Correlations between histological type, clinical behaviour, and prognosis in thymoma

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ABSTRACT Seventy four cases of thymoma were reclassified into three histological categories—cortical (30), medullary (9), and mixed (34) (the remaining patient had an intrathymic thymoma)—for an investigation of the relation between histological type, clinical behaviour, and long term prognosis. There were significant differences between the histological types in the frequency of myasthenia gravis and of the different tumour stages, the mean age of the patients, and prognosis. Myasthenia gravis occurred more commonly in patients with cortical (33%) and mixed thymoma (35%) than in patients with medullary thymoma (11%). Five, 10, 15, and 20 year actuarial survival was 100% for medullary thymoma; 85%, 76%, 65% and 65% respectively for mixed thymoma; and 52%, 45%, 45%, and 45% for cortical thymoma. Medullary thymoma is a benign tumour arising late in life and there was no mortality in this series after surgery alone. Cortical thymoma usually presented in middle age and must be regarded as malignant; mortality was 50% at five years despite a multidisciplinary approach, with surgery and postoperative radiotherapy in all patients and chemotherapy in selected cases. Mixed thymoma had a better prognosis than cortical thymoma, but must be regarded as potentially malignant. One third of the total patients had died by 10 years despite radical tumour resection.

Introduction

Thymomas, defined as thymic epithelial neoplasms with minimal or no cytological atypia,1 are characterised by a remarkable morphological heterogeneity2 and variable clinical behaviour.14 Histological classifications in current use are based purely on morphology245 and have little prognostic or therapeutic value. In consequence, treatment is at present based solely on the surgical stage.

The classification recently proposed by Marino and Müller-Hermelink6 takes both morphology and histogenesis into account, and provides a new approach to management. In a preliminary investigation7 we found that the classification correlated well with the surgical stage of the tumour. In the present study we have analysed the relation between clinical features, long term prognosis, and histological type in a larger group of patients.

Methods

From January 1965 to December 1987 120 patients with thymoma were treated surgically in the department of surgery of the University of Rome. In 38 cases insufficient pathological material was available for review, and eight patients were lost to follow up. The remaining 74 cases form the basis of this investigation.

Sections 4 μm thick were stained for conventional histological examination with haematoxylin and eosin and the Giemsa, periodic acid-Schiff, and Gomori techniques, and examined by the same pathologists (CDB, EOP, LPR), all of whom have specific experience in the subject. Invasiveness was graded according to the criteria of Masaoka.8 Symptoms and signs, paraneoplastic syndromes, completeness of resection, and follow up were related to histological type. Survival was calculated for 73 patients by the actuarial method9—one patient with intrathymic thymoma was excluded. Student's t test and χ² test were used for statistical analysis.

HISTOLOGICAL CLASSIFICATION

In this study a classification based on that of Marino...
and Müller-Hermelink\textsuperscript{6} was used, but categorising thymomas into only three types—cortical, mixed, and medullary.

By this classification cortical thymomas are defined as tumours composed mainly of medium sized to large epithelial cells, with round or oval nuclei, finely dispersed chromatin, prominent central nucleoli, and an ill defined cytoplasm (fig 1). Lymphocytes are usually abundant, often with a blastic appearance. There is frequently a “starry sky” pattern, focal medullary differentiation with Hassal’s corpuscles, and perivascular epithelial cell palisading.

Medullary thymomas are composed of small to medium sized cells with irregular, often spindle shaped nuclei devoid of nucleoli (fig 2). Lymphocytes, generally present in small numbers, are of the mature

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**Fig 1**  
Cortical thymoma: epithelial cells of the cortical type with prominent nucleoli (arrows) and scattered lymphocytes.

**Fig 2**  
Medullary thymoma: spindle shaped epithelial cells of medullary type with a scanty lymphoid component.
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Fig 3 Mixed thymoma: (a) cortical (arrows) and medullary (arrowheads) type epithelial cells with a few lymphocytes; (b) dilated perivascular spaces, a typical feature of mixed thymoma.

Thymocytic type. Epithelial lined cysts are often present, and a storiform pattern is sometimes observed.

In mixed thymomas epithelial components of cortical and medullary type are both present, intermingled with a variable number of lymphocytes (fig 3a). Dilated and sometimes hyaline perivascular spaces are frequent (fig 3b). Mixed thymomas with a clear predominance of cortical or medullary differentiation were assigned to the cortical and the medullary group respectively.

Intrathymic thymoma, composed of abnormal
myasthenia
gravis, CT (30)
<
<
p


differs (34)

458

4

Intrathymic
Histological
MT (9)

Distribution differs significantly
thymoma.

microscopic islands of epithelial cells surrounded by
normal thymic parenchyma, appeared only in the
series and will not be considered further.

