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| 题目       | 横田腫瘍の成長に関する研究

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Growth of Yoshida Sarcoma in the Thymus
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INTRODUCTION

One of the most prominent features of the behavior of carcinomas is their great tendency to produce secondary growth in regional lymph nodes. They appear to have found a comfortable home in the nodes and enjoy there an abundance of nutrients for growth; lymphoid tissue containing carcinoma cells may function as a feeder for cancer cells.

There are published many reports on susceptibility of the spleen, liver, kidney and muscle to transplanted tumor, but none allows a comparison of tumor growth in the thymus with that in other organs.

In the present study, the thymus was chosen as one of the lymphatic apparatuses, since 92% of the cells composing the thymus tissue are considered to be lymphocytes. The present paper will deal with susceptibility of the thymus to YOSHIDA sarcoma cells comparing with other tissues or organs.

MATERIALS AND METHODS

The animals used were male Donryu rats, aged 30 days, weighing 45 to 55 g. They were supplied commercially and fed ad libitum on water and stock diet MF, manufactured by Oriental Kobo Co.

Ascites tumor cells of YOSHIDA sarcoma were obtained through the courtesy of the laboratory of TAKEDA Chemical Co., Osaka, and maintained in our laboratory by successive inoculation into the abdominal cavity of Donryu rats. Prior to each experiment, ascites cells were withdrawn from the rats' peritoneal cavity 4 to 5 days after transplantation and the cell suspension was prepared so as to contain 10,000 cells in an aliquot of 1 μl of TYRODE solution.

The thymus of Donryu rats was exposed under ether anesthesia through a upper median sternal incision and 1 μl of the cell suspension was injected into the thymus, using a microsyringe.
For the comparison of susceptibility to tumor, the subcutaneous tissue, liver, spleen
and peritoneal cavity were chosen and transplanted with an equal number of cells. In
the case of the liver and spleen, Aron Alpha, Sankyo Pharmaceutical Co., was applied to
the needle hole after injection to prevent leakage of inoculated cells into the peritoneal
cavity.

“Take” of tumor was determined by visible growth of tumor in the organs except
for the liver in which tumor mass was not clearly recognized but change of color disclosed
proliferation of tumor cells in the region of transplantation.

RESULTS

1. YOSHIDA Sarcoma Tumor in the Thymus:

YOSHIDA sarcoma cells proliferated in the thymus and became visible as a mass on
the 4th day after transplantation, involving the whole thymus by the 9th day and result-
ing in death of the animals. Necropsy showed an enlarged thymus being replaced wholly
by YOSHIDA sarcoma cells, which extended as far as the aorta and trachea (Plate 1).

Microscopic observation of specimens on the 7th day after transplantation showed that
YOSHIDA sarcoma invaded a wide area of the thymus, replacing normal tissue cells with tumor
cells. At the margin of tumor mass sarcoma cells seemed to have phagocytized thymo-
cytes, because some of the tumor cells contained, in their cytoplasm, fragments stained by
hematoxylin which were considered to have been derived from nuclei of destroyed thymocytes (Plates
3 and 1).

2. Relative Susceptibility of the Thymus to
Tumor Implantation:

YOSHIDA sarcoma cells were implanted into the thymus, subcutaneous tissue, spleen, liver and perito-
eal cavity, and incidence of “Take” and growth of
tumor in these organs of tissues were compared with those in the thymus on the 7th day after implantation. The data are indicated in Table 1 and 2. It can be seen in the tables that the thymus group showed higher incidence of “Take” and larger size of tumor than the other organs. In the organs mentioned above, except for the thymus, incidence of “Take” and size of tumor were nearly equal on the 7th day after implantation, and the inoculated tumor cells grew at almost the same rate.

Further observation of the subcutaneous

**Plate 3** Yoshida sarcoma cells are seen infiltrating into the thymus tissue. Pyknotic thymocytes remain like islands. Hematoxylin and eosin, × 600.

**Plate 4** Yoshida sarcoma cells, possibly phagocytizing thymocytes; some of the sarcoma cells are seen to contain fragments which are thought to be derived from the nuclei of destroyed thymocytes. Hematoxylin and eosin, × 1,500.

