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Metastases but not cardiovascular mortality reduces life expectancy following surgical resection of apparently benign pheochromocytoma

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Abstract

The treatment of choice for non-metastatic pheochromocytoma is surgical resection. Its goals are to abolish catecholamine hypersecretion, normalize blood pressure, and prevent further tumor growth or progression to metastatic disease. Data on long-term mortality and morbidity after pheochromocytoma surgery are limited. We here report a retrospective study on the long-term outcome after surgery for apparently benign pheochromocytoma at the Radboud University Nijmegen Medical Centre. Data on clinical presentation, treatment, post-surgical blood pressure and recurrence, metastasis and death were collected of 69 consecutive patients (January 1966–December 2000; follow-up: until death or January 2006). Survival was compared with survival of a matched reference population. Two patients died of surgical complications. All ten patients with metastatic disease (including three diagnosed at first surgery) died. At follow-up, 40 patients were alive and recurrence free and three patients were lost to follow up. Two patients experienced a benign recurrence. Mean ± s.d. follow-up was 10.2 ± 7.5 (median 9, range 1–38) years. Kaplan–Meier estimates for 5- and 10-year survival since surgery were 85.8% (95% CI: 77.2–94.4%) and 74.2% (95% CI: 62.0–86.4%) for patients versus 95.5 and 89.4% in the reference population (P < 0.05). Sixty-four percent of all patients with hypertension prior to surgery showed a significant decrease in blood pressure, but remained hypertensive after surgery. In conclusion, compared with the general population patients have a reduced life expectancy following pheochromocytoma surgery, due to their risk of developing metastatic disease. Only one-third becomes normotensive without antihypertensive medication. Therefore, lifelong follow-up is warranted.

Endocrine-Related Cancer (2008) 15 1127–1133

Introduction

Pheochromocytoma (PHEO) is a rare catecholamine producing tumor that derives from chromaffin cells of the adrenal medulla or extra-adrenal abdominal and thoracic paraganglia (DeLellis et al. 2004, Lenders et al. 2005, Pacak et al. 2005). Symptoms, signs and complications of PHEO are mainly related to hypersecretion of catecholamines. These include headache, palpitations, diaphoresis, and hypertension. Once the biochemical diagnosis of PHEO is established by plasma and/or urine measurements of catecholamines and their metabolites (Lenders et al. 2002), tumors are localized by anatomical or functional imaging, such as ¹²³I-MIBG scintigraphy (Ilias et al. 2005).
Treatment of non-metastatic PHEO consists of surgical resection following preoperative pharmacological sympathetic blockade (van der Horst-Schrivers et al. 2006). The aims of surgery are abolishment of catecholamine hypersecretion, normalization of blood pressure, and prevention of further tumor growth and progression to metastatic disease. The long-term outcome of patients after PHEO surgery has been addressed in a limited number of studies. Estimates of benign and malignant recurrence rate after surgery vary between 6 and 23% (Modlin et al. 1979, Scott & Halter 1984, van Heerden et al. 1990, Pruszczyk et al. 1991, Plouin et al. 1997). Conflicting results of previous cohort studies suggest that the life expectancy after PHEO surgery is either normal (Stenstrom et al. 1988) or decreased (Khorram-Manesh et al. 2005) when compared with the general population. Furthermore, though PHEO is generally considered to be a surgically curable cause of secondary hypertension, blood pressure remains elevated after surgery in a considerable number of patients without recurrent disease (Plouin et al. 1997, Khorram-Manesh et al. 2005).

The aim of this study was to investigate the long-term follow-up of patients after surgery for apparently benign PHEO. Survival was compared with results obtained from a period-, age-, and gender-matched Dutch reference population.

**Methods**

**Patients**

To evaluate long-term outcome, the study was restricted to patients diagnosed with PHEO before January 2001. From January 1966 until December 2000, 71 patients with a histologically confirmed diagnosis of PHEO were evaluated at the departments of General Internal Medicine and Endocrinology at the Radboud University Nijmegen Medical Centre. In all but four patients, initial PHEO surgery was performed at our centre. Two patients were excluded from the study: one patient was referred to our center because of metastases and in a female patient who died during Caesarian section the PHEO was found at autopsy. This left 69 patients (44 females, 25 males) for evaluation of follow-up after surgery for apparently benign PHEO.

