The diagnosis of Epstein-Barr virus-associated polymorphic B cell lymphoma in immunocompromised patients: Review of methods

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P Beauparlant, C Alfieri, J-H Joncas. The diagnosis of Epstein-Barr virus-associated polymorphic B cell lymphoma in immunocompromised patients: Review of methods. Can J Infect Dis 1991;2(3):109-115. Polymorphic B cell lymphoma and diffuse B cell lymphoproliferation associated with Epstein-Barr virus infection is increasingly reported in immunodeficient patients. Accurate diagnosis of these pathologies is essential because the appropriate treatment regimens for the patients in question differ from those for patients with other lymphoproliferative diseases. Two complementary techniques are currently used in the diagnosis and characterization of Epstein-Barr virus-associated B cell lymphomas and diffuse B cell lymphoproliferation. Immunofluorescence allows specific detection of Epstein-Barr nuclear antigens in lymphomatous tissue. Molecular hybridization with the *Bam* H1-W and/or *Bam* H1-NJ probes confirms the presence of the Epstein-Barr virus genome in tumour cells. The *Bam* H1-NJ probe is also useful in determining the clonality of the tumour and the replication mode, episomal or linear, of the viral genome. The polymerase chain reaction method allows detection of the Epstein-Barr virus genome within 24 h in these tumours and is more sensitive.

Key Words: B cell lymphoma, Epstein-Barr virus, Immunofluorescence, Molecular hybridization, Polymerase chain reaction

Diagnostic du virus d'Epstein-Barr associé au lymphome cellulaire B polymorphe chez les patients immunodéprimés: Revue des méthodes

RESUME: On rapporte des cas de plus en plus nombreux de lymphomes cellulaires B indifférenciés et de lympho-prolifération diffuse des lymphocytes B associés au virus d'Epstein-Barr chez les patients immunodéprimés. Il est essentiel de poser un diagnostic exact, vu que le régime thérapeutique requis n'est pas le même que pour les autres maladies lymphoprolifératives. Deux techniques complémentaires sont présentement utilisées pour le diagnostic et la caractérisation des lymphomes cellulaires B et de la lympho-prolifération diffuse des lymphocytes B, qui sont associés au virus d'Epstein-Barr. L'immunofluorescence...
permet de déceler les antigènes nucléaires codés par le virus d'Epstein-Barr dans les tissus lymphomateux, tandis que l'hybridation moléculaire utilisant les sondes Bam-H1-W ou Bam H1-NJ confère la présence du génome du virus d'Epstein-Barr dans les cellules tumorales. La sonde Bam H1-NJ permet également de déterminer la clonalité de la tumeur et le mode de réplication, épisomique ou linéaire, du génome viral. L'amplification de l'ADN viral par la réaction en chaine à la polymérase permet la détection du génome du virus Epstein-Barr dans ces tumeurs en 24 h et cette méthode est plus sensible.

**EPSTEIN-BARR VIRUS. THE CAUSATIVE AGENT OF infectious mononucleosis (1), is a DNA tumour virus associated with Burkitt's lymphoma and nasopharyngeal carcinoma (2). More recently, polymorphic B cell lymphomas and diffuse B cell lymphoproliferations associated with Epstein-Barr virus infection have been reported in patients with congenital (3) and acquired (4-9) immunodeficiencies. These pathological conditions have been reported in transplant patients (4-7), in patients receiving chemotherapy for leukemia (8) and, more frequently, in patients suffering from the acquired immune deficiency syndrome (AIDS) (9). These tumours may be monoclonal or poly- clonal, and monoclonal tumours may be associated with chromosomal abnormalities (10). The term 'polymorphic' refers to the histological appearance of these tumours, which may be described as consisting of a mixture of undifferentiated B cell lymphoblasts, immunoblasts and fully differentiated plasma cells (11). Because this virus has the property of immortalizing B cells in vitro, it is thought that the same phenomenon might occur in vivo, leading to B cell lymphomatous proliferations in the absence of adequate immune function. An accurate diagnosis of these tumours is essential because the treatment regimens appropriate for these patients differ from those for patients with other lymphoproliferative diseases. It was reported that two leukemic patients who developed polymorphic B cell lymphomas recovered when immunosuppressive drugs were discontinued, and their leukemia has not recurred to date, after periods of four and six years, respectively (12).

