CLINICAL STUDY

The outcome of surgery in 668 patients with acromegaly using current criteria of biochemical ‘cure’

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Abstract

Background and aim: The aim of this study was to illustrate the present role of transsphenoidal surgery as primary therapy in GH-secreting adenomas, and to compare the results concerning control of disease with previous series using older criteria of cure.

Method: We report on a consecutive series of 668 acromegalic patients treated over a time period of 19 years. Biochemical cure was defined as normalisation of basal GH level, suppression of GH levels to below 1 ng/ml during an oral glucose load and normalisation of IGF-I levels. Of the 506 patients undergoing primary transsphenoidal surgery, a total of 57.3% postoperatively fulfilled the criteria used.

Results: The rate of biochemical ‘cure’ correlated with the magnitude of the initial GH levels, the tumour size and invasion. The overall complication rate was below 2%. Mortality in this series was 0.1% (1 of 688). During a follow-up period of 10.7 years only two recurrences (0.4%) occurred. However, in the patients treated by transcranial surgery and by repeat surgery the cure rate was found to be relatively low (5.2 and 21.3% respectively).

Conclusions: These data suggest that surgery remains with very few exceptions the primary treatment of acromegaly for (i) a high cure rate, (ii) low morbidity, (iii) low recurrence rate and (iv) immediate decline of GH. Based on current criteria of cure, recurrences are uncommon. However, cure by surgery alone is improbable in patients harbouring extended, invasive tumours with high secretory activity, in whom further adjuvant treatment is mandatory.

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Introduction

Acromegaly is a rare but very serious condition usually caused by a growth hormone (GH)-producing pituitary adenoma. Ectopic GH or GH-releasing hormone production or a pituitary carcinoma may very infrequently also cause acromegaly. Several retrospective cohort studies suggest that mortality in acromegaly is at least twice of that in the general population (1–4). The cause of death is most commonly a vascular accident, but significant increases have been reported for both respiratory disorders and malignancies. These studies have also demonstrated that a reduction of GH levels significantly lowered mortality and that mortality rates similar to those of the general population are restored once remission is induced. Partial reduction of GH levels has been proven to be inadequate to restore normal life expectancy (1, 5). Therefore, aggressive management to lower serum GH levels is necessary once the diagnosis has been confirmed. Depot somatostatin analogues are now increasingly being prescribed as adjuvant or even primary therapy for the treatment of acromegaly. Studies have shown them to be both effective and safe with endocrinological remission achieved in up to 60% of cases (6–9). The newly developed GH receptor antagonist pegvisomant seems to be able to normalise insulin-like growth factor (IGF)-I in up to 97% of patients and may become an important therapeutic option in acromegaly (10, 11). However, the most recent consensus guidelines for the management of acromegaly suggest for the majority of patients surgery as the first-line therapy, either alone or in combination with medical treatment, conventional radiotherapy and/or radiosurgery (12). The aim of this study was to update the surgical results and the long-term outcome of the largest-ever reported consecutive series of acromegalic patients treated in the Department of Neurosurgery at the University of Erlangen-Nuremberg using the current internationally accepted criteria for biochemical ‘cure’ of the disease and to compare the results concerning control of disease with previous series using older criteria of cure.
Patients and methods

Patient series

Between December 1982 and December 2001, 688 operations (514 primary transsphenoidal, 28 primary transcranial and 146 secondary (repeat) surgery) in 646 patients harbouring GH-secreting pituitary adenomas were performed in the Department of Neurosurgery of the University of Erlangen-Nürnberg, the majority of them by the senior author (F.R.). Of these, 634 patients presented with signs and symptoms of acromegaly and 12 with gigantism, ranging in age from 12 to 76 years (mean 42.3±10.6 s.d. years). There were 338 females and 306 males (ratio female: male 1.1:1.0). However, patients lacking complete preoperative endocrinological data (n = 10), patients mainly referred from abroad, who were lost to follow-up (n = 5), and the patient who died following transsphenoidal surgery, were not included in the evaluation of the endocrinological outcome. Evaluation of the surgical results was performed on 506 patients following primary transsphenoidal, 26 patients following primary transcranial and 140 patients following secondary surgery (672 surgical procedures in 630 patients). The period of follow-up monitoring ranged from 12 to 228 months (mean 146±62 months). During the period of study 48 patients died.

