A COMPARISON OF EPIDEMIOLOGIC, HISTOLOGIC, AND VIROLOGIC STUDIES ON HODGKIN’S DISEASE IN WESTERN KENYA AND NAGASAKI, JAPAN

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Abstract. Rapid progress in molecular technologies has enabled the detection of several oncogenic viruses in various types of tumors. The pathogenesis of Hodgkin’s disease is suggested to have a strong association with Epstein-Barr virus (EBV). However, Hodgkin’s disease related to EBV shows a wide geographic variation in epidemiology. These variations among different populations suggest an interaction of environmental factors and a direct role of EBV infection. Therefore, we performed a comparative study on epidemiologic, histologic, and virologic features of Hodgkin’s disease among those in the western part of Kenya and in Nagasaki, Japan. The age distribution of Hodgkin’s disease showed a distinct peak in the 0–9-year-old age group in Kenya, and a higher and lower peak in the 60–69- and 30–39-year-old age groups, respectively, in Japan. The most common subtype of Hodgkin’s disease in both countries was mixed cellularity, followed by nodular sclerosis, lymphocyte depletion, and lymphocyte predominance. Mixed cellularity showed a significantly high prevalence among Kenyan children nine years of age or younger. Using the in situ hybridization method, EBV-encoded RNA (EBER-1) was detected in 79% of the Kenyan cases and 59% of the Japanese cases, with the mixed cellularity subtype showing a strong correlation with EBER-1. There was 100% positivity in both countries in those less than nine years old. These results suggest that EBV plays a more direct role in the pathogenesis of Hodgkin’s diseases in Kenya, especially in cases among young children and also in Japanese children. Environmental and/or genetic factors may have a role, in addition to EBV, in the pathogenesis of Hodgkin’s disease, especially in Nagasaki, Japan.

Epstein-Barr virus (EBV) infection is known to play an important role in oncogenesis of several neoplasias, in particular, African endemic Burkitt’s lymphoma and Chinese nasopharyngeal carcinoma in southeast Asia. Many reports also show that other diseases, such as Hodgkin’s disease, gastric cancer, and esophageal cancer, have some relationship with EBV infections. Hodgkin’s disease is a malignant lymphoproliferative disorder with characteristic features regarding epidemiology and histopathology. In industrialized countries, this disease shows a bimodal age distribution with peaks in young adults and in the elderly, with a predominance of nodular sclerosis. However, in developing countries, such as tropical Africa, central and south America, Hodgkin’s disease is seen more frequently in children, with a predominance of mixed cellularity and lymphocyte depletion subtypes.

The EBV genome has been detected in patients with Hodgkin’s disease by the polymerase chain reaction (PCR), Southern blot analysis, and in situ hybridization methods, and the rates of EBV detection vary depending on the geographic location under study or on the research methods used. The EBV positivity rate in patients with Hodgkin’s disease is reported to be much higher in developing countries, such as Peru, Honduras, and Kenya, than in industrialized countries, such as the United States and the European countries.

We have performed studies to compare the differences of epidemiologic and histologic patterns of Hodgkin’s disease and to investigate the presence of EBV in tissues of patients with this disease from the western part of Kenya and Nagasaki, Japan. We have chosen these areas for the following reasons. 1) The western part of Kenya, especially the basin area along Lake Victoria, is a highly predominant area of EBV-associated African endemic Burkitt’s lymphoma. 2) There are few reports of Hodgkin’s disease from sub-Saharan African countries, especially those correlating this disease with EBV. 3) Hodgkin’s disease is a relatively uncommon disease among the malignant lymphomas in Japan. 4) There are few reports of Hodgkin’s disease in association with EBV in Asian countries.

MATERIALS AND METHODS

We studied all consecutive 48 biopsy materials of cases of Hodgkin’s disease among the specimens that were submitted for pathologic diagnosis from various hospitals in the western part of Kenya (Western, Nyanza, and Rift Valley provinces; 1985–1995) to the Department of Histopathology, Rift Valley Provincial General Hospital and the 49 cases of Hodgkin’s disease recorded at the Cancer Registry of Nagasaki prefecture (1979–1991). This was a retrospective study and specimens used were archival specimens. This study was authorized by the Government of Kenya (research permit no. OP13/001/8C224/36) and was reviewed and approved by the Committee of Cancer Registry of Nagasaki Prefecture. In cases where more than one biopsy was taken from a patient, only the first sample was used. We divided the cases into three age groups: children (0–14 years old), young adults (15–44 years old), and adults (> 45 years old). The data on the age of 19 cases from Kenya were not available.