Results

Thirty six patients were male and 38 female, their ages
ranging from 14 to 82 years, with a mean of 48 years.
Thirty had a cortical thymoma and nine a medullary
thymoma, and in 34 cases the tumour was of mixed
type. The patients with medullary thymidium were
older (mean age 69 years—p < 0·001) than patients
with mixed thymoma (47 years) and cortical thymoma
(43 years). Sixteen patients were symptom free at
presentation. Myasthenia gravis was present in 24
(32%), and its incidence was significantly lower in
patients with medullary thymoma (table 1). There was
a significant difference in stage at presentation among
the three types, most of the cortical thymomas being at
stage III and most of the medullary and mixed
tumours at stage I or II (table 1). Three patients with
cortical thymoma had superior vena caval occlusion
and one had erythroid hypoplasia.

Sixty nine patients had undergone surgery on one
occasion (preceded by mediastinoscopy in five instances);
four had had two operations and one three
operations. Surgical procedures are detailed in table 2.
In one patient with extension of tumour to the
diaphragm and in one with pleural and diaphragmatic
metastases resection was completed by thoracotomy
through the sixth intercostal space after the bulk of the
tumour had been removed via a median sternotomy.
Reconstruction of the superior vena cava by a syn-
thetic prosthesis was necessary in one patient with
cortical thymoma, and an upper lobectomy was
carried out because of direct tumour infiltration in one
case of mixed thymoma. Major postoperative com-
lications occurred in five patients and two died,
giving an operative mortality of 2·7%. Radiotherapy

Table 1  Relation between histological type of thymoma,
myasthenia gravis, and surgical stage

<table>
<thead>
<tr>
<th>Histological type (n)</th>
<th>Stage*</th>
<th>Myasthenia gravis†</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>I</td>
<td>II</td>
</tr>
<tr>
<td>CT (30)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>MxT (34)</td>
<td>12</td>
<td>15</td>
</tr>
<tr>
<td>MT (9)</td>
<td>8</td>
<td>1</td>
</tr>
<tr>
<td>Intrathymic (1)</td>
<td>1</td>
<td>—</td>
</tr>
<tr>
<td>Total</td>
<td>21</td>
<td>22</td>
</tr>
</tbody>
</table>

*Distribution differs significantly between the groups (χ² = 41·684,
p < 0·001).
†Frequency differs significantly between the groups (χ² = 158·78,
p < 0·001).
CT—cortical thymoma; MxT—mixed thymoma; MT—medullary
thymoma.

was given routinely to patients with stage III and IV
disease, and chemotherapy was combined with radio-
therapy in three patients with stage 4 disease.

Radical resection was possible in 60 of the 74 cases;
palliative resection was performed in six and biopsy
alone in eight. All patients with medullary thymomas
underwent radical resection, whereas in 10 of the 30
patients with cortical thymoma and in four of 34 with
mixed thymoma only palliative surgery or biopsy was
possible.

All patients with medullary thymoma are alive two
to 22 years after surgery; for those with mixed and
cortical thymoma the prognosis was significantly
worse, with actuarial survival rates of 65% and 45% at
20 years (table 3 and fig 4). Six patients with stage I or
II cortical or mixed thymoma died of the disease. All
of the patients with relapsing disease that required
further surgery had a cortical thymoma; three had
stage III or IV tumours and had undergone radio-
therapy after the first operation. In two patients stage
II lesions were present and no radiotherapy was
administered. One patient with medullary thymoma
was followed by computed tomography for five years
(fig 5); the tumour remained localised throughout this
period.

Discussion

Neoplasms of the thymus differ in histogenesis, and
epithelial, lymphoid, neuroendocrine, germ cell, and
mesenchymal tumours have all been described.23 The
use of the term thymoma is restricted to thymic
epithelial tumours with minimal or no atypia; those
displaying obvious histological and clinical malign-

Table 2  Surgical procedures performed in the 74 patients
with thymoma

<table>
<thead>
<tr>
<th>No (%) of procedures</th>
<th></th>
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<tbody>
<tr>
<td>Median sternotomy</td>
<td>51 (59)</td>
</tr>
<tr>
<td>Thoracotomy</td>
<td>20 (23)</td>
</tr>
<tr>
<td>Sternal split</td>
<td>7 (9)</td>
</tr>
<tr>
<td>Mediastinoscopy</td>
<td>5 (6)</td>
</tr>
<tr>
<td>Sternotomy and thoracotomy</td>
<td>2 (2)</td>
</tr>
<tr>
<td>Anterior mediastinoscopy</td>
<td>1 (1)</td>
</tr>
</tbody>
</table>


Table 3  Significance (p) of correlation between five, 10, 15,
and 20 year survival and histological type of thymoma