Endocrinological evaluation

Preoperatively, basal fasting level of GH, IGF-I (determined routinely since 1989), prolactin (PRL), cortisol, gonadotrophins, testosterone, estradiol and thyroid hormones were measured in the serum of all patients. Preoperatively, in 638 (95%) cases an oral glucose tolerance test (OGTT) was performed. In all other cases (n = 34) the test was not possible due to severe diabetes mellitus. Postoperatively, OGTT was performed at day 7 in all but one case (patient died at day 4 following surgery) and was repeated at 3 months in 640 of 646 (99%) and then in annual or biannual intervals in 630 (97.5%) of the cases.

IGF-I levels were routinely available from 1989. Of the 242 surgically treated patients before 1989, in 103 cases the preoperative IGF-I levels could be provided by the referring endocrinologists. For the rest of the patients no preoperative IGF-I levels were available. Postoperatively, in 158 out of 242 patients IGF-I levels were again provided by their referring doctors. For the rest, IGF-I levels were measured by the authors in a later follow-up visit and were correlated to the results of OGTT at 3 months after surgery. Due to incomplete data seven patients had to be excluded from this study.

Pituitary stimulation as a combination of adrenocorticotropic hormone (ACTH), thyrotrophin-releasing hormone (TRH) and luteinizing hormone (LH)-releasing hormone tests or the short ACTH test alone combined with basal measurements of all other pituitary hormones and with clinical signs and symptoms was used to reveal preoperative pituitary deficiencies. Hormonal deficiencies of the corticotroph and the thyrotrroph axis were corrected prior to surgical intervention. Corticosteroids were not given routinely after surgery except in cases of known hypocortisolism. The same tests as performed preoperatively were repeated 7 days after surgery. Further endocrinological and radiological controls were performed at 3 and 12 months following surgery.

Of the 630 patients, 389 (61.7%) were followed up by the authors. Long-term data of all other patients were provided by the referring endocrinologists. Follow-up examinations were held mostly at annual or biannual visits.

The assays used for hormone measurements changed over the very long time period of the study. Between December 1982 and April 1989 RIAs (CIS Isotopen-diagnostik; Dreieich, Germany) were used. From May 1989 until December 2000 ELISAs were used (Pharmacia & Upjohn, Erlangen, Germany). Since December 2000 the luminometric method was used (Immulite; DPC-Biermann, Nauheim, Germany). IGF-I assays became routinely available in 1989. Until February 2000 an RIA was used (Bio-Merieux, Marcy l’Etoile, France). Since then ELISA has been the preferred method of determination (Pharmacia & Upjohn). For the definition of ‘normality’ using different GH/IGF-I assays no allowances were made. IGF-I was always evaluated according to the age-related diagrams provided by the manufacturer of the assays.

In the present study, the data were analysed according to the most recent definition of cure resulting from the International Consensus Conferences in Cortina d’Ampezzo 1999 and Monte Carlo 2000 as published by Giustina et al. (13) (Table 1). Early remission was achieved if all three criteria were fulfilled during the endocrinological evaluation 3 months after surgery. Furthermore, the results of the annual or biannual follow-up visits were analysed to estimate the recurrence rates.

In addition to hormonal evaluation by the mentioned tests, clinical signs and symptoms of hypopituitarism were evaluated by means of a standardised questionnaire and physical examination. In females, assessment of menstrual cycle abnormalities (e.g. oligomenorrhoea or amenorrhoea) and in males, loss of facial and body hair and loss of libido and impotence, combined with low basal levels of gonadotrophins and/or a less than 3-fold increase in LH and a less than 2-fold increase in
follicle-stimulating hormone (FSH) after administration of gonadotrophin-releasing hormone led to the diagnosis of secondary hypogonadism. In the case of postmenopausal women, this was diagnosed when the serum gonadotrophins were low or low normal. Secondary adrenal failure was diagnosed in cases of antecedent malaise, circulatory collapse, unexplained vomiting, weight loss and insufficient stimulation of the serum cortisol levels after ACTH administration (e.g. increase in cortisol of less than 7 μg/dl over the basal value or a value of less than 18 μg/dl during ACTH stimulation). Secondary hypothyroidism was associated with constant feeling of coldness, coarsening, seeding of hair, thickening of the skin and low peripheral triiodothyronine and thyroxine with lowered basal thyrotrophin (TSH) and/or insufficient stimulation after TRH administration in the absence of severe non-thyroid illness or of replacement therapy with thyroid hormones for goitre prophylaxis. Postoperative GH deficiency was not assessed in these patients.

