CASE STUDY

Human herpes virus-8 associated primary effusion lymphoma of the pleural cavity in HIV-negative elderly men


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ABSTRACT: Human herpes virus-8 (HHV-8)-associated primary effusion lymphoma (PEL) is an unusual lymphoma confined to the body cavities, which primarily affects human immunodeficiency virus (HIV)-positive men at high risk for Kaposi’s sarcoma (KS). We describe two HIV-negative elderly Italian men, who developed pleural HHV-8-positive PEL in association with other diseases (systemic hypertension, colonic carcinoma, chronic obstructive airways disease, dilated cardiomyopathy), but without KS. Thoracic computed tomography revealed unilateral pleural effusion and pleural thickening. Thoracentesis disclosed large lymphoma cells, with no T- or B-cell associated antigens, clonal rearrangement of the immunoglobulin heavy chain gene and the presence of HHV-8 but not Epstein-Barr virus deoxyribonucleic acid sequences.

Primary effusion lymphoma (PEL) is a rare non-Hodgkin’s lymphoma of B-cell origin, in which tumour cells contain deoxyribonucleic acid (DNA) sequences of a new herpes virus known as Kaposi’s sarcoma (KS)-associated herpes virus or human herpes virus-8 (HHV-8) [1]. The disease has been predominantly described in homosexual and bisexual men with advanced human immunodeficiency virus (HIV) infection, but rarely in HIV-negative individuals, in Europe and in the USA [1–4]. Peculiar features of PEL include: 1) a propensity to proliferate in the fluids of body cavities with infiltration of the serosal linings, usually in the absence of tumour mass(es) and/or lymph node involvement; 2) large-cell morphology with null immunophenotype; and 3) a B-cell genotype. Recent studies indicate that the normal counterpart of PEL tumour cells is the mature B-cell, of preplasma cell type. Recent studies have shown that HHV-8 infection is not ubiquitous in the peripheral blood mononuclear cells of healthy blood donors [7], serological studies have shown that HHV-8 infection is not ubiquitous in most populations [8]. In Italy, however, HHV-8 seroprevalence and incidence of classic KS are higher, as compared to those recorded in the USA and other Northern European countries [8–10].

We describe two elderly Italian HIV-negative patients with HHV-8-positive pleural PEL, and review the few recent reports of non-AIDS-related HHV-8-positive PEL. Our study suggests a possible relationship between immunosenescence and the development of non-AIDS PEL. Its frequency is probably underestimated, at least in Italy, a geographical area at high risk for KS, and thus conceivably also for PEL.

Methods

Patients

Two elderly HIV-negative individuals developing non-Burkitt’s lymphomatous effusions in the absence of extranodal or node-based lymphoma were identified among 3,418 with serous effusions examined at the Dept of Pathology of the Policlinico Hospital (Rome, Italy) between January 1991 and December 1997.

Samples

Ailquets of pleural fluids were received following diagnostic thoracentesis, and cell pellets were examined using Papanicolaou and May-Grünewald Giemsa-stained cytopsinss. Cells recovered from patient 1 were cryopreserved in 10% dimethyl sulphoxide/20% foetal calf serum in minimal essential medium.

Immunophenotypic characterization

The immunophenotypic profile of the cell populations was determined by means of immunoperoxidase-staining of acetone-fixed cytopsins using a labelled streptavidin/biotin method and monoclonal antibodies directed against CD5, CD15, CD19, CD20, CD22, CD45, CD45R0, CD68, CD30 (Ki-1), epithelial membrane antigen (EMA), vimentin and cytokeratin (clone CKMNF116). All reagents were supplied by Dako (Glostrup, Denmark), except CD22 and anti-CD15 (Becton-Dickinson, San Jose, CA, USA). Flow cytometry, a more desirable means of determining immunophenotype in suspected lymphomas, was not performed because of the scarcity of the specimens.
Molecular studies

Genomic DNA was extracted from cryopreserved cell suspension in patient 1, and from alcohol-fixed cells of previously Papanicolaou-stained slides in patient 2, since only a limited amount of cells was obtained from the pleural fluid in this patient. Polymerase chain reaction (PCR) to detect HHV-8 DNA sequences was performed using the primers and conditions described elsewhere [11]. PCR experiments for HHV-8 detection included the coamplification of DNA extracted from a HHV-8-positive KS biopsy as positive control. In addition, in order to assess the integrity of the DNA, a 215-base pair (bp) fragment of the noncoding 5′ region of the Bcl-6 gene was coamplified using the primers 5′-AGGAAGGAGGGAATTAG-3′ and 5′-AACGATTCTTCAAGGCGAG-3′. PCR analysis of Epstein-Barr virus (EBV) DNA sequences was performed using primers representative of the Epstein-Barr virus-associated nuclear antigen-2 region [12]. In case 1, the genomic configurations of the immunoglobulin heavy chain (IgH), c-myc, Bcl-2 and Bcl-6 genes were investigated by means of Southern blot analysis using the probes and experimental conditions reported elsewhere [13, 14]. In case 2, the IgH gene configuration was analysed via PCR amplification using the following primers: FR3A (5′-ACACGCGGGTGATTATCGT-3′), LH (5′-TGAGAGCAGGCGGTGACC-3′), and VLJH (5′-GTGCCAGGGTACTTCTGGCCCCAG-3′).

