found that do not harbor one of the common mutations in codons 609, 611, 618, 620, and 634 (these showed only the wild-type allele in this family) (18).

The presence of pheochromocytoma in 20% of vHL families is often used to classify the disease into two types; families without pheochromocytoma are described as vHL type 1, and those with pheochromocytoma as vHL type 2 (19–21). Interesting correlations between this phenotype expression and the types of mutations in the vHL disease tumor suppressor gene have been reported (9, 21). In short, missense mutations (in contrast to deletions, insertions, and nonsense mutations) are found significantly more often in families with vHL type 2.

The family described in this manuscript is the first reported family with a mutation in the vHL disease tumor suppressor gene, but with pheochromocytoma as the only manifestation of vHL disease (i.e., without angiomas, hemangioblastomas, and renal cell carcinomas). Therefore, the family cannot be assigned to one of the two above-mentioned types of vHL disease. The C775G mutation detected in this family has not been described previously, but it is interesting to note that it is also a missense mutation. Furthermore, the mutation identified is at the very 3'-end of the vHL disease tumor suppressor gene, and the substitution of leucine by valine is supposed to cause little conformational change in the putative vHL protein. The absence of other vHL manifestations in all seven members of the family is unlikely to be coincidental given the prevalence and mean age at onset of the other clinical features of the disease (22). Therefore, the association of this particular mutation with such a pheochromocytoma-only variant of vHL disease might contribute to an understanding of the genetic mechanisms of vHL disease.

In conclusion, we have shown that in this family, the sole occurrence of pheochromocytoma is a minor variant of vHL disease. To emphasize both the relationship to and the distinction from the full blown syndrome, we propose that this pheochromocytoma-only variant be called vHL type 3.

Note added in proof:
Since the writing of the manuscript, the numbering of the vHL coding sequence has been changed: codon 259 is now counted as codon 188 (Il-

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Acknowledgments
We are indebted to W. Mraz, M.D., and K. Jacob, Ph.D., Institute for Clinical Chemistry at the Klinikum Grosshadern, for determinations of calcitonin and urinary catecholamines.

References