Neuropathological examination

The neuropathological evaluation was performed at the Department of Neuropathology of the University of Erlangen-Nürnberg. All specimens were fixed in 10% zinc-formalin, embedded in paraffin and stained by the haematoxylin-eosin method. Immunohistochemistry with antibodies against GH, PRL, ACTH, FSH, LH, TSH and the alpha-subunit of the glycoproteins was performed using the avidin-biotin peroxidase method. Electron microscopy was not used routinely.

Surgical treatment

The vast majority of the surgical procedures (n = 506) were performed by the transsphenoidal route. In 26 cases with invasive, asymmetrical suprasellar, retrosellar and/or suprachiasmatic parasellar extension, transcranial surgery via a pterional approach was used and only in one case (with an invasive adenoma in the interhemispheric region) was a bifrontal approach required. Combined transsphenoidal and transcranial approaches were used to treat 16 patients.

Secondary (repeat) surgery was carried out exclusively by the transsphenoidal route in 140 patients, mostly referred from other endocrinological and neurosurgical departments for persistence of acromegaly following primary surgery (n = 98). In 42 patients the first operation was performed at the authors’ department.

Statistical analysis

Data are presented as means ± s.d. Statistical analysis of the data concerning the influence of the presence of invasion, tumour size, preoperative GH levels and of the results of the neuropathological evaluation on remission rate was performed using a chi-squared test, Student’s t-test and analysis of variances (ANOVA). The level of significance was set at P < 0.05.

Results

Depending on the growth characteristics, degree and direction of extrasellar expansion, GH-secreting adenomas were classified as microadenomas (tumour diameter < 10 mm), macroadenomas (tumour diameter 11–39 mm) and giant adenomas (tumour diameter ≥ 40 mm). Macroadenomas were further divided into intrasellar tumours (is), parasellar and/or sphenoidal tumours (ps/sphe), suprasellar tumours without visual compromise (s1) and suprasellar tumours with visual compromise (s2).

Primary transsphenoidal surgery

The overall rate of endocrinological remission in the group of 506 patients undergoing primary transsphenoidal surgery was 57.3% (290 of 506) with 27% of the tumours having an invasive character. The best results were achieved in the microadenoma and intrasellar macroadenoma groups with 75.3% (107 of 142) and 74.2% (78 of 105) respectively, considering all three criteria for biochemical cure. Remission rate tended to drop in the more extensive macroadenomas: 44.5% (37 of 83) for suprasellar tumours without and 33.3% (8 of 24) for suprasellar tumours with visual compromise and 41.5% (59 of 142) for tumours with parasellar and/or sphenoidal expansion. In giant adenomas, biochemical cure was achieved only in one case (1 of 10) by surgery alone. An overview of the remission rate in the different categories following primary transsphenoidal surgery is shown in Table 2.

Table 2: An overview of the remission rate in the different categories of tumours following primary transsphenoidal surgery (n = 506); 23% of the tumours were found to be invasive.

<table>
<thead>
<tr>
<th>Macroadenomas</th>
<th>Microadenomas</th>
<th>is</th>
<th>ps/sphe</th>
<th>s1</th>
<th>s2</th>
<th>Giant adenomas</th>
<th>Σ</th>
</tr>
</thead>
<tbody>
<tr>
<td>Number of cases</td>
<td>142</td>
<td>105</td>
<td>105</td>
<td>83</td>
<td>24</td>
<td>10</td>
<td>290 (57.3%)</td>
</tr>
<tr>
<td>Remission* rate (n)</td>
<td>107 (75.3%)</td>
<td>78 (74.2%)</td>
<td>142 (98.6%)</td>
<td>59 (41.5%)</td>
<td>37 (44.6%)</td>
<td>8 (33.3%)</td>
<td>1 (10.0%)</td>
</tr>
</tbody>
</table>

* Basal GH < 2.5 ng/ml, OGT GH < 1 ng/ml, normal IGF-I.