In order to facilitate the diagnosis of Epstein-Barr virus-associated lymphomatous proliferations, the authors report two case studies in which the lymphomas were Epstein-Barr virus positive and monoclonal as assessed by immunofluorescence and molecular hybridization.

**MATERIALS AND METHODS**

**Immunofluorescence for detection of viral antigens in tissue:** Detection of the Epstein-Barr nuclear antigen on imprints and frozen sections of biopsy specimens was performed by anticomplement immunofluorescence with positive and negative control antisera (13). These reference sera contained no antinuclear antibodies as determined by anticomplement immunofluorescence on the Epstein-Barr virus negative Molt-4 cell line.

**Determination of tumour clonality by molecular hybridization:** Molecular hybridization with a specific Epstein-Barr virus probe (Bam H1-NJ) was used to assess whether the tumour was polyclonal or monoclonal (14). Briefly, the 7 kilobase-pair cloned Epstein-Barr virus restriction fragment, Bam H1-NJ, which was constructed by fusion of the outermost 3' and 5' Bam H1 fragments of the linear genome while deleting the terminal repeat segments, was labelled with phosphorus-32 by the random priming method (15) with modifications (16). Ten micrograms of tumour DNA was digested with Bam H1, electrophoresed and blotted onto nitrocelullose by the method of Southern (17). After drying for 2 h in an 80°C oven, blots were prehybridized for 3 h at 42°C in 50% formamide, 6x SSC, 0.2% polyvinylpyrrolidone, 0.2% ficoll, 50 mM sodium phosphate (pH 7.0) and 100 μg/mL of sheared, freshly denatured salmon sperm DNA. The radioactive probe (Epstein-Barr virus Bam H1-NJ) was then added to the prehybridization solution, and the blots were incubated in a 42°C waterbath overnight. After a series of washes (10 mins in 2x SSC at 20°C; 60 mins in 0.1x SSC plus 0.1% sodium dodecylsulphate at 60°C; 3 mins in 0.1x SSC at 20°C), blots were dried in a 60°C oven for 30 mins. Autoradiography was performed with Kodak XAR film placed between two Cronex screens.

**RESULTS**

The following case studies, used here to illustrate the authors' diagnostic approach, have been discussed in detail elsewhere (12), and will only be briefly described below.

**Case 1:** A girl suffering from congenital combined immune deficiency associated with adenosine deaminase deficiency died at the age of three years from a monoclonal (IgM and gamma-light chain positive) primary B cell lymphoma of the brain (Figure 1).

**Case 2:** A 4-year-old boy, on maintenance chemotherapy for acute lymphoblastic leukemia in remission, subsequently developed primary
Epstein-Barr virus infection and intestinal tumours. These tumours were polymorphic in appearance: one was positive for IgA-kappa and another for IgM-gamma, suggesting multi­clonality.

Immunofluorescence studies using a human Epstein-Barr nuclear antigen positive antiserum revealed Epstein-Barr nuclear antigens in the fresh tumour cells (imprints and frozen sections) from both cases (Figure 2). No fluorescence was observed with the negative control serum.

Molecular hybridization using the BamH1-NJ probe was performed on DNA extracted from control laboratory cell lines P3HR-1, B95-8, Raji and Molt-4, as well as from tumour biopsies obtained from cases 1 and 2 (Figure 3). Molt-4 is a T cell leukemia line which is negative for Epstein-Barr virus. The Raji cell line, which is nonproductive for virus replication, contains approximately 50 copies of episomal Epstein-Barr virus DNA per cell but no linear Epstein-Barr virus DNA (18). The virus-producing P3HR-1 and B95-8 cell lines contain linear Epstein-Barr virus genomes in addition to episomal copies (19–21). Hybridization results reveal the presence of one high molecular weight band for both Raji and the tumour biopsies (Figure 3). Thus, one may deduce that the BamH1-NJ probe identified the large fused terminal regions of

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**Figure 1** Computed tomography scan of a primary B cell lymphoma of the brain (case 1)

**Figure 2** Anticomplement immunofluorescence with positive (left) and negative (right) control antisera on fresh tumour cells from case 1
the Epstein-Barr virus episome consisting of the
Bam H1-N fragment plus the Bam H1-J fragment
linked by the presence of a clonally defined num­
ber of terminal repeat segments (each consisting
of 500 basepairs) (14). These findings confirmed
the monoclonality of the tumours.

