Comparative study of histopathology and immunohistochemistry of indefinite round cell cutaneous tumors and characterization of canine lymphoma

[Estudo comparativo de histopatologia e imuno-histoquímica de tumores indefinidos de células redondas e caracterização de linfoma canino]

L.H.A. Machado¹, M.I.P. Palumbo¹*, F.S. Zahn¹, R.L. Amorim¹, M.R. Farias², J. Werner³, R. Torres Neto¹, J.C. Rodrigues³, F.C. Oliveira¹

¹Faculdade de Medicina Veterinária e Zootecnia – Universidade Estadual Paulista – FCAV-unesp – Botucatu, SP
²Pontifícia Universidade Católica do Paraná – PUCPR – Curitiba, PR
³Autônomo

ABSTRACT

With the purpose of shedding light on some doubts in veterinary oncology, the present article intends to compare the results of histopathological and immunohistochemical examinations of unspecific round cell neoplasia, to realize immunophenotyping of canine lymphoma cases, to establish the T or B origin of neoplastic cells, and to determine the degree of proliferation and apoptosis of lymphomas by immunohistochemistry. Of 11 animals presenting immunohistochemical diagnosis of lymphoma, five had been diagnosed as Lymphoma by HE staining of histopathological slides and six had been classified as unspecific round cell neoplasia. All cases submitted to immunohistochemical examination were T-cell lymphomas. There was a positive correlation between cell proliferation and apoptosis. The comparison among histopathological and immunohistochemical results obtained in the cases examined in the present study suggested that immunohistochemistry is essential for the differentiation of round cell neoplasia.

Keywords: dog, histopathology, immunohistochemistry, round cell neoplasia, lymphoma

INTRODUCTION

Lymphoma is one of the most frequent neoplasias in dogs, which may also be found under the names lymphosarcoma and malignant lymphoma, and represents approximately 7-24% of all canine neoplasia and 83% of hematopoietic tumors (Teske et al., 1994). One of the methods used to classify canine lymphoma is immunohistochemistry, which demonstrates the presence of cell type marking antigens (Milner et al., 1996). Its use in Veterinary Medicine is still restricting because of the high cost and the absence of specific markers in some cases (Fisher et al., 1995), although this reality is changing (Soares and Arias 1999).
Immunohistochemistry has proved to be an important tool in precise diagnosis (Dobson et al., 2001), making it possible to classify type B or T origin and characterize the degree of maturation of lymphoid neoplastic cells.

The technique has been used successfully in histological slides of paraffin- included tissue, making B-cell lymphomas with monoclonal antibody anti-mb1 (CD79a) and T-cell lymphomas with polyclonal antibody anti-CD3 (Bacchi and Gown, 1993; Fournel- Fleury et al., 1997; Fournel- Fleury et al., 2002; Bhang et al., 2006; Cardoso et al., 2006).

Because canine lymphoma has similarities with non-Hodgkins human lymphoma, it is possible to use its cytohistological and immunophenotypic classifications in canine lymphoma, associating these data with clinical findings, thus improving treatment and the determination of prognosis and survival time in animals (Fisher et al., 1995; Kiupel et al., 1999). Also, the results obtained in animals may be used as experimental models for human oncology (MacEwen, 1990).

Tumor growth is determined by three main factors: cell cycle length, percentage of proliferating cells and the number of cells lost by apoptosis (Franks, 1990). Cell proliferation, with mass formation, is one of the characteristics of neoplasia, which does not depend on the primary cause, as a direct consequence of disturbance in the control of cell cycle (Bacchi and Gown, 1993). Detection and quantification of proliferating cells have been considered important prognostic parameters in oncology, and may be used to identify neoplasia or to evaluate its malignancy (Bacchi and Gown, 1993; Rabenhorst et al., 1993). The mouse anti-Ki-67 monoclonal antibody clone MIB-1 has been used as a marker of cell proliferation in many tumors, both in humans and animals. The evaluation of cell proliferation by Ki-67 index is highly predictive of the behavior of several tumors (Abadie et al., 1999; Sakai et al., 2002).

In canine mast cell tumors, the immunoreactivity for Ki-67 is correlated to the histological grade, being useful in the evaluation of cell proliferation and in the determination of the degree of cell differentiation (Teske et al., 1994). Histocytooma is one of the most important and challenging differential diagnosis of lymphoma (Bhang et al., 2006; Cardoso et al., 2006), corresponding to 19.6% of cutaneous tumors (Gross et al., 1992).