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primary transphenoidal surgery is provided in Table 2. In a number of patients discrepancies concerning normalisation criteria were observed. Of the 152 patients with persistence of elevated IGF-I levels, in 49 cases (32.2%) suppression of GH to below 1 ng/ml during OGTT was observed. In the group of patients lacking normalisation of GH secretion during OGTT \( n = 64 \), 19 patients (29.8%) with normalised IGF-I levels were observed. Even the last group was considered as not being in remission.

**Primary transcranial surgery**

Transcranial surgery was performed in only 26 cases, mostly for extensive and invasive asymmetrical lesions extending into the suprasellar area or as a combined approach before or after transphenoidal surgery. In line with the characteristics of these lesions, the overall remission rate was only 7.7\% \( (n = 2) \). The transcranial group consisted of suprasellar macroadenomas with visual compromise \( (s, n = 16) \) and giant adenomas \( (n = 2) \). Transcranial surgery yielded remission rates of 16.2\% \( (n = 2) \) and 0\% respectively.

**Secondary (repeat) surgery**

Re-operations were performed in 140 cases in this series. In this group the overall remission rate was 27.1\% \( (n = 38) \). In the subgroup of patients \( (n = 62) \) with non-invasive tumours and an initial GH level below 40 µg/l prior to the first surgical approach, the remission rate was found to be as high as 38.7\% \( (n = 24) \).

**Surgical complications**

Mortality was rare. Only one patient (0.1\%), harbouring a giant adenoma, died on day 4 after surgery due to haemorrhage into the suprasellar tumour parts, which due to invasion of the diaphragm could not be removed by the transphenoidal approach used in this case. Morbidity was very low: meningitis and cerebrospinal fluid leaks respectively occurred in 1.8 and 0.8\% of the cases respectively. In this series, there were no cranial nerve injuries resulting in permanent neuro-ophthalmological deficits. Carotid artery injury occurred in one patient and was survived without sequelae. The complication rate in repeat transphenoidal surgery was not significantly different from that found in primary procedures.

**Impact of primary surgery on pituitary function**

For the evaluation of the surgical intervention on pituitary function only patients who underwent primary transphenoidal \( (n = 506) \) or transcranial \( (n = 26) \) surgery were taken into consideration. There was evidence of some degree of hypopituitarism preoperatively in 285 of 506 (56.3\%) of the primary transphenoidal and in 20 of 26 (76.9\%) patients of the primary transcranial group. Postoperative clinical and endocrinological workup revealed impaired pituitary function in 235 (46.4\%) after 3 months for all patients who underwent transphenoidal surgery. In contrast, the total amount of patients with endocrine deficits increased following primary transcranial surgery: hypopituitarism was encountered postoperatively in up to 24 (94\%) patients 3 months after surgery. At 3 months following primary transphenoidal surgery, anterior pituitary function was normalised in 55 (19.3\%), improved in 84 (29.4\%), remained unchanged in 141 (49.5\%) and worsened in 5 (1.8\%) of the 285 patients with some degree of preoperative hypopituitarism. Endocrine deficiencies in patient with normal preoperative pituitary function occurred in 10 of 241 (4.1\%) cases. At 3 months following transcranial surgery, anterior pituitary function was not normalised in any patient but was improved in 3 (15\%), remained unchanged in 12 (60\%) and worsened in 5 (25\%) of the 20 patients with preoperative impairment of the pituitary function. In four of the six transcranially operated patients presenting without endocrine deficits, deterioration of the pituitary function was encountered. None of the patients of the primary surgery group presented with diabetes insipidus. Six of the 506 (1.1\%) transphenoidal treated and 3 of the 26 (11.5\%) transcranially treated patients developed permanent diabetes insipidus. The results are summarised on Table 3.

Of the 532 patients undergoing primary surgery, hyperprolactinaemia was demonstrated in 182 (34.2\%) patients. In 71.9\% of these cases \( (n = 131) \) the PRL levels were found below 150 ng/ml and in 39 of them the tumour stained positive for PRL. In all but two patients with PRL levels \( > 150 \) ng/ml \( (n = 49) \) the adenoma stained positive for PRL.

