Ki-67 labelling index and invasiveness among anterior pituitary adenomas: analysis of 103 cases using the MIB-1 monoclonal antibody

Luciano Mastronardi, Antonio Guiducci, Cristina Spera, Fabrizio Puzzilli, Fabio Liberati, Giulio Maira

Abstract

Aims—To investigate the relation between proliferative activity of anterior pituitary adenomas, quantified by the Ki-67 labelling index, and their invasive behaviour.

Methods—Expression of Ki-67 was evaluated in 103 anterior pituitary adenomas consecutively operated on in a 36 month period and correlated with surgical evidence of invasiveness.

Results—Non-invasive (n = 65) and invasive (n = 38) adenomas were identified from surgically verified infiltration of sellar floor dura and bone. The wall of the cavernous sinus was infiltrated in 16 cases. Forty one adenomas were non-functioning and 62 functioning (24 prolactin, 21 growth hormone, 10 ACTH, seven mixed). The overall mean (SD) Ki-67 labelling index was 2.64 (3.69) per cent (median 1.5). The mean index was 3.08 (4.59) per cent in functioning and 1.97 (1.78) per cent in non-functioning tumours; 5.47 (9.52) per cent in ACTH adenomas and 2.33 (2.42) per cent in others (p = 0.01); 3.71 (5.17) per cent in invasive and 2.01 (2.45) per cent in non-invasive adenomas (p = 0.027); and 5.58 (7.24) per cent in cavernous sinus infiltrating and 2.10 (2.39) per cent in cavernous sinus non-infiltrating adenomas (p = 0.0008). To identify a value of labelling index beyond which adenomas should be considered invasive and another beyond which cavernous sinus infiltration should be suspected, normality Q–Q plots were obtained: a threshold labelling index of 5.58 (7.24) per cent in cavernous sinus infiltrating adenomas was obtained; a threshold labelling index of 3.71 (5.17) per cent in invasive and 2.01 (2.45) per cent in non-invasive adenomas was defined, with statistically significant differences (p = 0.02 and p = 0.004, respectively).

Conclusions—The Ki-67 labelling index can be considered a useful marker in determining the invasive behaviour of anterior pituitary adenomas.

(J Clin Pathol 1999;52:107–111)

Keywords: invasiveness; Ki-67; pituitary adenomas

Patients who can be considered surgically cured from patients at high risk of clinical and radiological recurrence, several experimental and clinicopathological studies have been undertaken to identify markers of tumour invasiveness.1–14

As with other neoplasms, determination of cell proliferation activity seems to be prognostically useful in anterior pituitary adenomas. Ki-67, in particular, is a nuclear antigen expressed in the G1, S, G2, and M phases of the cell cycle,15–21 and its labelling index, now characterised by the monoclonal antibody MIB-1,22 is widely considered to be a marker of cellular proliferation (growth fraction).

Our aim in this study was to investigate the relation between the proliferative activity of anterior pituitary adenomas, quantified by the Ki-67 labelling index, and their invasive behaviour.

Methods

PATIENT CHARACTERISTICS

Between July 1994 and July 1997, 103 patients suffering from an anterior pituitary adenoma were operated on in our neurosurgical division. In all cases an evaluation of the Ki-67 labelling index of the surgical specimens was performed using the monoclonal antibody MIB-1.22

The patients’ age ranged from 17 to 77 years (mean (SD) 42.8 (15.2) years, median 39): 10 cases (9.8 per cent) were under 25 years, 57 (55.3 per cent) were between 26 and 50 years, and 36 (34.9 per cent) were over 50 years. The M/F ratio was 1:1.39 (60 females and 43 males). In 18 cases a previous operation had been performed elsewhere for pituitary adenoma, and in two cases two previous operations. A functioning adenoma, with clinically expressed hormonal dysfunction, was present in 62 patients (59 per cent): in 27 cases (43.3 per cent) the disturbances were related to high serum prolactin, in 25 (40.3 per cent) there was acromegaly, and in 10 (16.1 per cent) Cushing’s syndrome. The preoperative serum hormone concentration was defined as the highest value before surgery. Reduced visual acuity or campimetric disturbances were present in 47 cases (45.6 per cent), mainly in patients with non-functioning adenomas (55.8 v 38.7 per cent). Raised intracranial pressure was observed in six cases (5.9 per cent).