**Initial work up, surgery, and follow-up**

For the biochemical diagnosis of PHEO, plasma and 24-h urine were assayed by HPLC for concentrations of catecholamines (plasma $n=59$, urine $n=38$) and urine fractionated metanephrines ($n=12$), according to previously described procedures (Willemsen et al. 1995, 2003). CT ($n=57$), MRI ($n=4$), abdominal ultrasound ($n=36$) and $^{123}$I-MIBG scintigraphy ($n=42$) were employed to localize the tumor(s). Except for three laparoscopic adrenalectomies, tumors were resected by open procedures (including four that initially started laparoscopically). Five patients with bilateral tumors underwent staged open surgery. Surgery was performed before 1990 in 43 per cent (29 out of the 68) of patients. Plasma and urine concentrations of catecholamines and metanephrines were reassessed after a minimum of 1 month post-surgery.

**Data collection**

Data on clinical presentation, biochemical, radiological and histological diagnosis, and surgical and pharmacological treatment were retrospectively collected from the patients' medical files in a standardized fashion. Post-surgery follow-up data, including blood pressure values, were collected until the end of December 2005. In patients whose most recent visit to our clinic took place before January 2005, information on the current health status, blood pressure, and antihypertensive treatment was updated by telephone interviews of the patients and their primary physicians.

**Definitions**

Hypersecretion of (nor)epinephrine was defined as elevation of plasma and/or urine levels of (nor)-epinephrine and/or its metabolite (nor)metanephrine above the upper range of normal. Malignancy was defined as histological evidence of tumor cells at sites where chromaffin tissue is normally absent and/or by evidence of distant metastases documented by imaging studies (Linnoila et al. 1990). Recurrence was defined as the reappearance of the disease as documented at reintervention or by combined biochemical and imaging tests, after complete eradication of the tumor. Hypertension was defined as repeated office readings of systolic and/or diastolic blood pressure exceeding 140/90 mmHg and/or the use of antihypertensive medication.

**Data analysis and statistics**

Survival of patients who underwent successful resection of apparently benign PHEO was estimated using the Kaplan–Meier method and compared with the life expectancy of a Dutch reference population that was matched for age, sex, and period. Demographic data
Results
Clinical presentation and diagnosis

Signs and symptoms at presentation are summarized in Table 1. Symptoms occurred episodically in 82% (55 out of the 70) of patients, with a frequency of at least daily in 42% (23 out of the 55). In two patients, an adrenal tumor was found incidentally during work up of colon carcinoma and chest pain respectively. Seventy-seven percent (54 out of the 70) of patients had hypertension. Apart from pre-surgical α- and β-blockade, 64% (43 out of the 67) were on antihypertensive drugs. Age at initial diagnosis of PHEO was 46.1 ± 15.6 (range 21–78) years. Hypersecretion of norepinephrine only occurred in 35% (20 out of the 57), epinephrine only in 7% (4 out of the 57), and both epinephrine and norepinephrine in 58% (33 out of the 57) of patients. PHEOs were located as follows: left adrenal (n = 33), right adrenal (n = 22), bilateral adrenal (n = 6), and extra-adrenal (n = 8: 5 retroperitoneal, three in the bladder). Ninety-three percent (43 out of the 46) of tumors detected by anatomical imaging (CT, MRI, and ultrasound) were positive on 123I-MIBG scintigraphy. Histological findings in all tumors were consistent with PHEO. Fourteen patients had an underlying familial tumor syndrome: five neurofibromatosis, eight multiple endocrine neoplasia (MEN) type 2A, and one MEN type 2B.

Follow-up

Of the 69 patients, two died within 4 weeks after surgery (Table 2). In three patients, metastatic lesions were detected intra-operatively. The remaining 64 patients were included in the follow-up analysis. Follow-up of these patients was 11.0 ± 7.4 (median 9, range 1–38) years. In 20 patients, the most recent clinical visit took place more than one year before the final analysis (median 5, range 1–7 years). Three patients were lost to follow-up at 2, 8, and 27 years after surgery.

Two patients experienced a benign recurrence (one with MEN 2B). In one patient, initial surgery was incomplete, and plasma catecholamines only normalized after resection of the rest tumor.

Seven additional patients developed metastases, within 1–14 years after the initial diagnosis (Table 3). There were no significant differences between patients with metastatic and non-metastatic PHEO with respect to sex, age at diagnosis, pre-surgical plasma catecholamine levels, and extra-adrenal tumor location (Table 4). All patients with metastases died. Survival since the diagnosis of metastases was 2.6 ± 0.8 (range 0–6) years. Twelve additional patients died during follow-up, all of whom were considered disease free although the cause of death is unknown in 5 out of these 12 patients (Table 3).

Survival

For the 64 patients who underwent successful surgery for non-metastatic PHEO, Kaplan–Meier estimates showed 5- and 10-year survival rates of 85.8% (95% CI:

<table>
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<th>Table 1 Symptoms and signs at initial presentation</th>
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<td>Symptoms and signs</td>
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<td>Headache</td>
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<td>Palpitations</td>
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<td>Diaphoresis</td>
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<td>Pallor</td>
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<td>Anxiety/panic attacks</td>
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<td>Weight loss</td>
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\[\text{Lenders et al. 2005.}\]
and at most recent post-surgical follow-up, mean blood pressure had decreased from $164\pm28/95\pm12$ to $143\pm21/87\pm11$ mmHg ($P<0.001$ for systolic and diastolic blood pressure).