<table>
<thead>
<tr>
<th>Postoperative pituitary function</th>
<th>Normal</th>
<th>Improved</th>
<th>No change</th>
<th>Worse</th>
</tr>
</thead>
<tbody>
<tr>
<td>Transphenoidal surgery ( n = 285 )</td>
<td>55 (19.3%)</td>
<td>84 (29.5%)</td>
<td>141 (49.5%)</td>
<td>5 (1.8%)</td>
</tr>
<tr>
<td>Transcranial surgery ( n = 20 )</td>
<td>0</td>
<td>3 (15.0%)</td>
<td>12 (60.0%)</td>
<td>5 (25.0%)</td>
</tr>
</tbody>
</table>

Table 3 An overview of the endocrinology of the patients with impaired preoperative pituitary function. New endocrine deficiencies in patients with normal preoperative pituitary function occurred in 10 of 241 (4.1\%) of cases.
**Remission rate in invasive tumours**

Pituitary adenomas are considered invasive if they have infiltrated or perforated the normal anatomical confines of the pituitary gland, namely the sellar diaphragm, basal dura, clivus or sphenoidal and cavernous sinuses. Although invasion tends to be more common with increasing tumour size, microadenomas may also have an invasive character and conversely large tumours may reach a considerable size by simply displacing the adjacent anatomical structures without actually invading them. The evaluation of preoperative magnetic resonance imaging (MRI) and intraoperative findings during primary transsphenoidal and transcra-nial surgery (n = 532) yielded the following results concerning the distribution of invasive tumours in this series: 7 of 142 microadenomas (5%), 124 of 378 macroadenomas (32%) and 12 of 12 giant adenomas (100%). In this series, the overall remission rate of non-invasive adenomas following primary surgery was 66.5% and dropped to 23.7% for invasive tumours. The results of each adenoma group are demonstrated separately in Table 4.

**Remission rate and tumour size and secretory activity**

Tumour size was measured from the preoperative MRI (and computed tomography) images and estimated intraoperatively by the surgeon by comparison with microinstruments. A spherical volume distribution was assumed. On the basis of the estimated diameters, tumours were classified into group A with microadenomas less than 10 mm (tumours were classified into group A with microadenomas more than 10 mm in diameter (n = 378, 70.7%) and group C with giant adenomas with more than 40 mm extension at least in one plane (n = 12, 2.2%). A remission rate of 75.3% in group A, 48.6% in group B and 8.3% in group C was observed. The results are summarised in Table 5. A significant correlation (r = 0.55, P < 0.001) was found between tumour size and remission rate in acromegaly. But also, tumour size correlated with remission rate in acromegaly. But also, tumour size correlated with the preoperative GH level particularly in large tumours with clearly elevated hormone levels. Values are means ± s.d.

**Table 4** Demonstration of the remission rates in invasive (i.) and non-invasive (n.i.) pituitary adenomas according to tumour size. In this series, the overall remission rate of non-invasive adenomas following primary surgery was 72.2% and dropped to 21.6% for invasive tumours.

<table>
<thead>
<tr>
<th>Microadenoma</th>
<th>Macroadenoma</th>
<th>Giant adenoma</th>
<th>Σ i/n.i.</th>
</tr>
</thead>
<tbody>
<tr>
<td>Number of cases</td>
<td>135/7</td>
<td>254/124</td>
<td>389/143</td>
</tr>
<tr>
<td>Remission* rate</td>
<td>100/6 (74%/85%)</td>
<td>181/24 (71.2%/19.3%)</td>
<td>281/31 (72.2%/21.6%)</td>
</tr>
</tbody>
</table>

*Basal GH < 2.5 ng/ml, OGT GH < 1 ng/ml, normal IGF-I.

**Table 5** A significant correlation was found between tumour size and remission rate in acromegaly. But also, tumour size correlated with the preoperative GH level particularly in large tumours with clearly elevated hormone levels. Values are means ± s.d.

