Cyclic Adenosine 3',5'-Monophosphate-binding Proteins in Human Ovarian Cancer: Correlations with Clinicopathological Features

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ABSTRACT

The regulatory subunits of protein kinase A, or cyclic AMP-binding proteins, were measured in a series of 107 human ovarian tumors (89 malignant, 7 borderline, and 11 benign tumors) and related to tumor clinicopathological features and patient survival.

Total cyclic AMP-binding protein levels were not significantly different between malignant tumors and either borderline or benign tumors. However, serous tumors showed significantly higher levels of total cyclic AMP-binding proteins than other malignant tumors \((P = 0.007)\). Poorly differentiated tumors also possessed significantly higher levels of binding proteins as compared with well/moderately differentiated tumors \((P < 0.01)\). Retrospective analysis of follow-up data also revealed a significant trend for patients with high tumor cyclic AMP-binding proteins to have poorer survival \((P = 0.03)\).

Individual binding proteins were identified by photoaffinity labeling, and the RI \((M_r 48,000)\) protein was expressed as a percentage of total cyclic AMP-binding proteins detected. The percentage of the RI protein was not significantly different among malignant, borderline, or benign pathologies and was not associated with tumor stage, differentiation, or debulk status. The percentage of RI was significantly increased in serous tumors compared to other common epithelial malignancies \((P = 0.01)\). In malignant tumors there was a significant positive correlation between the percentage of the RI protein and total cyclic AMP-binding proteins \((P = 0.01)\).

These data indicate that high tumor levels of cyclic AMP-binding proteins are associated with serious histology, poor differentiation, and poor patient survival.

INTRODUCTION

Cyclic AMP exerts its major effects on cellular proliferation and differentiation via activation of cyclic AMP-dependent protein kinase A \((1)\). The regulatory subunits of protein kinase A, often referred to as cyclic AMP-binding proteins, exist as different subtypes, types I and II, also known as RI and RII \((2)\).

High levels of tumor cyclic AMP-binding proteins may be associated with poor prognosis in patients with breast cancer \((3, 4)\). The mechanisms by which high tumor cyclic AMP-binding protein levels confer poor prognosis is not clear, but it has been suggested that the RI \((M_r 48,000)\) and RII \((M_r 52,000)\) regulatory proteins differentially regulate systems that control cellular proliferation and differentiation \((6, 7)\).

Ovarian cancer is the most fatal of the gynecological malignancies, with a 5-year survival rate of around 20–30% \((8)\), reflecting advanced disease at diagnosis. The limited success of conventional chemotherapeutic regimens has led to the investigation of novel targets for the treatment of this disease. Modulation of the expression of intracellular signal-transducing proteins provides the possibility of a biological approach to disease control. As we have recently reported, ovarian tumors exhibit differing levels of cyclic AMP-binding proteins and variations in the patterns of binding \((5)\), and it was of interest to relate the levels and types of cyclic AMP-binding proteins to tumor clinicopathological features and patient survival.

MATERIALS AND METHODS

Patients. Tissue samples from 107 patients were collected at initial debulking surgery for suspected ovarian malignancy. Upon collection the samples were stored in liquid nitrogen prior to subsequent analyses.

Clinicopathological Details. Tumor pathology, as obtained from patient records, was confirmed on hematoxylin and eosin stained sections. Tumors were classified as either malignant \((n = 89)\), borderline \((n = 11)\) or benign \((n = 7)\). Higher-grade tumors were classified as one steroid cell tumor, one malignant mixed mesodermal tumor, two tumors of mixed epithelial origin, and three unclassified tumors. Of the seven borderline tumors, four were of serous origin and three of mucinous origin. Eleven benign tumors were classified into five mucinous cystadenofibromas, two serous cystadenofibromas, two fibroma/thecomas, one mature cystic teratoma, and one Brenner tumor.

Information was recorded on tumor stage \((\text{International Federation of Gynecology and Obstetrics})\) and differentiation. Early stage tumors were comprised of those presenting at stages I and II, and late stage were comprised of those presenting at stages III and IV. Well-differentiated and moderately differen-

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Cyclic AMP-binding Proteins in Ovarian Cancer

Amersham Life Science) in the absence or presence of varying concentrations of radioinert cyclic AMP (Sigma) to give final concentrations of 0, 10, 20, 40, 80, and 10,000 nmol/liter. Bound cyclic AMP was separated from free nucleotide by filtration at 2 h at 37°C, and radioactivity was measured on a Tricarb liquid scintillation counter (Packard). The number of binding sites and dissociation constants of binding were determined by Scatchard analysis (9), and binding was expressed as fmol of binding protein per mg of cytosol protein (10).