In all cases the neuroradiological diagnosis was obtained by contrast enhanced cerebral computed tomography and magnetic resonance imaging (MRI); in 20 cases (19.4 per

Department of Neurological Sciences, Unit of Neurosurgery, Civilian Hospital, Terni, Italy

L Mastronardi
C Spera
F Puzzilli
G Maia

Institute of Pathological Anatomy, Civilian Hospital, Terni

A Guiducci
F Liberati

Correspondence to: Dr Luciano Mastronardi, Via Archimede 120, 00197 Rome, Italy. email: mastro@tin.it

Accepted for publication 21 September 1998
The surgical specimens were routinely processed, fixed in neutral buffered formalin, and embedded in paraffin. To evaluate the Ki-67 antigen staining, 5 µm sections, previously mounted onto glass slides and dried, were incubated overnight at 4°C in the MIB-1 antibody (Immunotech). Immunostaining was performed using the avidin–biotin–peroxidase method. Ten fields were selected in regions in which there were the highest concentrations of MIB-1 positive nuclei and were examined at high power (×400); each field corresponded to 700 to 1000 cells, depending on the cellularity of the tumour specimen. Areas of necrosis, normal adenohypophysal cells, and endothelial cells were excluded from the evaluation. On the basis of a manual count of 1000 cells, the Ki-67 labelling index was defined as the proportion of MIB-1 positive cells (dense brown precipitate restricted to the nuclei). The histopathological diagnosis and the Ki-67 labelling index evaluation was performed by a single neuropathologist (AG).

STATISTICAL ANALYSIS

Computer assisted data analysis was performed with commercially available software (SPSS 6.0). The $\chi^2$ test (1 or 2 degrees of freedom, continuity correction) and analysis of variance (ANOVA) were used to identify the statistical significance of differences of Ki-67 labelling index observed in relation to age, sex, previously operated $v$ newly diagnosed patients, functioning $v$ non-functioning adenomas, type of hormone secreted, presence or absence of preoperative visual disturbances, neuroradiologically estimated volume of tumour, surgical invasiveness, and infiltration of cavernous sinus. Values are expressed as the mean (SD); for each comparison, a $p$ value was obtained and significance was assumed at $p \leq 0.05$. To identify a value of MIB-1 beyond which adenomas are likely to invade the sella floor dura and bone, and another beyond which cavernous sinus infiltration is likely, the expected normal values obtained from normality Q-Q plots (figs 1 and 2) were calculated as thresholds.

Results

A patient with a huge, invasive, cavernous sinus infiltrating adenoma died in the postoperative period from pulmonary embolism. Morbidity consisted of a cerebrospinal fluid fistula in five patients (4.8 per cent of 105 surgical procedures), treated successfully with spinal cerebrospinal fluid drainage through a lumbar subarachnoid catheter for three to four days.

In 18 cases the adenoma was functioning and invasive, in 44 functioning and non-invasive, in 20 non-functioning and invasive, and in 21 non-functioning and non-invasive ($p = 0.022$). Table 1 summarises the mean Ki-67 labelling indices of 103 anterior pituitary adenomas.

The overall mean (SD) Ki-67 labelling index was 2.64 (3.69) per cent (median 1.5; range 0 to 31). In five cases the labelling index was $\geq 10$ per cent. All these were secreting tumours (two ACTH, one growth hormone, one prolactin, and one growth hormone–prolactin), three were invasive, and two had infiltrated the cavernous sinus. In none of these cases was there any histological evidence of pituitary carcinoma.

There were no differences in the index in relation to age, sex, previous operations, presence of preoperative visual disturbances, or

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**Figure 1** Normality plot of MIB-1 values in relation to invasiveness of sellar floor dura and bone.

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**Figure 2** Normality plot of MIB-1 values in relation to infiltration of the cavernous sinus.
neuroradiological estimation of the volume of the tumour. The Ki-67 labelling index was 3.08 (4.59) per cent in functioning tumours and 1.97 (1.78) per cent in the non-functioning tumours (NS). ACTH secreting adenomas had a higher labelling index than the others, at 5.47 (9.52) v 2.33 (2.42) per cent; p = 0.01. We found no significant differences among the other types of hormone releasing adenomas.