Histologic evaluation. Specimens were fixed in 10% formalin saline solution, embedded in paraffin, sectioned, and stained with hematoxylin and eosin, periodic acid–Schiff, silver impregnation, and Mallory’s stain. Hodgkin’s disease was diagnosed based on results with these histologic stains along with immunohistochemical staining using monoclonal antibodies CD15 and CD30. According to the Rye classification, all cases were classified into four histologic subtypes (lymphocyte predominance, mixed cellularity, nodular sclerosis, and lymphocyte depletion) by three pathologists (MK, KT, and HI).
In situ hybridization. This method, which uses a digoxigenin-labeled EBV-encoded RNA (EBER-1) 30-base oligonucleotide probe\textsuperscript{25-26} (kindly provided by Dr. M. Tokunaga, Kagoshima City Hospital, Kagoshima, Japan), was performed on specimens from 48 cases from Kenya and 46 cases from Nagasaki. Briefly, 5-\(\mu\)m sections of formalin-fixed, paraffin-embedded tissue were mounted on aminopropyl-triethoxy silane (APS)–coated slides, deparaffinized, rehydrated, predigested with pronase (protease, lot 45H00841; Sigma, St. Louis, MO), and hybridized overnight at 37\(^\circ\)C with 0.5 \(\mu\)g of digoxigenin-labeled probe. After washing with 0.5 \(\times\) SSC (0.075 M NaCl, 0.0075 M sodium citrate), hybridization was detected by an antidigoxigenin antibody–alkaline phosphatase conjugate (code 1093 274 150; Boehringer Mannheim, Mannheim, Germany), then counterstained with methyl green. Lymph node sections from a patient with infectious mononucleosis were used as a positive control and a sense probe for EBER-1 was used as a negative control for each procedure.

Immunohistochemistry. Immunohistochemical staining was performed on 48 cases from Kenya and 46 cases from Nagasaki. The following monoclonal antibodies were used: CD15 (C3D-1, lot 124; Dako, Glostrup, Denmark), CD30 (Ber-H2, lot 044; Dako), and EBV-encoded latent membrane protein (LMP-1) (CS1-4, lot 013; Dako).\textsuperscript{8,17,19-21,33,34} All sections were cut at a thickness of 5 \(\mu\)m and mounted on APS-coated slides. After dewaxing with xylene and rehydrating with alcohol, slides were placed in 0.01 M citrate buffer solution and pretreatment procedures to unmask the antigens were performed in a microwave oven for 10 min. Containers were cooled at room temperature for 20 min. Endogenous peroxidase activity was blocked with 3% hydrogen peroxide. The sections were then washed with xylene and dehydrating with alcohol, slides were placed in 0.01 M citrate buffer solution and pretreatment procedures to unmask the antigens were performed in a microwave oven for 10 min. Containers were cooled at room temperature for 20 min. Endogenous peroxidase activity was blocked with 3% hydrogen peroxide. The sections were then washed in phosphate-buffered saline (PBS) and processed for immunohistochemistry with a labeled streptavidin biotin-based method (LSAB kit, lot 0754; Dako). After incubation with a second blocking reagent for 5 min, primary antibody was applied and incubated at 4\(^\circ\)C overnight. Sections were washed three times in PBS (5 min/wash) and the secondary antibody was applied for 10 min. After the sections were rinsed three times in PBS (5 min/wash), streptavidin-peroxidase reagent was applied and incubated for 10 min. The sections were washed three times in PBS (5 min/wash) and reactions were developed with diaminobenzidine–activated with hydrogen peroxide. Sections were then washed with water, counterstained with hematoxylin, dehydrated, and mounted. Sections from a patient with infectious mononucleosis were used as a positive control and sections from a patients with reactive lymphadenitis were used as a negative control.

Statistical analysis. A statistical comparison was performed for EBER-1 positivity rates between the cases in Kenya and those in Nagasaki using the chi-square test. A \(P\) value of 0.05 was chosen as the significance level.

RESULTS

Age and sex. The mean age of the patients from Kenya was 21 years (range = 4–65) and that of the patients from Nagasaki was 52 years (range = 6–86). Hodgkin’s disease showed a bimodal age distribution in both countries, with the first distinct peak in the 0–9-year-old age group and a second lower peak in the 50–59-year-old age group in Kenya, and a higher peak in the 60–69-year-old age group and a lower peak in the 30–39-year-old age group in Nagasaki (Figures 1 and 2). The male to female ratio was almost 2:1 in both countries.