**Photoaffinity Labeling.** Individual binding proteins were determined by photoaffinity labeling using 8-azidoadenosine 3',5'-cyclic [32P]imonomophosphate (ICN Radiochemicals; Ref. 11). Cytosol samples (50 μl), prepared as described above, were incubated with 8-N3-[32P]cyclic AMP (15 μl, 0.4 μmol/liter) and 0.27 M morpholino ethane sulfonic acid with 53 mM magnesium chloride (15 μl; Sigma) in 96-well microtiter plates at room temperature for 1 h in the dark. The reaction mixtures were then irradiated for 30 s at 254 nm by placing a Mineralight UVS-11 hand lamp directly over the plate. The reactions were stopped by the addition of SDS buffer (3% SDS, 15% 2-mercaptoethanol, 30 mM Tris, 30% glycerol, and 1% saturated bromophenol blue). The samples were heated to 90°C for 3 min, and the proteins were resolved electrophoretically on 12% SDS-PAGE with 14C-labeled molecular weight markers for 3 to 4 h at 35 mA. Following electrophoresis, the gels were fixed in 40% methanol, 10% acetic acid, 10% glycerol overnight, dried in a gel drier under vacuum, exposed to preflashed X-ray film (Kodak X-OMAT or Fuji) for 5 to 15 h at ~80°C in autoradiography cassettes fitted with intensifier screens and processed in Kodak X-ray developer and fixer. Autoradiograms were scanned by densitometry, and expression of the M148,000 protein was expressed as a percentage of the total scan.

**Statistics.** Relationships between variables were analyzed using the Mann-Whitney U and Kruskal-Wallis nonparametric tests. Correlations were analyzed using the Spearman rank test. Differences in survival were determined using the Kaplan-Meier method, and groups were compared using the log rank test and χ2 test for trend.

**RESULTS**

**Total Cyclic AMP-binding Proteins, Tumor Clinico-pathological Features, and Patient Survival.** Cyclic AMP-binding proteins were detectable in all ovarian tumors studied irrespective of phenotype, levels varying between 284 and 41,458 fmol/mg protein. Although there was a subgroup of malignant tumors that contained very high levels of cyclic AMP-binding proteins, there was no significant difference in the median levels of binding proteins observed between malignant tumors and either borderline (P = 0.053) or benign tumors (P = 0.07; Fig. 1). Within the malignant tumor group, those of serous histology showed significantly higher levels of total cyclic AMP-binding proteins than endometrioid (P < 0.01) or clear cell (P < 0.05) tumors (Fig. 2). Serous tumors also expressed significantly higher levels of total cyclic AMP-binding proteins than other malignant tumor histologies combined (P = 0.007; data not shown). A significant association was observed between total cyclic AMP-binding protein levels and tumor differentiation, with poorly differentiated tumors displaying higher binding protein levels (P < 0.01; Fig. 3). Cyclic AMP-binding protein levels were not significantly different between early and late stage tumors (P = 0.27) or between debulked and nondebulked tumors (P < 0.08).

In order that tumor cyclic AMP-binding proteins might be related to patient survival, tumor cyclic AMP-binding protein...
Total cyclic AMP-binding protein levels in histological subgroups of malignant ovarian tumors. Serous tumors expressed significantly increased levels of cyclic AMP-binding proteins compared to tumors of endometrioid (P < 0.01) or clear cell (P < 0.05) histology. Horizontal bars, median values.

Fig. 2

Total cyclic AMP-binding protein levels in well/moderately and poorly differentiated ovarian tumors. Poorly differentiated tumors expressed significantly increased levels of cyclic AMP-binding proteins compared to tumors of well/moderate differentiation (P < 0.01). Horizontal bars, median values.

Fig. 3

Type I Protein Kinase A, Total Cyclic AMP-binding Proteins, and Tumor Clinicopathological Features. Photoaffinity labeling of cyclic AMP-binding proteins was performed in 79 malignant ovarian tumor cytosols. A significant positive correlation was observed between the percentage of binding detected as the RI protein, which migrated with a Mr of 48,000, and total cyclic AMP-binding proteins (P = 0.01) in the malignant tumor group (Fig. 5).