Preoperative treatment with bromocriptine had no effect on the labelling index in prolactin secreting adenomas or mixed adenomas with predominant prolactin expression.

Invasive adenomas had a Ki-67 labelling index of 3.71 (5.17) per cent, v 2.01 (2.45) per cent in non-invasive tumours (p = 0.027). The index was 5.58 (7.24) per cent in cavernous sinus infiltrating and 2.10 (2.39) per cent in non-infiltrating adenomas (p = 0.0005).

We derived normality Q-Q plots with the observed values of MIB-1 (figs 1 and 2). For invasive adenomas a threshold value of 3.5 per cent was identified and for cavernous sinus infiltrating adenomas, 5 per cent. The results obtained are summarised in tables 2 and 3. The observed differences were statistically significant (p = 0.02 and p = 0.004, respectively).

After follow up ranging from six to 42 months, a reoperation for the treatment of a clinical and radiological tumour recurrence was performed in two patients after eight and 14 months, respectively. In these patients the first intervention was performed by a trans-sphenoidal approach and the second by craniotomy. At the first operation, the tumour was found to be invasive in both cases, and in one it invaded the cavernous sinus. In these cases the Ki-67 labelling indices of the first specimens were 6.1 per cent and 3.8 per cent, respectively. Both the relapsed tumours had invaded sellar floor dura and bone and the wall of the cavernous sinus; the Ki-67 labelling indices of the recurrent tumours were 10 per cent and 5.5 per cent, respectively.

### Table 1 Mean Ki-67 labelling indices (LI) of 103 anterior pituitary adenomas

<table>
<thead>
<tr>
<th>Cases</th>
<th>n</th>
<th>Mean LI</th>
<th>SD</th>
<th>p Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>All patients</td>
<td>103</td>
<td>2.64</td>
<td>3.69</td>
<td></td>
</tr>
<tr>
<td>&lt;25 years</td>
<td>10</td>
<td>2.67</td>
<td>2.01</td>
<td></td>
</tr>
<tr>
<td>25–50</td>
<td>57</td>
<td>2.30</td>
<td>2.27</td>
<td></td>
</tr>
<tr>
<td>&gt;50</td>
<td>36</td>
<td>3.15</td>
<td>5.60</td>
<td>0.57</td>
</tr>
<tr>
<td>Previous operations</td>
<td>Yes</td>
<td>2.48</td>
<td>2.30</td>
<td></td>
</tr>
<tr>
<td>No</td>
<td>83</td>
<td>2.08</td>
<td>4.05</td>
<td>0.83</td>
</tr>
<tr>
<td>Visual disturbances</td>
<td>Yes</td>
<td>2.65</td>
<td>2.49</td>
<td></td>
</tr>
<tr>
<td>No</td>
<td>56</td>
<td>2.02</td>
<td>4.58</td>
<td>0.97</td>
</tr>
<tr>
<td>Functioning adenoma</td>
<td>Yes</td>
<td>3.08</td>
<td>4.59</td>
<td></td>
</tr>
<tr>
<td>No</td>
<td>41</td>
<td>1.97</td>
<td>1.78</td>
<td>0.14</td>
</tr>
</tbody>
</table>

- ACTH, adrenocorticotrophic hormone; PRL, prolactin; TSH, thyroid stimulating hormone.

### Table 2 Comparison of invasive behaviour among 103 anterior pituitary adenomas with measurement of Ki-67 labelling index (LI, MIB-1), using a threshold value of 3.5% (see text)

<table>
<thead>
<tr>
<th>LI ≤3.5%</th>
<th>Invasive</th>
<th>Non-invasive</th>
</tr>
</thead>
<tbody>
<tr>
<td>24</td>
<td>54</td>
<td></td>
</tr>
<tr>
<td>LI &gt;3.5%</td>
<td>14</td>
<td>11</td>
</tr>
<tr>
<td>Total</td>
<td>38</td>
<td>65</td>
</tr>
<tr>
<td>p Value</td>
<td>0.02</td>
<td></td>
</tr>
</tbody>
</table>

