Proliferating Cell Nuclear Antigen Expression in the Survival of Astrocytoma Patients

L.C. Ang, M. Plewes, L. Tan, H. Begley, A. Agranovich and D. Shul

Abstract: The PC10, a monoclonal antibody against proliferating cell nuclear antigen (PCNA), is known to show immunoreactivity in routinely processed paraffin embedded tissue. This antibody was applied to 72 astrocytic tumours from surgical biopsy material obtained in a ten year period. The PCNA labelling index (LI) obtained by image analysis was compared with patient's survival, age at diagnosis, and Karnofsky score as well as the histological grade of tumour. The survival analysis shows that patients with tumour PCNA LI of more than 6% have significantly poorer prognosis than those with 6% and below. In addition, there is also good correlation between PCNA LI with age, Karnofsky and tumour grade. This study suggests that although the PCNA expression of astrocytoma could be a useful predictor of patient's outcome, it is not an independent prognostic factor but has derived its statistical association with survival secondarily through its relationship with tumour grade, age and Karnofsky score.

Resume: Expression de l'antigène nucléaire des cellules en prolifération comme prédicteur de la survie chez les patients atteints d'astrocytome. Le PC10, un anticorps monoclonal dirigé contre l'antigène nucléaire des cellules en prolifération (PCNA) démontre une immunoréactivité dans les tissus inclus dans la paraffine par la technique de routine. Cet anticorps a été appliqué à 72 tumeurs astrocytaires provenant de biopsies chirurgicales obtenues sur une période de dix ans. L'indice de marquage par le PCNA obtenu par analyse d'image a été analysé en fonction de la survie des patients, à l'âge au moment du diagnostic, au score de Karnofsky et au stade de la tumeur. L'analyse de survie montre que les patients ayant un indice de plus de 6% ont un pronostic significativement plus sombre que ceux dont l'indice est de 6% ou moins. De plus, il existe également une bonne corrélation entre l'indice de marquage par le PCNA et l'âge, le score de Karnofsky et le stade de la tumeur. Cette étude suggère que, bien que l'expression du PCNA de l'astrocytome puisse être un prédicteur utile du devenir du patient, ce n'est pas un facteur pronostique indépendant parce que son association statistique avec la survie est secondaire à sa relation avec le stade de la tumeur, l'âge et le score de Karnofsky.


Tumours of astrocytic lineage are the most common primary neoplasm of the central nervous system. The histological grading is considered as an important factor in predicting the outcome of patients with such tumours. But there are various systems used in grading these tumours rendering comparisons between different treatment centres very difficult. Even if one uniform system is used in grading, the results could be very subjective depending on the experience and training of individual pathologists. Because of these reasons, various markers of cellular proliferative activities are introduced with the hope of grading these tumours more objectively. The PC10 a monoclonal antibody against proliferating cell nuclear antigen (PCNA) could be successfully applied to various tumour tissues, including archival material, which have been already fixed in formalin and then embedded in paraffin.

PCNA, an auxiliary protein of DNA-delta-polymerase at the site of DNA replication during cell cycle, attains the highest concentration during the proliferative phases of the cycle. PCNA expression in a number of tumours such as gastric carcinoma, gut lymphoma and haemangiopericytoma has also been linked to the prognostic outcome of such tumours. A handful of previous reports have also studied the correlation of PCNA LI (labelling index) with the grades of astrocytoma but these results appeared rather conflicting. Only one other study with smaller number of astrocytomas (42 cases) has looked into the correlation of PCNA LI and survival of astrocytoma patients. In the present study we examine the correlation of PCNA LI

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with astrocytoma patients’ survival and 3 other prognostic factors namely 1) age at diagnosis, 2) histological grade and 3) Karnofsky score. A program using the IBAS image analysis system was designed to measure the PCNA LI in order to obtain more objective results.

MATERIALS AND METHODS

A total of 72 cases of astrocytoma including glioblastoma multiforme were identified from the surgical biopsy files of the Royal University Hospital of Saskatoon through the period of 1 July 1980 to 30 June 1990. All these cases had sufficient material for PC10 immunohistochemistry and were fixed in 4% buffered formaldehyde for approximately 24 hours before processing. These 72 tumours were reviewed and graded histologically using the method proposed by Daumas-Duport and divided into Grade I (12 cases), Grade II (14 cases) and Grade IV (46 cases). This method is based upon the presence or absence of four morphological criteria: nuclear atypia, mitoses, endothelial proliferation and tumour necrosis. The summary score for a tumour is translated into a grade as follows: 0 criteria = grade I, 1 criterion = grade II, 2 criteria = grade III, 3 or 4 criteria = grade IV. We had also excluded cases with diagnoses of juvenile pilocytic astrocytoma, optic glioma, subependymal giant cell astrocytoma, pleomorphic xanthoastrocytoma, gangliogioma and mixed glioma with a prominent oligodendrogliomatous component. All these types of primary glial tumours were excluded because they have distinct pathological features and different biological behaviour compared to the ordinary astrocytoma.