Histologic classification. The most common histologic subtype in both countries was mixed cellularity, followed by nodular sclerosis, lymphocyte depletion, and lymphocyte predominance (Tables 1 and 2). However, among the subtypes, the prevalence of mixed cellularity was significantly higher among Kenyan children nine years of age or younger. In both countries, the mean age of patients with nodular sclerosis was lower than that of any subtype of Hodgkin’s...
Hodgkin's disease in Kenya and Nagasaki, Japan

**Table 1**

Histologic classification of Hodgkin's disease patients in Kenya

<table>
<thead>
<tr>
<th>Histologic type</th>
<th>Sex</th>
<th>Total</th>
<th>Mean age (years)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Male</td>
<td>Female</td>
<td>No.</td>
</tr>
<tr>
<td>LP</td>
<td>2</td>
<td>1</td>
<td>3</td>
</tr>
<tr>
<td>MC</td>
<td>23</td>
<td>10</td>
<td>33</td>
</tr>
<tr>
<td>NS</td>
<td>4</td>
<td>3</td>
<td>7</td>
</tr>
<tr>
<td>LD</td>
<td>4</td>
<td>1</td>
<td>5</td>
</tr>
</tbody>
</table>

*L P = lymphocyte predominance; MC = mixed cellularity; NS = nodular sclerosis; LD = lymphocyte depletion.

**Table 2**

Histologic classification of Hodgkin's disease patients in Nagasaki

<table>
<thead>
<tr>
<th>Histologic type</th>
<th>Sex</th>
<th>Total</th>
<th>Mean age (years)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Male</td>
<td>Female</td>
<td>No.</td>
</tr>
<tr>
<td>LP</td>
<td>3</td>
<td>4</td>
<td>7</td>
</tr>
<tr>
<td>MC</td>
<td>16</td>
<td>4</td>
<td>20</td>
</tr>
<tr>
<td>NS</td>
<td>8</td>
<td>7</td>
<td>15</td>
</tr>
<tr>
<td>LD</td>
<td>4</td>
<td>3</td>
<td>7</td>
</tr>
</tbody>
</table>

*For definitions of abbreviations, see Table 1.

**Discussion**

Kenya is located in the lymphoma belt along the low latitudinal zone of the African continent. The western part of Kenya shows an especially high frequency of EBV-related endemic Burkitt's lymphoma. In our study, Hodgkin's disease in Kenya demonstrated the typical age distribution seen in other developing countries; a distinct peak in childhood, with a predominance of the mixed cellularity subtype, and the lower peak in adult groups. The series of patients with Hodgkin's disease from Nagasaki showed a higher peak in the adult group and a lower peak in the young adult group, which was relatively similar to data from other industrialized countries.

The expression rate of the EBV genome in Hodgkin and Reed-Sternberg (RS) cells of tissues from patients with Hodgkin's disease ranges from 26% to 100%, depending on geographic location or the method that researchers used, such as PCR and in situ hybridization methods or Southern blot analysis. Relatively higher positive rates have been reported from Central and South America and Africa, examples of which are 100% in Honduras, 94% in Peru, and 92% in Kenya. Lower rates (26–48%) have been reported from the United States and European
countries and intermediate rates (64–76%) from Asian countries. Among the several methods that detect EBV, the PCR is more sensitive than the others. However, one disadvantage of the PCR is that false-positive results can easily be obtained when the case harbors the virus in non-neoplastic lymphoid cells. One advantage of in situ hybridization is that it definitively localizes virus RNA in the neoplastic cells morphologically. Therefore, we used in situ hybridization in our study to detect the EBV genome in tissues from patients with Hodgkin’s disease. Strong signals were revealed in the nuclei of RS cells and their variants in 79% of the cases from Kenya and in 59% from Nagasaki.

As in many previous reports, our results showed a strong association between EBV and the mixed cellularity and lymphocyte depletion histologic subtypes. In the mixed cellularity subtype, EBER-1 positivity was 85% in Kenya and 67% in Nagasaki, while that for the lymphocyte depletion subtype was 80% in Kenya and 71% in Nagasaki.