Another important variable related to the biological behavior of proliferative lesions is the index of apoptosis (Sano et al., 2004). Apoptosis has been studied by the expression of caspase, which are enzymes directly related to apoptosis, being present in most cells’ cytoplasm in the inactive form, as a single chain of polypeptides that is broken when apoptosis occurs (Sano et al., 2004). The inhibition of caspase activity may delay or impair cell death by apoptosis (Nicholson, 1999).
The most studied caspase is caspase-3, known to be related to apoptosis in many hematopoietic disturbances in humans; both the level of caspase-3 expression and the form of its intracytoplasmic marking are related to tumoral progression. Highly aggressive neoplasia present lower caspase-3 cytoplasmic expression than low grade neoplasia (Porter and Jänicke, 1999).

The present article aimed to compare the results of histopathological and immunohistochemical examinations of unspecific round cell neoplasia, to realize immunophenotyping of canine lymphoma cases, to establish the T or B origin of neoplastic cells, to determine the degree of proliferation and apoptosis of lymphomas by immunohistochemistry.

MATERIAL AND METHODS

Cases of cutaneous lymphoma and unspecific round cell neoplasia were selected in histopathological archives. Initial selection, as well as initial histopathological examinations were performed at Werner and Werner Veterinary Pathology Laboratory and immunohistochemical examinations were performed at the Service of Veterinary Pathology of FMVZ – UNESP – Botucatu.

The fragments were identified and fixed in 10% buffered formalin, then submitted to histological routine procedures and included in paraffin. Histological cuts of 3µm were stained by Hematoxylin-Eosin (HE) (Luna, 1968) and observed under light microscopy.

The study of tumor cell lines was performed using standardized techniques at the laboratory of the Service of Veterinary Pathology of FMVZ – UNESP – Botucatu, with the following panel of antibodies: Anti-CD3, Anti-CD79a, Caspase-3 and MIB-1. For Anti-CD3 and anti CD-79a antibodies, only positive and negative criteria were established; for caspase-3 and MIB-1, cells were counted in all slides, whether immunolabeled (positive) or not (negative).

Data were submitted to Spearman correlation coefficient, which is applicable to non-parametric variables, and to Pearson coefficient. The results of the Pearson test were not presented because no correlation was observed.

RESULTS

The results of histological and immunohistochemical examinations of the confirmed cases of lymphoma and non-confirmed cases of cutaneous lymphoma are presented in Table 1 and 2, respectively (Table 1, Table 2).

Table 1. Results of histological and immunohistochemical evaluation of confirmed cases of canine lymphoma

<table>
<thead>
<tr>
<th>Dog</th>
<th>HE</th>
<th>HE</th>
<th>MIB</th>
<th>Caspase</th>
<th>CD3</th>
<th>CD79</th>
<th>Immunohistochemical diagnosis</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>Lymphoma</td>
<td>223</td>
<td>53</td>
<td>10</td>
<td>171</td>
<td>X</td>
<td>T-cell lymphoma</td>
</tr>
<tr>
<td>2</td>
<td>Lymphoma</td>
<td>55</td>
<td>159</td>
<td>0</td>
<td>0</td>
<td>X</td>
<td>T-cell lymphoma</td>
</tr>
<tr>
<td>3</td>
<td>IRCN</td>
<td>28</td>
<td>110</td>
<td>40</td>
<td>184</td>
<td>X</td>
<td>T-cell lymphoma</td>
</tr>
<tr>
<td>4</td>
<td>IRCN</td>
<td>2</td>
<td>387</td>
<td>4</td>
<td>253</td>
<td>X</td>
<td>T-cell lymphoma</td>
</tr>
<tr>
<td>5</td>
<td>IRCN</td>
<td>57</td>
<td>270</td>
<td>110</td>
<td>160</td>
<td>X</td>
<td>T-cell lymphoma</td>
</tr>
<tr>
<td>6</td>
<td>IRCN</td>
<td>125</td>
<td>274</td>
<td>9</td>
<td>242</td>
<td>X</td>
<td>T-cell lymphoma</td>
</tr>
<tr>
<td>7</td>
<td>IRCN</td>
<td>110</td>
<td>159</td>
<td>8</td>
<td>223</td>
<td>X</td>
<td>T-cell lymphoma</td>
</tr>
<tr>
<td>8</td>
<td>IRCN</td>
<td>11</td>
<td>315</td>
<td>11</td>
<td>264</td>
<td>X</td>
<td>T-cell lymphoma</td>
</tr>
<tr>
<td>9</td>
<td>Lymphoma</td>
<td>51</td>
<td>142</td>
<td>35</td>
<td>74</td>
<td>X</td>
<td>T-cell lymphoma</td>
</tr>
<tr>
<td>10</td>
<td>Lymphoma</td>
<td>193</td>
<td>215</td>
<td>7</td>
<td>263</td>
<td>X</td>
<td>T-cell lymphoma</td>
</tr>
<tr>
<td>11</td>
<td>Lymphoma</td>
<td>113</td>
<td>195</td>
<td>24</td>
<td>230</td>
<td>X</td>
<td>T-cell lymphoma</td>
</tr>
<tr>
<td>Total</td>
<td></td>
<td>968</td>
<td>2279</td>
<td>258</td>
<td>2064</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