### Table 3 Comparison of cavernous sinus infiltration among 103 anterior pituitary adenomas with measurement of Ki-67 labelling index (LI, MIB-1), using a threshold value of 5% (see text)

<table>
<thead>
<tr>
<th>Cavernous sinus infiltration</th>
<th>Invasive</th>
<th>Non-invasive</th>
</tr>
</thead>
<tbody>
<tr>
<td>LI ≤5%</td>
<td>10</td>
<td>80</td>
</tr>
<tr>
<td>LI &gt;5%</td>
<td>6</td>
<td>7</td>
</tr>
<tr>
<td>Total</td>
<td>16</td>
<td>87</td>
</tr>
<tr>
<td>p Value</td>
<td>0.004</td>
<td></td>
</tr>
</tbody>
</table>

ACTH, adrenocorticotrophic hormone; FSH, follicle stimulating hormone; GH, growth hormone; PRL, prolactin; TSH, thyroid stimulating hormone.

Discussion

A tendency for juxtasellar expansion of some anterior pituitary adenomas has been observed from the first reports. In the early 1970s, Jefferson identified a group of pituitary tumours that he defined “invasive” adenomas, in which an extrasellar spread occurred, sometimes with infiltration of the cavernous sinus.Martins et al similarly defined an invasive pituitary adenoma as a tumour which extends beyond its capsule or involves contiguous structures. The incidence of invasiveness among these tumours varies among different anterior pituitary adenoma subtypes and also in relation to the criteria used for assessment. Infiltrating behaviour has been demonstrated surgically in about 35 per cent of pituitary adenomas and histologically (by microscopic infiltration of dura mater) in about 90 per cent. From our experience and in agreement with Thapar et al, the most reliable criterion of invasiveness seems to be the surgeon’s impression during the operation.

Until now, unfortunately, no routine markers have been available to identify invasive anterior pituitary adenomas. The usual morphological findings typical of histological aggressiveness are quite uncommon among these neoplasms. Recently, however, several cell cycle specific nuclear antigens have been recognised, using various immunohistochimical methods, which allow reliable evaluation of tumour growth characteristics. One in particular, Ki-67, is a nuclear antigen readily identified by the monoclonal antibody MIB-1, and is typically expressed in proliferating cells during the G1, S, G2, and M phases of the cell cycle. It has been found to be useful in assessing several human neoplasms, providing information about the cell proliferation rate and thus about long term prognosis.
Table 4: Proposal for a grading system for invasiveness of pituitary adenomas in relation to mean Ki-67 labelling indices (LI) observed in 103 cases, and in one case with pituitary carcinoma (not included in this report).

<table>
<thead>
<tr>
<th>Grade</th>
<th>Mean Li-67 LI</th>
</tr>
</thead>
<tbody>
<tr>
<td>0</td>
<td>Non-invasive pituitary adenomas</td>
</tr>
<tr>
<td>1</td>
<td>Invasive pituitary adenomas</td>
</tr>
<tr>
<td>2</td>
<td>Caverno-sus sinus infiltrating pituitary adenomas</td>
</tr>
<tr>
<td>3</td>
<td>Pituitary carcinoma (1 case)</td>
</tr>
</tbody>
</table>

MIB-1 immunostaining is a practical method that can be used in the routine histological evaluation of brain tumours and also of anterior pituitary adenomas. Though automated analysis is faster and easier, manual cell counting is equally reliable and is applicable everywhere. A high growth fraction expressed by a high Ki-67 labelling index should suggest the need for careful clinical and radiological follow up.