<table>
<thead>
<tr>
<th>Microadenoma</th>
<th>Macroadenoma</th>
<th>Giant adenoma</th>
</tr>
</thead>
<tbody>
<tr>
<td>Diameter (mm)</td>
<td>142 (27%)</td>
<td>378 (70.7%)</td>
</tr>
<tr>
<td>Average diameter</td>
<td>2–9</td>
<td>10–37</td>
</tr>
<tr>
<td>GH level (μg/l)</td>
<td>7.6±1.75</td>
<td>16.7±5.6++</td>
</tr>
<tr>
<td>Average GH</td>
<td>1–142</td>
<td>4–357</td>
</tr>
<tr>
<td>level (μg/l)</td>
<td>16.4±15.2</td>
<td>42.6±38.4++</td>
</tr>
<tr>
<td>Remission rate</td>
<td>107 (75.3%)</td>
<td>186 (48.6%)</td>
</tr>
</tbody>
</table>

10 μg/l the remission rate was as high as 89.5%. If a basal GH level of 50 μg/l is defined as an arbitrary threshold for the secretory activity of an adenoma, then the overall cure rate drops from 74% in cases with preoperative GH levels below this threshold to 25% in cases with preoperative GH levels above this threshold. No patient with GH level above 125 μg/l could be surgically cured.

**Remission rate and neuropathological evaluation**

Since the electron microscopy was not used routinely, it was not possible to evaluate the influence of the ultrastructural characteristics of the lesions on the endocrinological outcome. Therefore only their immunohistochemical profiles in terms of positive staining were taken into consideration. Of the 532 patients who underwent primary transsphenoidal or transcra-nial surgery, tumour specimens could be obtained in 526 cases. In six cases and due to very small tumours no material could be preserved. Three hundred and forty-seven (65.9%) tumours stained positive only for GH, In 159 (30.2%) tumours a dual secretion of GH and PRL was noted. Fourteen (2.6%) tumours stained positive for GH and TSH and six (1.1%) for GH, PRL and TSH. The remission rates were found to differ significantly between the pure GH-secreting and those tumours secreting more than one hormone and was found to be 73.7% (n = 256), 31.4% (n = 50), 0% and 0% for the four groups respectively.
Recurrence rate

In the present series, a mean follow-up period of more than 120 months (mean 126 ± 74 months) has been observed. During this period and using all three criteria for the definition of biochemical cure, only two patients (0.4%) with recurrent acromegaly presented 3 and 6 years after initial surgery.

Discussion

The goals of surgical therapy in acromegaly include initial endocrinological remission with normalisation of the dynamic GH secretion pattern as well as normalisation of basal IGF-I to give regression of the signs and symptoms of the disease and prevention of recurrence (long-term remission), performance of selective adenomectomy (preservation of normal pituitary functions) and elimination of mass effects resulting in restoration of normal neurological function.

As a result of the difficulties in defining postoperative endocrinological remission and the variations in criteria used by different investigators (13), the success rates in terms of achievement of these goals vary significantly in the published series from different institutions (Table 6).

In early series, the criterion for remission was a postoperative GH level below 5 μg/l. Ross & Wilson (14) in 1998 analysed the results of 30 published series and yielded an overall ‘cure’ rate of 56% in 153 cases. In another multicentre study published in 1987 by Zervas (15) the overall ‘cure’ rate was 66% in 1256 cases. Similar results were found in other series published by Losa et al. 1989 (16) and Valdemarsson et al. in 1991 and 2000 (17, 18).

Table 6 Results of primary transsphenoidal surgery for GH-secreting pituitary adenomas.