HE = hematoxylin/eosin. IRCN = inespecific round cell neoplasia. Same letter in a row indicates positive correlation between variables (r=0.66059, P=0.0269).
Table 2. Results of histological and immunohistochemical evaluation of non-confirmed cases of cutaneous lymphoma

<table>
<thead>
<tr>
<th>Cão</th>
<th>HE diagnosis</th>
<th>MIB</th>
<th>Caspase</th>
<th>CD3</th>
<th>CD79</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td>Negative</td>
<td>Positive</td>
<td>Negative</td>
<td>Positive</td>
</tr>
<tr>
<td>1</td>
<td>Round cell neoplasia</td>
<td>80</td>
<td>290</td>
<td>14</td>
<td>330</td>
</tr>
<tr>
<td>2</td>
<td>Round cell neoplasia</td>
<td>53</td>
<td>615</td>
<td>6</td>
<td>387</td>
</tr>
<tr>
<td>3</td>
<td>Plasma cell neoplasia</td>
<td>112</td>
<td>315</td>
<td>20</td>
<td>392</td>
</tr>
<tr>
<td>4</td>
<td>Round cell neoplasia</td>
<td>423</td>
<td>156</td>
<td>13</td>
<td>360</td>
</tr>
</tbody>
</table>

**DISCUSSION**

The evaluation of fine needle aspirates by a qualified pathologist may be adequate to diagnose canine lymphoma, but conclusive histological confirmation is still recommended (Vail, 2004). From 11 animals presenting immunohistochemical diagnosis of lymphoma, five had been diagnosed as Lymphoma by HE staining of histopathological slides and six had been classified as unspecific round cell neoplasia (IRCN), thus demonstrating that, in many cases, more specific diagnostic techniques are necessary to allow better treatment and prognosis establishment.

Immunohistochemistry has shown to be an important tool in precise disease diagnosis (Dobson et al., 2001), making it possible to classify either T or B-cell Lymphoma and also the degree of neoplastic lymphoid cell maturation. The technique has been successfully applied in paraffin-included histological material, marking B-cell lymphomas with monoclonal antibody anti-μB1 (CD79a) and T-cell lymphomas with polyclonal antibody anti-CD3 (Fournel-Fleury et al., 1997; 2002; Bhang et al., 2006; Cardoso et al., 2006; Arespacochaga et al., 2007). According to the literature, most canine lymphomas are type B (Teske et al., 1994; Fournel-Fleury et al., 1997; Arespacochaga et al., 2007), but in the present study, all cases submitted to immunohistochemical examination were T-cell lymphomas, as they were marked only by polyclonal antibody anti-CD3. In a study of 40 cases of canine lymphoma, similar proportions (42.5% each) of T and B-cell cases were reported and 15% of mixed T/B lymphomas were also observed (Suzano et al., 2008). This difference among reports may be due to the fact that in the present study, all the 11 cases were cutaneous lymphomas, while in the other reports extracutaneous lymphomas, considered to have better prognoses, were also included.

The mouse anti-Ki-67 monoclonal antibody clone MIB-1 has been applied as a marker for cell proliferation in many tumors, both in humans and animals. Apoptosis has been evaluated by caspase expression, which is directly related to apoptosis and is present in most cells’ cytoplasm in inactive forms that, when the single polypeptides chain is broken, initiates apoptosis (Luna, 1968). In the present study there was a positive correlation between cell proliferation and apoptosis. Further studies are already planned in order to associate immunohistochemical results and clinical response to chemotherapy in order to establish whether this correlation could reflect in prognosis. The presence of apoptosis may facilitate this response, making the tumor more susceptible to chemotherapy.

**CONCLUSION**

The comparison among histopathological and immunohistochemical results obtained in the cases examined in the present study suggested that immunohistochemistry is essential for the differentiation of round cell neoplasia. More studies on cutaneous lymphomas are necessary to determine which are the most involved cell types, their proliferative and apoptotic characteristics and the association of those findings with clinical evolution of the patient.
REFERENCES