Figure 3. A, Epstein-Barr virus-encoded RNA in situ hybridization showing strong signals in nuclei of Reed-Sternberg (RS) cells and their variants (in situ hybridization with methyl green, magnification × 400). B, immunohistochemical staining of latent membrane protein-1 (LMP-1) showing strong staining in the membrane and cytoplasm of RS cells and their variants (LMP-1 immunohistochemical staining, magnification × 400).
HODGKIN’S DISEASE IN KENYA AND NAGASAKI, JAPAN

TABLE 3

Epstein-Barr virus (EBV)–encoded RNA positivity of Hodgkin’s disease patients in Kenya

<table>
<thead>
<tr>
<th>Group†</th>
<th>No. of cases</th>
<th>Histologic subtype*</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td>LP</td>
</tr>
<tr>
<td>Children</td>
<td></td>
<td></td>
</tr>
<tr>
<td>(&lt;15)</td>
<td>18</td>
<td>1/1</td>
</tr>
<tr>
<td>(≥15)</td>
<td>12</td>
<td>0/0</td>
</tr>
<tr>
<td>Young adults (15–44)</td>
<td>8</td>
<td>0/1</td>
</tr>
<tr>
<td>Adults</td>
<td></td>
<td></td>
</tr>
<tr>
<td>(≥45)</td>
<td>6</td>
<td>0/0</td>
</tr>
<tr>
<td>EBV + (%)</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

* For definitions of abbreviations, see Table 1.
† Age group values are in years.
‡ Includes one age unknown group.

nagasaki. On the other hand, of the four subtypes, nodular sclerosis had the lowest rate of EBER-1 positivity: 57% in Kenya and 36% in Nagasaki.

Interestingly, our results showed higher EBER-1 positivity in children in both Kenya (89%) and Nagasaki (100%) compared with the other groups. Strikingly, EBER-1 positivity was 100% in those nine years of age and less in both countries. This phenomenon has already been reported by several investigators, and our results support these findings.18–23 The adult group (≥ 45 years old) had the second highest rate of EBER-1 positivity, with young adult groups having the lowest rate in both countries. In our study, certain histologic subtypes seemed to be predominant in specific age groups that showed EBER-1 positivity. Children and adults showed a predominance of mixed cellularity, while young adults showed a predominance of nodular sclerosis. Mixed cellularity had a strong association with EBV, while nodular sclerosis had the lowest correlation with EBV. This result is similar to those of many previous reports;3,8,10,13,14,20–22 however, Leoncini and others noted a high expression of EBER-1/2 in Kenyan cases with nodular sclerosis.6 Since these investigators used the REAL classification of Hodgkin’s disease subtyping,35 further study is needed to confirm these data.

The reason for the difference in these epidemiologic features has not yet been determined; however, it may be speculated that the environment of poor economic conditions and poor hygiene in which many African children live causes a secondary immunodeficiency status.15,28

Latent membrane protein-1 is known to have transforming activity and has also been reported to be an oncogenic protein that protect cells from programmed cell death.4,8,13,17,19,21,34 Although the precise role of LMP-1 is still obscure, strong staining of LMP-1 in the membrane and cytoplasm of Hodgkin and RS cells from tissues of patients with Hodgkin’s disease in our study indicates that EBV plays an important role in the pathogenesis of this disease with its oncogenic potential.4,5,7,9,11–14,16–23 We have also observed four cases that were EBER-1 positive and LMP-1 negative and eight cases that were EBER-1 negative and LMP-1 positive. The group of individuals who were EBER-1 positive and LMP-1 negative can be explained by the lower sensitivity of the immunohistochemical method compared with in situ hybridization.5,7 There is a possibility that the group of eight cases who were EBER-1 negative and LMP-1 positive harbor a type 2 strain in the neoplastic cells. This is consistent with the results of several investigators,6,7,10,12,18,23 who detected EBV type 2 strains in cases of Hodgkin’s disease.

The results of our study suggest that EBV plays a more direct role in the pathogenesis of Hodgkin’s disease in Kenya than in Nagasaki, and that this disease in children is more strongly linked with EBV infection than in adults. In addition to that of EBV, environmental and/or genetic factors also have a role in the pathogenesis of Hodgkin’s disease, especially in Nagasaki, Japan.

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TABLE 4

Epstein-Barr virus (EBV)–encoded RNA positivity of Hodgkin’s disease patients in Nagasaki

<table>
<thead>
<tr>
<th>Group†</th>
<th>No of cases</th>
<th>Histologic subtype*</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td>LP</td>
</tr>
<tr>
<td>Children</td>
<td></td>
<td></td>
</tr>
<tr>
<td>(&lt;15)</td>
<td>4</td>
<td>0/0</td>
</tr>
<tr>
<td>(≥15)</td>
<td>2</td>
<td>0/0</td>
</tr>
<tr>
<td>Young adult (15–44)</td>
<td>13</td>
<td>2/2</td>
</tr>
<tr>
<td>Adult</td>
<td>28</td>
<td>2/4</td>
</tr>
<tr>
<td>EBV + (%)</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

* For definitions of abbreviations, see Table 1.
† Age group values are in years.
‡ Includes one age unknown case.
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