<table>
<thead>
<tr>
<th>Series</th>
<th>No. cases</th>
<th>Total cure rate (%)</th>
<th>Microadenomas</th>
<th>Macroadenomas</th>
<th>Definition of ‘cure’</th>
</tr>
</thead>
<tbody>
<tr>
<td>Losa et al. 1989 (16)</td>
<td>29</td>
<td>55</td>
<td>n.d.</td>
<td>n.d.</td>
<td>GH &lt; 1 μg/l and normal IGF-I level</td>
</tr>
<tr>
<td>Fahlbusch et al. 1992 (19)</td>
<td>222</td>
<td>57</td>
<td>72</td>
<td>65</td>
<td>GH &lt; 2 μg/l OGT</td>
</tr>
<tr>
<td>Tindall et al. 1993 (20)</td>
<td>91</td>
<td>82</td>
<td>n.d.</td>
<td>n.d.</td>
<td>GH &lt; 5 μg/l and/or normal IGF-I level</td>
</tr>
<tr>
<td>Davis et al. 1993 (36)</td>
<td>174</td>
<td>52</td>
<td>n.d.</td>
<td>n.d.</td>
<td>GH ≤ 2 μg/l (basal or OGT)</td>
</tr>
<tr>
<td>Sheaves et al. 1996 (28)</td>
<td>100</td>
<td>42</td>
<td>61</td>
<td>23</td>
<td>GH &lt; 2.5 μg/l</td>
</tr>
<tr>
<td>Aboch et al. 1998 (41)</td>
<td>254</td>
<td>76</td>
<td>75</td>
<td>71</td>
<td>GH &lt; 5 μg/l</td>
</tr>
<tr>
<td>Freda et al. 1998 (23)</td>
<td>115</td>
<td>61</td>
<td>88</td>
<td>53</td>
<td>GH &lt; 2.5 μg/l OGT or normal IGF-I level</td>
</tr>
<tr>
<td>Swearingen et al. 1998 (4)</td>
<td>162</td>
<td>57</td>
<td>91</td>
<td>48</td>
<td>Random GH &lt; 2.5 μg/l</td>
</tr>
<tr>
<td>Gittoes et al. 1999 (42)</td>
<td>160</td>
<td>64</td>
<td>86</td>
<td>52</td>
<td>GH &lt; 2.5 μg/l or GH &lt; 1 μg/l (OFT)</td>
</tr>
<tr>
<td>Ahmed et al. 1999 (39)</td>
<td>139</td>
<td>67</td>
<td>91</td>
<td>45</td>
<td>Mean GH &lt; 2.5 μgl</td>
</tr>
<tr>
<td>Laws et al. 2000 (21)</td>
<td>117</td>
<td>67</td>
<td>87</td>
<td>50.5</td>
<td>GH &lt; 2.5 μg/l, GH ≤ 1 μg/l, normal IGF-I level</td>
</tr>
<tr>
<td>Kreutzer et al. 2001 (22)</td>
<td>57</td>
<td>70</td>
<td>n.d.</td>
<td>n.d.</td>
<td>Normal IGF-I level</td>
</tr>
<tr>
<td>Beauregard et al. 2003 (1)</td>
<td>103</td>
<td>52</td>
<td>82</td>
<td>60</td>
<td>Basal GH ≤ 2.5 μg/l (OFT), normal IGF-I level</td>
</tr>
<tr>
<td>This series</td>
<td>506</td>
<td>57</td>
<td>75</td>
<td>50</td>
<td>Basal GH ≤ 2.5 μg/l (OFT), normal IGF-I level</td>
</tr>
</tbody>
</table>

n.d., not done.
no definite ‘cure’ of the disease is achieved. Even following ‘successful’ surgical treatment, GH levels remain detectable and pulses of GH secretion are not normal in frequency and size (25). In acromegaly the most accurate term to be used is ‘safe’ GH secretion patterns, meaning relief of symptoms, restoration of normal life expectancy and minimal risk of recurrence of the disease. IGF-I seems to play a very important role in predicting long-term outcome. But more studies are still needed in order to precisely evaluate the value of IGF-I levels as a long-term outcome criterion. Together with the development of more sensitive ELISAs in the near future we may then be able to better define ‘cure’ in acromegalic patients.

The evaluation of the present series led to detection of several predictors of the surgical result, such as tumour size, invasiveness, extrasellar growth, secretory activity and the neuropathological findings. Regarding the growth characteristics of the lesions, microadenomas have a more favourable outcome than macroadenomas, particularly those with extrasellar extension. In microadenomas, a remission rate of as high as 75.3% could be achieved. With increasing tumour diameter and extrasellar extension, remission drops to 0% for giant adenomas. Similar results in 30 published surgical series and estimated an operative mortality rate of 1.04% (14). Due to specific precautions taken in the high-risk group of acromegalic patients, the complication rate in the presented series was comparable to that of non-acromegalic patients undergoing transsphenoidal surgery for other types of pituitary adenomas. Preoperative evaluation included assessment of cardiac function, blood pressure, respiratory function and metabolic disturbances. Morphological abnormalities, such as macroglossia and the enlargement of the jaw were taken into consideration. In several cases, fibre-optic endotracheal intubation was necessary and many patients with severe sleep apnoea were postoperatively monitored in the intensive care unit. All these problems must be carefully managed throughout the perioperative period in order to keep the complication rate as low as possible. In cases of markedly reduced general condition, short-term medical treatment with octreotide should be considered, which has proven very helpful in bringing the patient to a better condition before surgery (29, 30).