Molecular hybridization on B95-8 and P3HR-1
tumour DNA demonstrated multiple DNA fragments
of molecular weights ranging from 4.4 to 9.4
kilobasepairs (Figure 3). The high molecular
weight fragment corresponds to the fused region
of the Epstein-Barr virus episome, as described
above. Fragments of low molecular weight cor­
respond to the individual unfused Bam H1-N and
Bam H1-J segments of the linear genome, each
attached to one or more terminal repeat units (14).

**DISCUSSION**

Epstein-Barr virus infects and transforms B
lymphocytes both in vitro and in vivo. The result­
ing polyclonal lymphoblastoid cell lines generally
carry multiple episomal (circular) copies of the
genome and express only latent Epstein-Barr

**Figure 3** Left Molecular hybridization using Bam H1-NJ probe performed on DNA extracted from the laboratory cell
line B95-8 (1), Raji (2), Molt-4 (3) and P3HR1 (4). Right Molecular hybridization using the Bam H1-NJ probe performed
on DNA extracted from tumour biopsies obtained from cases 1 (1) and 2 (2). MW Molecular weight in kilobasepairs

virus antigens. In the occasional cell, Epstein­
Barr virus is spontaneously reactivated leading to
lytic antigen expression. In cases in which lytic
antigen expression culminates in the production
of virions, the presence of linear genomes can be
demonstrated (22). In vitro studies have shown
that acyclovir is effective only in reducing the
number of linear replicative Epstein-Barr virus
genomes (19,23). Acyclovir, when used therapeu­
tically in the treatment of polymorphic B cell
lymphoma associated with Epstein-Barr virus, is
believed to prevent viral replication and thus to
block the spread of infection to other B lympho­
cytes (24), thereby limiting extension of the
lymphoproliferative process. Acyclovir has been
reported to influence the outcome of polyclonal,
but not monoclonal, tumours (11). However, the
major factor influencing the outcome of this
tumour appears to be the relief of an immunosup­
pressive state (25).

A monoclonal tumour is one that originates
from a single infected cell, whereas a polyclonal
tumour evolves from multiple infected cells. Mul­
Polyclonal tumours are clonally distinct tumours, all of which originate from a single infected cell. Epstein-Barr virus replication is associated with the linearization of the viral genome at the terminal repeat segment. Virions differ in the number of terminal repeat segments contained in their genomes (14). Thus, if each clone of cells results from the proliferation of one cell infected by one virus particle, then the number of terminal repeat segments in all of the episomes in the clone will be identical, yielding only one upper band in a Southern blot of the DNA when probed with Bam H1-NJ. This upper band is composed of the terminal Bam H1-N and Bam H1-J fragments joined, in a given episome, by means of a determined number of terminal repeat segments. A polyclonal population will, on the other hand, give rise to multiple upper bands, because the number of terminal repeat elements varies among the clonal populations comprising the polyclonal tumour. This occurs as a result of the mode of replication of the virus (26). Based on this information, one may conclude that the Raji cell line is monoclonal, as well as the tumours in cases 1 and 2 in which no virus replication can be demonstrated by Southern blot analysis. Because of virus replication in the B95-8 and P3HR-1 cell lines, these appear polyclonal by Southern blot although they are monoclonal by immunoglobulin gene rearrangement; in these lines the mode of replication of the virus by the rolling circle model gives rise to additional longer fragments which have been shown to disappear following acyclovir inhibition of viral replication (26). Therefore, in the absence of viral replication within a tumour, as judged by the absence of short Bam H1-N and Bam H1-J bands, the presence of one longer Bam H1-NJ band confirms the monoclonality of the tumour. The significance of clonality in this type of tumour is presently unclear (12).

Another point to be highlighted is the capacity of this hybridization technique to reveal whether Epstein-Barr virus is replicating in tumour cells. Because acyclovir acts only on the replicative form (19,23), the degree of replication is believed to correlate with the expectation of a clinical response to acyclovir treatment (24). The genome of replicating Epstein-Barr virus is linear, implying that the Bam H1-N and -J fragments are separated from each other. Thus, the molecular hybridization results should reveal two or more fragments of low molecular weight (Figure 4). Molecular hybridization of the three laboratory cell lines indicated that P3HR-1 and B95-8, but not Raji, harboured replicative linear forms of Epstein-Barr virus. The tumour biopsies from the present two patients were like Raji in that they only contained latent episomal copies of the Epstein-Barr virus genome. However, a percentage of these tumours has been shown to contain replicative copies of the Epstein-Barr virus genome (27), theoretically justifying the use of acyclovir in such cases to prevent infection of additional cells.