The percentage expression of the RI protein was not significantly different among malignant, borderline, or benign tumor pathologies, and in malignant tumors was not associated
Cyclic AMP-binding Proteins in Ovarian Cancer

Fig. 4 Relationship between tumor cyclic AMP-binding protein levels and survival in patients with ovarian cancer. There was a significant trend for poor survival to be associated with high tumor cyclic AMP-binding protein levels (P = 0.03).

Table 1 Total cyclic AMP-binding proteins and clinicopathological features

<table>
<thead>
<tr>
<th>Variable</th>
<th>Classification</th>
<th>&lt;5000</th>
<th>5000-7000</th>
<th>&gt;7000</th>
<th>P*</th>
</tr>
</thead>
<tbody>
<tr>
<td>Histology</td>
<td>Serous</td>
<td>20</td>
<td>7</td>
<td>11</td>
<td>0.01</td>
</tr>
<tr>
<td></td>
<td>Rest</td>
<td>38</td>
<td>10</td>
<td>3</td>
<td></td>
</tr>
<tr>
<td>Stage</td>
<td>I/II</td>
<td>22</td>
<td>3</td>
<td>4</td>
<td>0.29</td>
</tr>
<tr>
<td></td>
<td>II/IV</td>
<td>36</td>
<td>13</td>
<td>10</td>
<td></td>
</tr>
<tr>
<td>Grade</td>
<td>Well/moderate</td>
<td>33</td>
<td>3</td>
<td>2</td>
<td>0.001</td>
</tr>
<tr>
<td></td>
<td>Poor</td>
<td>23</td>
<td>14</td>
<td>11</td>
<td></td>
</tr>
<tr>
<td>Debulk status</td>
<td>&lt;2 cm</td>
<td>43</td>
<td>7</td>
<td>6</td>
<td>0.01</td>
</tr>
<tr>
<td></td>
<td>≥2 cm</td>
<td>14</td>
<td>9</td>
<td>7</td>
<td></td>
</tr>
</tbody>
</table>

* P values determined by χ² test for trend.

with stage, differentiation, or debulk status (Table 2). Expression of the RI protein was not significantly increased in the serous group compared to other malignant histological subgroups combined (P = 0.057; Table 2); however, this did achieve significance in the major subgroup of common epithelial tumors (excluding seven tumors of nonepithelial origin; P = 0.01). No significant association was observed between expression of the RI protein and patient survival (data not shown).

DISCUSSION

This is the first report in which cyclic AMP-binding proteins have been correlated with clinicopathological features of ovarian cancer. These data indicate that high tumor levels of cyclic AMP-binding proteins are associated with a number of indicators of poor prognosis, including serous histology, poor tumor differentiation, and reduced patient survival, suggesting that therapeutic approaches to modulate cyclic AMP-binding protein levels may be beneficial in this disease.

Although there was no significant difference in the median levels of cyclic AMP-binding proteins among malignant, borderline, or benign tumors, there was a subgroup of malignant tumors in which cyclic AMP-binding protein levels were very high. With the exception of one very unusual steroid cell tumor, all tumors which expressed very high levels of cyclic AMP-binding proteins were of serous histology. Indeed, tumors of serous histology showed higher levels of cyclic AMP-binding proteins than did other histological subgroups. Poorly differentiated tumors, many of which were of serous histology, also displayed higher cyclic AMP-binding protein levels than did those of well/moderate differentiation, but no significant differences were observed between levels of binding proteins and tumor stage.

Interestingly, serous tumors also expressed a significantly higher percentage of the RI protein than other common epithelial malignancies, although this did not reach statistical significance compared to all other malignant ovarian tumors. A significant positive correlation was observed between the RI protein and total cyclic AMP-binding proteins in malignant ovarian tumor cytosols, where high binding was associated with a high proportion of the RI phenotype. This is in keeping with previous findings that malignant tumors overexpress RI at the
The prognosis for patients with stage I ovarian cancer, in which the disease is localized to the ovaries and surgery is usually effective, is good. In this study, three stage I patients who did not survive showed tumor cyclic AMP-binding protein levels in excess of 7000 fmol/mg protein, suggesting that, even in stage I disease, high cyclic AMP-binding protein levels may be associated with aggressive tumor behavior.