Postoperative pituitary deficiencies, particularly panhypopituitarism, are considered dreaded complications of pituitary surgery because they dramatically affect the patient’s quality of life by necessitating hormonal substitution therapy and lead to increased mortality. Following the introduction of transsphenoidal microsurgery in the management of patients with pituitary tumours, selective adenomectomy was aimed at, implying preservation of as much normal pituitary tissue as possible. Several studies demonstrated that preservation of pituitary tissue was associated with no further loss of pituitary function in the majority of patients. Only recently, several groups reported that impaired pituitary function could actually be restored by selective surgical resection of the adenoma (31–33). This observation could be confirmed in this study in which a large patient material was homogeneously treated by the same surgical techniques and evaluation in one laboratory. Residual anterior pituitary function was more frequently preserved after transsphenoidal surgery (49.5%) and could even be normalised or improved in 48.7% of cases with preoperative endocrine deficiencies. In contrast to most reports in the literature and despite surgical manipulations necessary to remove all tumour parts, only a moderate increase in anterior pituitary deficiency (4.1%) was observed as an untoward effect of the operation.

The frequency of tumour recurrence is strongly associated with the remission criteria used. It is paramount to distinguish between postoperative normalisation of GH secretion using all three criteria and those cases with lower postoperative basal GH levels but inadequate response to OGTT and/or abnormal IGF-I levels. Using a postoperative basal GH below 5 μg/l as the remission criterion, the overall ‘recurrence’ rate in 61 patients was 7% over a mean follow-up period...
of 6 years (19). Similar results were published by Losa et al. (16) and Arafah et al. (34). However, using suppression of GH levels below 2.0 µg/l as the criterion, long-term remission can be achieved as observed by our group in 1991 (35). In other published series, rates of recurrences up to 17.8% are encountered using other criteria for biochemical cure (36). In the present series, a mean follow-up period of more than 10 years has been observed. During this period and using all three criteria for biochemical cure, only two patients (0.4%) with recurrent acromegaly presented 3–6 years after initial surgery. This fact stresses the paramount importance of stringent endocrinological remission criteria in acromegaly.

The management of patients with persistent acromegaly after the first operation is a matter of ongoing discussion. Several meetings have been held to obtain a consensus about how these patients should ideally be treated. All treatment modalities concerning this disease have been discussed in detail. Many of these patients are candidates for repeat surgery. Re-exploration was found to be particularly successful in those cases in which neither surgical nor histological invasion could be demonstrated during the first surgery. In this series, re-explorations have been performed in a number of selected cases, mainly in patients with MRI studies that demonstrated residual adenomatous tissue. As expected, the overall remission rate was worse than in the primary surgery group, at 27.1%. Nevertheless, a remission rate of 38.7% could be achieved in patients with non-invasive tumours and an initial GH level below 40 µg/l prior to the first surgical approach. To evaluate the effectiveness of secondary transnasal microsurgery, 28 patients with persistent or recurrent acromegaly were analysed by Abe & Ludecke in 1998 (37). The overall endocrinological remission rate was found to be 57.1% (16 of 28 patients) without serious morbidity and no mortality. Kurosaki et al. (38) reported even higher remission rates: overall remission 60%, remission in transnasally respectable tumours 81.3%. The absence of major surgical complications and significant endocrinological deterioration of other pituitary functions combined with the possibility of long-term control of the disease supports re-exploration as a treatment option in a number of selected patients with persistent acromegaly.

The results in this series, once more strongly support the role of surgery as the primary management option of acromegalic patients. Surgery is very efficacious in restoring normal GH secretion patterns, acts rapidly and carries a low therapy-associated morbidity. It is also clear that surgical experience, skill and knowledge play an important role in how successful the primary surgery is (39, 40). The results presented above concern only those series published by authors with major expertise in this field and cannot be achieved by surgeons with little experience in pituitary surgery. Supraregional specialisation in a few neurosurgical centres with equally experienced endocrinologists and laboratory staff may lead to even better surgical results in acromegaly.

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References


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