Other Epstein-Barr virus restriction fragments may also be used as probes for the presence of the Epstein-Barr virus genome in lymphoma tissue. The Epstein-Barr virus Bam H1-W segment (termed Bam H1-V in some publications) is perhaps the best choice where sensitivity is concerned, because it is repeated several times within the IR1 region of the genome (28); however, it gives no information concerning the state of the Epstein-Barr virus genome (linear or episomal) or the clonality of the tumour.

In cases in which it is only necessary to determine the presence (or absence) of Epstein-Barr virus DNA in tissue specimens, dot blot hybridization is a fast and simple alternative (12). Whole cells or DNA extracted from these cells may be spotted directly onto nitrocellulose filters, which are then hybridized with the probe of choice.

Recently, at the National Centre for Epstein-Barr Virus at the Sainte-Justine Hospital Pediatric Research Centre, the polymerase chain

![Figure 4](image-url)
reaction method of DNA amplification allowed detection of the Epstein-Barr virus genome within 24 h by ethidium bromide staining of an amplified 110 basepair viral DNA segment in the same number of tumours that were found to be viral DNA positive by conventional hybridization (12). These results were subsequently confirmed within three days using a phosphorus-32-labelled oligoprobes spanning the amplified segment located within the Bam H1-W fragment. An additional tumour sample which was doubtful by the conventional method was definitely positive by the labelled oligoprobe confirmatory test. All other tumour samples which were negative by the conventional method also gave negative results by polymerase chain reaction (29).

The mechanism of B cell transformation induced by Epstein-Barr virus has yet to be determined. However, it is presently believed that three Epstein-Barr virus proteins are involved in this process. Two are nuclear proteins, namely Epstein-Barr nuclear antigens 1 and 2 (30-33), and the other, referred to as latent membrane protein 1 (34,35), is associated with the cytoplasm and plasma membrane. The authors have previously reported the presence of Epstein-Barr nuclear antigen in Epstein-Barr virus-associated polymorphic B cell lymphoma (12), and emphasize the diagnostic value of the sensitive Epstein-Barr nuclear antigen anticomplement immunofluorescence technique for detecting the presence of Epstein-Barr virus antigens in the tumour cells. Other laboratories have used immunofluorescence with monoclonal antibodies to Epstein-Barr nuclear antigen 2 and latent membrane protein 1 to demonstrate expression of these antigens in tumour tissue (36). Because immunofluorescence can be rather subjective and, therefore, understandably dependent on the experience of the observer, molecular hybridization is useful to confirm the diagnosis. This technique also allows detection of the Epstein-Barr virus genome in tumour tissue since the use of the restriction fragment Bam H1-NJ as a probe indicates tumour clonality. In addition, the presence of linear copies of the genome, which are associated with Epstein-Barr virus replication, may warrant the use of acyclovir to limit further replication of the virus and recurrent lymphoproliferation (24).

Polymorphic B cell lymphomas have been reported to occur in 1 to 3% of organ transplant patients (11,25,37). Since the majority of these lymphomas are associated with primary Epstein-Barr virus infection or reactivation, the authors have monitored Epstein-Barr virus infection in transplant patients at their institution. Primary Epstein-Barr virus infection in young children was frequent, reaching an incidence of over 50% (in a total of 29 cases) during the first six to 12 months following transplantation (unpublished data). These patients were followed between 1984 and 1990 with serial serological tests for Epstein-Barr virus IgG and IgM antibodies to viral capsid antigens and IgG antibodies to early antigens, in addition to virus isolation from saliva. The diagnosis of primary Epstein-Barr virus infection was confirmed by the appearance of Epstein-Barr viral capsid antigen IgM antibody and stable seroconversion after the disappearance of passively transferred antibodies from blood and/or other biological products. None of the children has yet developed lymphoma. These results may be explained by the small population sample, the length of follow-up, and the fact that immunosuppressive protocols were relatively mild compared to those used in other centres.

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