Formalin-fixed, paraffin-embedded blocks of tumour tissue were cut at 6 μm, attached to coated slides (3 -aminopropyltriethoxy-silane - Sigma), blotted, and air dried overnight at room temperature. These slides were hydrated and washed in PBS. Non-specific background staining was reduced by the application of 20% normal horse serum. The sections were incubated overnight at 4°C with monoclonal anti-PCNA antibody (clone PC10-Novocastra Laboratories Ltd., Newcastle-upon-Tyne, UK) diluted 1:200. The specificity of this antibody to PCNA in vertebrate species has been established in Western blot, immunoprecipitation and enzyme-linked immunosorbent assays (ELISA).

Slides were washed in PBS, then incubated 60 minutes at room temperature with biotinylated anti-mouse IgG 1:225 (Vector Laboratories), then treated with 2% osmium tetroxide for 10 minutes, washed in water and counterstained with 1% methyl green. For positive controls specimens of tonsils resected surgically were used and normal cerebral cortices were the negative control.

The PCNA immunostained sections of the tumours were examined qualitatively under the light microscope (L.C.A) and areas (non-hemorrhagic and non-necrotic) with highest concentration of positive nuclei were selected for image analysis. The morphometric measurements were performed (H.B.) with the IBAS system (Kontron Electronic, Germany) utilizing an Axioplan microscope (Zeiss, West Germany) and a MTI camera (Dage-MTI Inc., USA). Images of each tumour at 200x magnification were captured on the computer screen and subsequently processed by scaling the grey level distribution with 0 representing the darkest and 255 representing the lightest level. Grey levels (0 to 95) were selected to enhance all the nuclei in the field and then the nuclei were discriminated and counted. Lower grey levels (0 to 73) were selected to enhance only the PCNA stained nuclei which were also discriminated and counted. In each case at least 500 nuclei (mean 850) were counted and the PCNA LI was calculated as a percentage of positively stained nuclei over the total number of tumour cell nuclei counted.14-17

The clinical data from these 72 cases were obtained from the medical records of the Saskatoon Cancer Clinic and Royal University Hospital of Saskatoon. The data examined in this study included age at diagnosis, sex, Karnofsky score at admission, presenting symptoms, location of tumours, and survival. The age distribution of the various patients with the different grades of astrocytoma is shown in the Table. All patients were followed up for a minimum of two years with the longest follow-up being 9 years.

Table. Age distribution of astrocytoma patients.

<table>
<thead>
<tr>
<th>Age Group (Years)</th>
<th>Grade II</th>
<th>Grade III</th>
<th>Grade IV</th>
<th>Total Number</th>
</tr>
</thead>
<tbody>
<tr>
<td>&lt;10</td>
<td>3</td>
<td>1</td>
<td>0</td>
<td>4</td>
</tr>
<tr>
<td>10-19</td>
<td>1</td>
<td>0</td>
<td>2</td>
<td>3</td>
</tr>
<tr>
<td>20-29</td>
<td>1</td>
<td>1</td>
<td>0</td>
<td>2</td>
</tr>
<tr>
<td>30-39</td>
<td>3</td>
<td>3</td>
<td>1</td>
<td>7</td>
</tr>
<tr>
<td>40-49</td>
<td>3</td>
<td>2</td>
<td>2</td>
<td>7</td>
</tr>
<tr>
<td>50-59</td>
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<td>3</td>
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<td>16</td>
</tr>
<tr>
<td>&gt;=60</td>
<td>1</td>
<td>4</td>
<td></td>
<td>28</td>
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</table>

RESULTS

As previously reported the PC10 immunostaining in the nuclei of tumour cells could be either diffuse, granular or mixture of both.6,14-17 The staining intensity is also highly variable and there is marked regional variation in the distribution of immunostaining within the same tumour.6 The range of PCNA LI in this study is between 0.2% to 61%. Examples of immunoreactivity in grade II, grade III and grade IV astrocytoma are shown in Figure 1 (a, b and c).

The mean PCNA LI for patients diagnosed with Grade II, Grade III and Grade IV astrocytoma are 3.17% ± 0.81 (SEM), 9.71% ± 2.47(SEM) and 18.01% ± 1.98(SEM) respectively. Hence, the higher grade tumours tend to have higher labelling index. A scattergram of the relationship between tumour grade and PCNA LI with the regression line drawn in shows significant correlation with coefficient of correlation of + 0.452 which is statistically significant (P < 0.001) (Figure 2).

The survival after diagnosis of patients with PCNA LI of tumour above 6% (46 patients) is compared to the group of
patients with 6% or less (26 patients) (Figure 3). The difference in survival between these 2 groups is highly significant statistically (P < 0.0001). The median survival for patients with PCNA LI more than 6% is about 5 months compared to 6 years (73 months) in those with 6% or less. In the first group none of the patients were alive at the time of study and in the second group more than 35% were alive. Apparently, PCNA LI is a good predictor of patients’ outcome.