Observations in breast cancer have also implicated a link between tumor cyclic AMP-binding protein levels and eventual outcome of the disease (3, 4). In these studies, patients who had breast tumor cyclic AMP-binding proteins in excess of 8000 fmol/mg cytosol protein (9–12% of the patients) had an increased chance of disease recurrence compared to patients with lower cyclic AMP-binding protein levels. In light of the findings in breast cancer, our observations in ovarian cancer merit a prospective study to confirm the implication that high tumor cyclic AMP-binding protein levels confer poor prognosis in this disease.

Interestingly, patient survival was not associated with the percentage of total cyclic AMP-binding which was detected as the RI protein and total cyclic AMP-binding proteins, this was by no means absolute, and in a retrospective trawl of the data, and the fact that patient numbers involved in the current survival study are small, it would be premature to comment on the potential prognostic significance of cyclic AMP-binding protein levels in ovarian cancer. It is of particular interest, however, that an association was observed between survival status and high cyclic AMP-binding protein levels in patients with stage I disease. The prognosis for patients with stage I ovarian cancer, in which the disease is localized to the ovaries and surgery is usually effective, is good. In this study, three stage I patients who did not survive showed tumor cyclic AMP-binding protein levels in excess of 7000 fmol/mg protein, suggesting that, even in stage I disease, high cyclic AMP-binding protein levels may be associated with aggressive tumor behavior.

Table 2 Percentage of expression of RI (Mr 48,000 protein) and tumor clinicopathological features

<table>
<thead>
<tr>
<th>Clinicopathology</th>
<th>Median (range)</th>
<th>P*</th>
</tr>
</thead>
<tbody>
<tr>
<td>Pathology</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Malignant</td>
<td>35.9 (8.5–85.2)</td>
<td>0.08</td>
</tr>
<tr>
<td>Borderline</td>
<td>37.0 (28.8–51.7)</td>
<td>0.08</td>
</tr>
<tr>
<td>Benign</td>
<td>42.3 (5.9–70.4)</td>
<td>0.08</td>
</tr>
<tr>
<td>Histology</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Serous</td>
<td>43.4 (11.0–72.0)</td>
<td>0.06</td>
</tr>
<tr>
<td>Rest*</td>
<td>34.1 (8.5–85.2)</td>
<td>0.06</td>
</tr>
<tr>
<td>Stage</td>
<td></td>
<td></td>
</tr>
<tr>
<td>I/II</td>
<td>34.5 (11.0–85.2)</td>
<td>0.93</td>
</tr>
<tr>
<td>III/IV</td>
<td>37.6 (8.5–83.3)</td>
<td>0.93</td>
</tr>
<tr>
<td>Differentiation</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Well/moderate</td>
<td>33.4 (11–85.2)</td>
<td>0.59</td>
</tr>
<tr>
<td>Poor</td>
<td>38.2 (8.5–72.0)</td>
<td>0.59</td>
</tr>
<tr>
<td>Debulking status</td>
<td></td>
<td></td>
</tr>
<tr>
<td>&lt;2 cm</td>
<td>36.7 (11.0–85.2)</td>
<td>0.29</td>
</tr>
<tr>
<td>≥2 cm</td>
<td>36.0 (14.9–63.8)</td>
<td>0.29</td>
</tr>
</tbody>
</table>

* P values were determined by Kruskal-Wallis and Mann-Whitney U tests.

* Rest, all malignant tumor histologies excluding serous.

Serous versus all other common epithelial tumors, P = 0.01.

Fig. 5 A significant positive correlation was observed between total cyclic AMP-binding proteins and the percentage of the RI (Mr 48,000) protein as detected by 8-azidoadenosine 3',5'-cyclic [32P]monophosphate (P < 0.02).
light of this, modulation of the expression of cyclic AMP-binding proteins may provide a possible biological approach to the control of this disease. Strategies to interfere with cyclic AMP-binding proteins, e.g., 8-chloro cyclic AMP, a cyclic AMP analogue which modulates the RI subunit (7), have been shown to cause growth inhibition in a number of cancer cell lines (17, 18) and also in nude mice xenograft models (12, 19, 20). Phase I clinical trials of this drug are currently under way (21, 22). Down-regulation of RI using antisense oligodeoxynucleotides has also produced growth inhibition both in vitro (23) and in vivo (24). We are currently exploring these methodologies as potential therapeutic strategies in ovarian cancer.

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