**Table 1** Incidence of “Take” of Y. S. tumor on the 7th day after implantation

<table>
<thead>
<tr>
<th>Organ and tissue</th>
<th>Incidence of “Take”</th>
<th>%</th>
</tr>
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<tbody>
<tr>
<td>Thymus</td>
<td>14/11</td>
<td>100</td>
</tr>
<tr>
<td>Subcutaneous tissue</td>
<td>21/18</td>
<td>50</td>
</tr>
<tr>
<td>Spleen</td>
<td>7/16</td>
<td>44</td>
</tr>
<tr>
<td>Liver</td>
<td>3/10</td>
<td>30</td>
</tr>
<tr>
<td>Peritoneal cavity</td>
<td>6/11</td>
<td>55</td>
</tr>
</tbody>
</table>

**Table 2** Average size and weight on the 7th day after implantation

<table>
<thead>
<tr>
<th>Organ and tissue</th>
<th>Number of animals</th>
<th>Average size of tumor (mm³)</th>
<th>Average weight of tumor (mg)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Thymus</td>
<td>5</td>
<td>2169</td>
<td>561</td>
</tr>
<tr>
<td>Subcutaneous tissue</td>
<td>11</td>
<td>136</td>
<td>32</td>
</tr>
<tr>
<td>Spleen</td>
<td>7</td>
<td>161</td>
<td></td>
</tr>
<tr>
<td>Liver</td>
<td>3</td>
<td>110</td>
<td></td>
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Size : (Length × Width)² / 2.

group indicated that the size of these tumors on the 9th day was as larger as those in the thymus on the 5th to 6th day when the mass became visible (Plate 2). This suggests that the thymus was more favorable place for growth of the tumor than the subcutaneous tissue.

**DISCUSSION**

DeLong and Coman have provided evidence of equal susceptibility of the rat to tumor transplantation in the muscle, kidney, liver and spleen. They found no difference among the weights of tumor and incidence of "Take" when fragments of Walker rat carcinoma 256 were implanted in those organs. The results of the present experiment, however, suggest higher susceptibility of the thymus to tumor transplantation than the subcutaneous tissue, spleen, liver and peritoneal cavity, i.e., the tumor grew more rapidly in the thymus and were generally larger in size than those in the other organs. What does this higher susceptibility of the thymus depend on? Existence of a certain growth-promoting substance like Promine or absence of immune reaction in the thymus have been suggested by some workers. Attention, however, should be directed to the fact that the thymus is a lymphoid organ and composed mostly of lymphocytes. The thymus is considered to be a lymphocyte producer and 92% of the cells composing the thymus are lymphocytes. Particular affinity of tumors to the lymphatic apparatus has tacitly been recognized by many investigators.

Leighton has suggested that lymphatic tissue might play a nutritive role for secondary growth of tumor as a feeding trough; biological interaction and passage of nutritive material like DNA have been observed experimentally between thymocytes or lymphocytes and other cells.

Fisher and Fisher, in an electron microscopic study of hepatic metastases of Walker tumor, have revealed that tumor cells, possibly by a pinocytotic mechanism, received various nutrients directly from the host's hepatic cells. Microscopic appearance of Yoshida sarcoma cells in the margin of the tumor mass suggested the same mechanism might be operating also in the present case.

**SUMMARY**

Yoshida sarcoma cells were implanted into the rat thymus and other tissues, and the growth of tumor was observed for the purpose of comparing tissue susceptibility. The thymus was more susceptible to implantation of Yoshida sarcoma as well as more favorable for growth of the tumor than the subcutaneous tissue, liver, spleen and peritoneal cavity.

**ACKNOWLEDGEMENT**

Grateful acknowledgement is made to Assistant Professor T. Kondo, Kobe University, for his constant guidance in this investigation.

Thanks are tendered to Emeritus Professor C. Araki and Professor I. Honjo of Kyoto University for their interest and encouragement.

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GROWTH OF YOSHIDA SARCOMA IN THE THYMUS

REFERENCES