To date, only a few investigators have reported their experience in evaluating anterior pituitary adenoma growth fraction based on the expression of the Ki-67 antigen, using both formalin fixed tissue and in vitro cell culture. Landolt et al examined the growth rate of 31 pituitary adenomas, the Ki-67 labelling index ranged from 0.1 to 3.7 per cent, with higher values in functioning adenomas (especially the ACTH secreting type) and in invasive adenomas verified radiologically, surgically, and histologically. In a series of 62 cases, Knosp et al reported proliferation activity ranging from 0.1 to 2.8 per cent, with a higher Ki-67 labelling index (p < 0.05) in adenomas with histological evidence of dural infiltration. Daita et al evaluated the correlation between histological invasiveness of the sellar floor dura and proliferative activity in 31 pituitary adenomas, showing a higher MIB-1 positive ratio with dural invasion than with non-invasive tumours (p < 0.05). Asano et al evaluated the MIB-1 index in 63 surgically removed anterior pituitary adenomas; values ranged from 0 to 6.5 per cent, with low values in growth hormone secreting tumours and high values in prolactin secreting tumours. Preoperative treatment with bromocriptine had no effect on the labelling index values. In our series we also found that growth hormone secreting adenomas had a low Ki-67 labelling index, whereas prolactin secreting tumours had a mean index above the mean for the cohort as a whole (table 1). Shibuya et al reported their experience of 65 pituitary adenomas in which, among other tests, the Ki-67 expression was evaluated; they obtained higher labelling index values in recurrent, non-functioning, and ACTH secreting adenomas. We confirmed the latter finding in our series (table 1, p = 0.01), as have other investigators, and the finding of a high mean Ki-67 labelling index in recurrent adenomas was also shown by Ekramullah et al. These latter workers evaluated MIB-1 positivity in 14 regrowing non-functioning pituitary adenomas and 19 cases without clinical or MRI signs of tumour recurrence; they observed a higher index (p< 0.01) in the group with recurrence (mean 0.86 per cent) than in the “cured” cases (mean 0.23 per cent). Though in our series we did not find any difference in Ki-67 labelling index between the 20 patients previously operated on elsewhere and the 83 newly diagnosed patients (table 1), the indices in the two patients who presented with recurrences were 6.1 per cent and 3.8 per cent, respectively, while in relapsed tumours invading the surrounding tissues and the cavernous sinus the indices were higher (10 per cent and 5.5 per cent, respectively). We cannot rule out the possibility that some of the patients in our series who underwent reoperation did not have a true recurrence of an aggressive tumour but regrowth of the residual portion of the tumour, which was not completely extirpated at the original operation. This would invalidate the low mean Ki-67 labelling index observed in the reoperated group (table 1).

In a series of 70 anterior pituitary adenomas and seven carcinomas, Thapar et al reported mean MIB-1 values of 1.37 per cent in non-invasive adenomas, 4.66 per cent in invasive adenomas, and 11.91 per cent in pituitary carcinomas (p < 0.01). As in our findings (table 1), functioning adenomas in their series had a higher (p = 0.03) mean labelling index (3.25 per cent) than non-functioning adenomas (2.06 per cent). These investigators established a value of 3 per cent as the threshold labelling index for distinguishing non-invasive from invasive anterior pituitary adenomas. In our study, using normal Q-Q plots of observed MIB-1 values (figs 1 and 2), we identified two different threshold values of the Ki-67 labelling index, one for invasive adenomas (3.5 per cent) and one for cavernous sinus infiltrating adenomas (5 per cent). The results obtained with these thresholds are summarised in tables 2 and 3, and the differences observed were statistically significant (p = 0.02 and p = 0.004, respectively). From the mean Ki-67 labelling indices, it appears that adenomas with surgically verified infiltration of the sellar floor dura and bone are less aggressive than those that also infiltrate the wall of cavernous sinus; the latter probably represents a second level of invasiveness (table 4), though the biological significance of this difference remains unclear. Thus an index ranging between 3.5 and 5 per cent could suggest infiltrating behaviour, while an index above 5 per cent also suggests the possibility of infiltration of the cavernous sinus wall. In both circumstances, careful clinical, hormonal, and MRI follow up seems necessary so as to identify the predicted recurrence of the tumour as early as possible.

CONCLUSION
It is well recognised that about one third of pituitary adenomas are “invasive,” with extension beyond the capsule and involvement of contiguous structures, sometimes with infiltration of the cavernous sinus. Evaluation of the growth fraction—identified by Ki-67 nuclear antigen, which is easily detected with MIB-1 monoclonal antibody—seems to provide additional insight into the infiltrative
Ki-67 labelling index and invasiveness in pituitary adenomas

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