<table>
<thead>
<tr>
<th>Histological types</th>
<th>5 y</th>
<th>10 y</th>
<th>15 y</th>
<th>20 y</th>
</tr>
</thead>
<tbody>
<tr>
<td>CT v MT</td>
<td>&lt; &lt; 0·001</td>
<td>&lt; &lt; 0·001</td>
<td>&lt; &lt; 0·001</td>
<td>&lt; &lt; 0·001</td>
</tr>
<tr>
<td>MT v MxT</td>
<td>0·016</td>
<td>0·003</td>
<td>0·003</td>
<td>0·003</td>
</tr>
<tr>
<td>CT v MxT</td>
<td>0·004</td>
<td>0·015</td>
<td>0·119*</td>
<td>0·119*</td>
</tr>
</tbody>
</table>

*Not significant. CT—cortical thymoma; MT—medullary
thymoma; MxT—mixed thymoma.
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Survival

<table>
<thead>
<tr>
<th>Years from operation</th>
<th>0</th>
<th>5</th>
<th>10</th>
<th>15</th>
<th>20</th>
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</thead>
<tbody>
<tr>
<td>CT</td>
<td>100</td>
<td>85</td>
<td>73</td>
<td>66</td>
<td>65</td>
</tr>
<tr>
<td>MxT</td>
<td>100</td>
<td>76</td>
<td>66</td>
<td>65</td>
<td>59</td>
</tr>
<tr>
<td>MT</td>
<td>100</td>
<td>100</td>
<td>100</td>
<td>100</td>
<td>100</td>
</tr>
<tr>
<td>Total</td>
<td>52</td>
<td>45</td>
<td>45</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

Fig 4 Actuarial survival of patients undergoing operation. MT—medullary thymoma; MxT—mixed thymoma; CT—cortical thymoma.

nancy are more appropriately termed thymic carcinoma. Individual thymomas differ widely in gross morphology and clinical behaviour. Some are apparently benign, well encapsulated lesions, whereas others behave as malignant tumours, with local infiltration and intrathoracic metastases. Histologically, they are generally classified on purely morphological grounds as predominantly lymphocytic, predominantly epithelial, or mixed. This descriptive approach has little prognostic value as all thymomas are epithelial neoplasms, and lymphoid cells, though often present, are an accessory, non-neoplastic component. Not surprisingly therefore studies based on such classifications fail to provide significant clinicopathological correlations.

In 1985 Marino and Müller-Hermelink proposed a new classification based on morphology and immunohistochemistry, in which a histogenesis based on the different subsets of thymic epithelial cells was suggested. Cortical thymomas were related morphologically and phenotypically to thymic cortical epithelial cells, and medullary thymomas to medullary epithelial cells. Mixed thymomas were also described, and further divided into three subgroups—the mixed common type, mixed with cortical predominance, and mixed with medullary predominance. This subdivision seemed to us difficult to apply and sometimes arbitrary. We therefore classified mixed tumours with cortical predominance with cortical thymomas and those with medullary predominance with medullary thymomas.

Analysis of survival data showed significant differences in prognosis among the three main types. Medullary thymoma is a comparatively rare, benign tumour, arising late in life and usually not associated with myasthenia gravis. It presents as an encapsulated tumour, which grows slowly and does not infiltrate surrounding structures or metastasise. It responds well to surgical resection. In our series, even though postoperative radiotherapy was not given, there were no tumour related deaths or recurrences.

Cortical thymoma, on the other hand, behaves as a malignant tumour. Six of our 30 patients had inoperable disease, five intrathoracic metastases, and five local recurrence requiring further surgery. It usually presents in the middle decades of life as a locally invasive tumour at stage III or IV. Resection should be regarded as the first step in a multidisciplinary approach to treatment including radiotherapy, chemotherapy, or both.

Mixed thymoma is intermediate in its behaviour between medullary and cortical thymoma. These tumours may present as stage I or II disease, but should be considered potentially malignant. Post-
operative radiotherapy should always be administered, as tumour related deaths occurred even in patients with stage I disease.

In conclusion, our study of a large series of patients with thymoma shows that medullary, cortical, and mixed tumours differ in both clinical behaviour and prognosis, thus confirming the findings of Müller-Hermelink et al. Medullary thymoma is benign, and may be treated by surgery alone. Cortical thymoma must be considered a malignant tumour, and surgical treatment should be performed as the first step of a multidisciplinary approach that includes postoperative radiotherapy in all patients and chemotherapy in selected cases. Mixed thymoma has a better prognosis than cortical thymoma, but the treatment must be that of a potentially malignant tumour and surgical resection must be followed by radiotherapy even in stage I or II lesions.

References