Plasma catecholamine levels normalized after surgery, except in three patients with slightly elevated levels, despite lack of recurrence on imaging studies.

### Discussion

The main finding of our study is that patients undergoing successful surgery for apparently benign PHEO have a reduced life expectancy when compared with a sex, age, and period-matched reference population. This can be ascribed fully to the patients who developed metastatic disease while survival was not reduced in the really benign patients. We also show that only one-third of recurrence-free patients become and remain normotensive on the long-term following surgery.
Only a limited number of publications of the past 10 years provided long-term follow-up data on morbidity and mortality after PHEO surgery (Plouin et al. 1997, Goldstein et al. 1999, Amar et al. 2005, Khorram-Manesh et al. 2005). Differences in study population, length of follow-up, definitions of recurrent and malignant disease, and analysis methods hamper comparison of these studies. The rates of benign recurrence after successful surgery of non-metastatic PHEO ranged from 3.3 to 6.1% (Plouin et al. 1997, Amar et al. 2005, Khorram-Manesh et al. 2005), versus 3% in the present study. The prevalence of malignant disease, ranged from 6.6 to 16% (Plouin et al. 1997, Goldstein et al. 1999, Amar et al. 2005; Khorram-Manesh et al. 2005), versus 14% in the present study. Our observation of a 5-year metastases-free survival of 88% after successful surgery for really benign PHEO is comparable with that of 92–94% found in these studies (Plouin et al. 1997, Amar et al. 2005).

Since there are no established pathology criteria to distinguish benign from malignant PHEO, clinical, biochemical, and temporal characteristics have been evaluated as surrogate predictors of malignant recurrence. Features that were implied to be associated with shorter PHEO-free survival after surgery include extra-adrenal tumor location, low epinephrine/epinephrine + norepinephrine ratio (Plouin et al. 1997, Amar et al. 2005, Table 4 Benign versus malignant PHEO

<table>
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<th>Benign PHEO</th>
<th>Malignant PHEO</th>
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<tr>
<td>N</td>
<td>59</td>
<td>10</td>
</tr>
<tr>
<td>Sex (M/F)</td>
<td>19/40</td>
<td>6/4</td>
</tr>
<tr>
<td>Age at diagnosis (y)</td>
<td>45.2±15.3</td>
<td>51.1±16.3</td>
</tr>
<tr>
<td>Pre-surgery E/(E+NE) ratio</td>
<td>0.21±0.22</td>
<td>0.14±0.24</td>
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<tr>
<td>Extra-adrenal tumor</td>
<td>12% (7/59)</td>
<td>10% (1/10)</td>
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Benign: no metastatic lesions during follow-up. Malignant: metastatic lesions during follow-up. M, male; F, female; E, epinephrine; NE, norepinephrine.

**Figure 1** Kaplan–Meier curve of survival after pheochromocytoma surgery of 64 patients with apparently benign disease compared with the Kaplan–Meier survival curve of a matched reference population.

**Figure 2** Kaplan–Meier curve of survival after pheochromocytoma surgery of ten patients (eventually) diagnosed with metastatic disease compared with the Kaplan–Meier survival curve of a matched reference population.

**Figure 3** Kaplan–Meier curve of survival after pheochromocytoma surgery of 57 patients with benign disease compared with the Kaplan–Meier survival curve of a matched reference population.
the long term (Plouin et al. 2005). All our patients were operated before 2001, mainly through open procedures. Secondly, the prognosis of patients with PHEO is largely determined by the underlying genotype. Genetic testing was not systematically performed in all patients, partly due to the fact that comprehensive genetic testing, including SDHB mutations, became available after 2001. The observed prevalence of disease-causing mutations of 20% might, in fact, be an underestimation, and the results may theoretically be influenced by the portion of unrevealed mutations.

In conclusion, patients who undergo successful surgery for apparently benign PHEO have a reduced life expectancy when compared with the general population. The excess mortality is explained by the development of metastatic disease during post-surgical follow-up in some of the patients. By contrast, the survival of patients who do not develop metastatic disease is not different from that of the general population. Hypertension persists in two-thirds of recurrence-free patients. Lifelong follow-up of patients who had surgery for PHEO is warranted.

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.

Funding
This research did not receive any specific grant from any funding agency in the public, commercial or not-for-profit sector.

Acknowledgements
The authors regret to announce that AAJ Verhofstad is now deceased.

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