Results

Report of two cases

Case 1. An 89-yr-old man, with a remote history of malaria (1936), systemic hypertension, surgery for lumbar intervertebral disk herniation (1984) and large bowel adenocarcinoma (1994), was admitted to the hospital in June 1995 because of dyspnoea. Laboratory data disclosed: red blood cell count 3.4×10^{6} cells·μL^{-1}, haemoglobin concentration 10 g·100 mL^{-1}, white blood cell count 7.9×10^{3} cells·μL^{-1} (CD4+ count not performed), erythrocyte sedimentation rate (ESR) 70 mm·h^{-1}, platelet count 191×10^{3} cells·μL^{-1}, iron concentration 18 μg·dL^{-1}, γ-globulin concentration 1.8 g·dL^{-1} (26%, normal range 9.0–18.0%), and serum lactate dehydrogenase (LDH) activity 239 IU·L^{-1} (normal range 9.0–18.0%).

CT scan showed contraction of the left hemithorax, thickening of the pleura and pleural effusion. Thoracentesis fluid showed malignant lymphoma (not shown). Pleural fluid LDH activity was 957 IU·L^{-1} (pleural fluid LDH/serum LDH= 5.9). The patient is currently under clinical surveillance without any therapy, and is alive at 12 months follow-up.

Both patients were Italian-born (birthplaces: Puglia region, Southern Italy (case 1); and Lazio region, Central Italy (case 2). Patient 1 had lived in Puglia (Foggia province) during his whole life, whereas patient 2 had lived in Sardinia for many years, and then in Lazio (Rome). They were both seronegative for HIV and hepatitis B and C virus. Cultures of pleural fluids for aerobic, anaerobic, mycobacterial and fungal organisms yielded negative results. There was no evidence of lympho-adenopathy or tissue-based lymphoma based on CT scan findings. Neither of them had a history of previous or concomitant KS.

Morphology and immunophenotype

Cytological examination in both cases showed a population of medium-to-large-sized lymphoid cells intermixed with small lymphocytes, macrophages, neutrophils, eosinophils and rare mesothelial cells (fig. 1). Tumour cells had centrally or peripherally located irregular nuclei, prominent nucleoli and abundant deep basophilic cytoplasm. Bi- and multinucleate cells and very large cells with multilobate nuclei were common. Nuclear fragmentation, apoptosis and numerous mitotic figures were additional findings. Tumour cells were CD45- or negative, B-cell-associated antigen (CD20, CD19, CD22)-, T-cell-associated antigen (CD5, CD45R0)-, cytokeratin-, CD68-, and CD15-negative and EMA (focal)- and CD30 (focal)-positive. In case 1, a few lymphoma cells were vimentin-positive. Small lymphocytes were mostly T-cells (CD5+/CD45R0+); other reactive cells were macrophages (CD68+) and mesothelial cells (cytokeratin-positive).

Molecular findings

The HHV-8 DNA sequences were detected in DNA extracts from lymphoma cells in both cases (fig. 2). Clonal rearrangement of the IgH gene was detected in both cases. Owing to the small amount of available tumour cell DNA,
case 2 was only analysable by means of the PCR. No variation from the germ line configuration of control DNA was found by analysing c-myc, Bcl-2 and Bcl-6 genes in case 1. The EBV genome was absent in both cases.

Discussion

According to the currently available literature, HHV-8-positive PELs occur rarely in HIV-negative individuals. To the best of the authors' knowledge, only six cases with a detailed description of clinicopathological features have been reported to date [1–4]. The main characteristics of our patients, and of the other described non-AIDS cases, are summarized in table 1.

Non-AIDS PELs share several features with the more common AIDS-associated PELs, such as large-cell morphology, null immunophenotype, clonal IgH gene rearrangements, the presence of KS and lack of c-myc rearrangements. Distinguishing features consist of an older age on presentation, apparent host immunocompetence, infrequent EBV coinfection and an apparently less aggressive clinical course.

In our cases, the lymphomatous nature of the pleural effusion was an unexpected finding. Patients had no evidence of node-based or extranodal lymphoma, and fluid accumulation occurred in the course of other diseases. Laboratory abnormalities, including anaemia, moderate iron deficiency, a high ESR, hypergammaglobulinaemia and elevated serum/plerural fluid LDH activities, were inconclusive. Relevant radiological findings included modest serosal thickening unaccompanied by pulmonary infiltrates and effusions. The diagnosis of PEL was initially suspected on the basis of the morphology (large cell) and immunophenotypic features of the pleural fluid population (CD45+, B-cell-associated antigen-negative-/T-cell-associated antigen-negative), and the absence of a nodal or extranodal lymphoma other than in the pleural cavity. A final diagnosis of HHV-8-associated PEL was established following molecular evidence of B-cell clonality and HHV-8 infection in tumour cells.