The coefficient of correlation between PCNA LI and age of patients at time of diagnosis is +0.40 which is statistically significant (P < 0.001). The significant positive coefficient of correlations suggests that PCNA LI tends to increase in tumours from older patients. There is also significant negative coefficient of correlation −0.336 (p < 0.004) which suggests that PCNA LI tends to decrease with increasing Karnofsky score. A comparison of PCNA LI for tumours arising from various anatomical sites such as the different lobes, thalamus, basal ganglia, brain stem, cerebellum etc. was not done because these tumours happened to fall into too numerous categories according to sites resulting in very small numbers in each category, making valid analysis not feasible.

Using the Cox Proportional Hazard model for univariate survival analysis which displays the degree of association between each of 5 factors (PCNA LI, age, histological grade, Karnofsky score and sex) and patients’ survival, the results are expressed as approximate chi-square to enter or remove and its p-value. As shown in Figure 4, the histological grading using the Daumas-Dupont criteria is the most significant prognostic factor, followed by age, then Karnofsky, and then the PCNA LI. Based on the multivariate approach with the Cox Proportional Hazard model, we found that once histological grade, age and Karnofsky have been entered into the survival model, PCNA LI is no longer a significant additional prognostic factor.

**DISCUSSION**

According to Bravo and MacDonald-Bravo there are at least two forms of PCNA antigens; a soluble form, not involved in cell proliferation and an insoluble form associated with cell proliferation including DNA replication. The PC10 monoclonal antibody when used on formalin-fixed normal tissue is supposed to label only the form associated with cell proliferation. But it is also quite obvious that in some neoplasms such as breast and
Although PC10 has been shown to be useful for studies on archival tissue embedded in paraffin, the interpretation of the results may be affected by duration of formalin fixation and the heterogeneity in the immunoreactivity. The fact that the PCNA immunostaining decreases significantly with prolonged formalin fixation makes it necessary to exclude tumour tissue fixed in formalin for longer than 24 hours before processing. The marked heterogeneity in distribution of immunoreactivity would mean that the PCNA LI is also largely dependent on biopsy sampling and in a small biopsy the labelling may not be fully representative of the entire tumour. The heterogeneity in intensity of immunostaining could also lead to interpreting areas with low reactivity as negative areas thus introducing an element of subjectivity in interpretation. In an attempt to minimize the regional variability in PC10 we had quantitated those areas with highest concentration of positive immunostaining in all cases and used the values obtained for calculating the PCNA LI. Moreover, the regions with the highest proliferative activity in a tumour would be clinically most relevant to the patient’s outcome in terms of the rate of growth, progression and spread to the tumour.

Another drawback is observer bias that would occur with visual analysis of PCNA LI and this could preclude its routine use and furthermore it is very time consuming to count such large numbers of nuclei manually. In order to eliminate observer bias, a program using image analysis was designed for quantifying the PCNA LI in this study.

Previous studies on glioma, however have not yielded consensus on the correlation of PCNA expression with tumour grading. Louis et al. using the 19F4 monoclonal antibody against PCNA which is applicable only to cryostat sections have found that though the PCNA LI for higher grade (grade IV) tumours were appropriately high but it could not distinguish between the grade III and the lower grade tumours (i.e. grade I and II). Furthermore, they found that the 19F4 antibody produced weaker and less consistent staining than Ki-67 in cryostat sections. On the other hand using the PC10 Allegranza et al., Karamitopoulou et al. and Theunissen and Blaauw have found that the PCNA LI correlated well with the histological grades of astrocytoma. Only the last study has correlated the other clinical parameters such as survival, age at diagnosis and Karnofsky score to the PCNA immunoreactivity.

Our study demonstrates a significant correlation existed between PCNA LI and certain known prognostic factors for astrocytoma such as histological grade, age at the time of diagnosis and Karnofsky score. PCNA LI is also well correlated with the survival of tumour patients, with patients having PCNA LI of 6% or less showing significant better outcome than those with labelling index above 6%. Because of the relatively small number of low grade astrocytomas and the short follow-up for some of them, we could not assess the value of PCNA LI in predicting which of these would remain as low grade tumours and which would progress to higher grades over a longer period. The univariate survival analysis using the Cox Proportional Hazard model has revealed that PCNA LI is less significant as a prognostic factor than histological grade. Furthermore, from the multivariate analysis we have concluded that PCNA LI is not an independent prognostic factor but it has derived its statistical association with survival secondarily through its relationship with tumour grade, age and Karnofsky score. Our findings support those of Theunissen and Blaauw in suggesting that while...
the PC10 immunoreactivity could be used as an additional method for assessing the prognosis of astrocytoma, it does not contribute any more reliable information on the outcome than histological grading based on the criteria of Daumas-Duport et al.3

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