The risk of developing PEL is possibly linked to decreased immunological surveillance, as suggested by its high frequency in AIDS individuals [1]. In the apparently immunocompetent host, the mechanisms underlying PEL development are less clear. However, the occurrence of PEL in elderly individuals (see table 1 and the present cases) suggests that a certain degree of immunodepression and/or immunodysregulation may also be present in these cases. Ageing is notoriouly associated with low T-cell immunoresponsiveness and a normal/increased humoral response, probably resulting from an imbalance in cytokine production [15]. The frequent comcomitance of other diseases (second primary malignancies or cardiac failure/systemic hypertension) observed in our cases and in previously reported subjects with non-AIDS PEL [1], as well as in patients with classic KS [9], suggests an additional potential risk factor. Thus, underlying chronic disorders and age-related immunosenescence [15] represent the background against which non-AIDS PEL development may occur, possibly through HHV-8-mediated cytokine dysregulation in HHV-8-infected individuals [16].

It should be pointed out that our PELs showed epidemiological analogies with KS, another HHV-8-associated disease [17, 18]. The birthplace and area of residence of our patients (South/central Italy and Sardinia) are considered as risk indicators for KS [17]. Moreover, the history of malaria in case 1 is in keeping with the fact that in places in which KS is endemic, malaria is common [18]. Both HHV-8 seroprevalence and the incidence of classic KS (1 per 100,000 males)

Table 1. Human herpes virus-8-related primary effusion lymphomas in non-acquired immune deficiency syndrome patients

<table>
<thead>
<tr>
<th>First author [Ref.]</th>
<th>Age yrs</th>
<th>Sex</th>
<th>Origin</th>
<th>Body cavity</th>
<th>Kaposis sarcoma</th>
<th>Underlying conditions</th>
<th>EBV</th>
<th>Outcome</th>
</tr>
</thead>
<tbody>
<tr>
<td>CARBONE [2]</td>
<td>69</td>
<td>F</td>
<td>Italy</td>
<td>Peritoneal</td>
<td>No</td>
<td>Cirrhosis</td>
<td>+</td>
<td>Died, 1 month</td>
</tr>
<tr>
<td>NADOR [1]</td>
<td>85</td>
<td>M</td>
<td>USA</td>
<td>Pleural</td>
<td>No</td>
<td>Congestive heart failure</td>
<td>+</td>
<td>Died, 6 months</td>
</tr>
<tr>
<td>SAD [3]</td>
<td>78</td>
<td>M</td>
<td>USA</td>
<td>Pleural</td>
<td>No</td>
<td>NR</td>
<td>-</td>
<td>Died, 6 months</td>
</tr>
<tr>
<td>STRAUCHEN [4]</td>
<td>84</td>
<td>F</td>
<td>NR</td>
<td>Artificial</td>
<td>Yes</td>
<td>Hypertension, large-bowel carcinoma</td>
<td>-</td>
<td>Died, 4 yrs</td>
</tr>
<tr>
<td>Present cases</td>
<td>89</td>
<td>M</td>
<td>Italy</td>
<td>Pleural</td>
<td>No</td>
<td>Malaria, hypertension, large-bowel carcinoma</td>
<td>-</td>
<td>Lost to follow-up</td>
</tr>
<tr>
<td></td>
<td>75</td>
<td>M</td>
<td>Italy</td>
<td>Pleural</td>
<td>No</td>
<td>Hypertension, COAD, dilated cardiomyopathy</td>
<td>-</td>
<td>Alive, 12 months</td>
</tr>
</tbody>
</table>

*: related to the capsule of a silicone implant. EBV: Epstein-Barr virus; F: female; M: male; NR, not reported; COAD, chronic obstructive airways disease.
are higher in Italy than in other Western countries [8, 9], particularly in Southern Italy [10]. Thus, it is conceivable that a sizeable proportion of healthy Italian adults are latently infected with HHV-8, and that the frequency of non-AIDS PEL (probably currently underestimated) might approximate to that of classic KS. Individuals of Southern/Eastern European ancestry [18] might also be at high risk.

Since the diagnosis of primary effusion lymphoma is based on the detection of human herpes virus-8 deoxyribonucleic acid (its genetic hallmark) besides the demonstration of clonal immunoglobulin heavy chain gene rearrangements, the application of diagnostic molecular techniques to body cavity-based lymphomatous effusions is recommended. The development of reliable serological assays for human herpes virus-8 based lymphomatous effusions is recommended. The development of reliable serological assays for human herpes virus-8 antibodies will probably contribute to the identification of those subjects with past exposure to the virus, who may be at risk for human herpes virus-8 related diseases, including primary effusion lymphoma.

**Addendum**


**References**