HUMAN IMMUNODEFICIENCY VIRUS-ASSOCIATED CHANGES IN GERMINAL CENTERS OF LYMPH NODES AND RELEVANCE TO IMPAIRED B-CELL FUNCTION

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The acquired immunodeficiency syndrome (AIDS) and AIDS-related conditions are characterized by a progressive loss of immunity. The spectrum of immunologic abnormalities is broad. One of the most prominent features of the disease is the functional depression and different degree of depletion of CD4+ lymphocytes. The causative agent, the human immunodeficiency virus type-1 (HIV-1), possesses a highly specific tropism for cells carrying the CD4 receptor (1-4). Thus, its main targets are lymphocytes and cells of the monocyte/macrophage lineage (5-10) including follicular dendritic cells (11-14) and Langerhans cells (15,16).

Because of the central role of CD4+ lymphocytes in the immune regulation, their loss and impaired function explain many immune disorders observed in HIV-1 infection. The CD4+ T-cell subset was originally designated as "helper" cells because they were found to be critical to in vitro activation, maturation of, and antibody secretion by B-cells. Paradoxically, HIV-1 infected individuals with diminished CD4+ lymphocytes show a variety of abnormalities indicating an in vivo B-cell activation (17-24). The patients have hypergammaglobulinemia; as peripheral B-cells secrete spontaneously increased levels of immunoglobulins. The mitogenic responses to T-dependent as well as T-independent mitogens are also impaired. In addition, they have an impaired ability to mount both primary and secondary humoral responses. Disturbance of CD4+ lymphocytes cannot account for these abnormalities alone because impaired B-cell function has been demonstrated even in asymptomatic individuals with preserved T-cell function and counts (23).

The mechanism(s) of this B-cell imbalance is not known, although, infection with Epstein-Barr virus (EBV) or cytomegalovirus (CMV) has been proposed. However, there are data suggesting that hypergammaglobulinemia can be independent of EBV infection or reactivation (24), and in HIV-infected persons, cells spontaneously secreting immunoglobulins are not EBV infected (25,26). Furthermore, in pediatric or transfusion-related HIV-infection, B-cell abnormalities have been observed even in the absence of EBV or CMV infection (27,28).

Recent studies indicate that crude viral preparations of HIV-1 or live HIV are potent B-cell activators in vitro (29-31). It is known that the most constant finding in HIV-1-induced persistent generalized lymphadenopathy (PGL) is an exuberant activation of the B-dependent zone of the lymph node (for review, see ref. 32). The histologic manifestation of this activation is a longstanding germinal center reaction with large irregularly shaped follicles. Therefore, the analysis
of germinal centers (GC) in PGL can offer valuable data bearing on the pathogenesis of the disease.

**Presence of HIV-1 in GC**

In 1984, Armstrong and Horne reported that GC from patients with PGL or AIDS-related Kaposi's sarcoma harbor retrovirus particles (33). Soon after, we confirmed this finding (34). The virions are located between the cell processes of follicular dendritic cells (FDC) (Figs. 1-4). This consistent finding has led to emphasis on the diagnostic value of electron microscopic examination of lymph nodes from patients with PGL or AIDS (32,35,36). Virions can be seen in GC early in the course of infection. We examined lymph nodes obtained 2 to 6 weeks after seroconversion. At this time, GC already contained a few virions and no ultrastructural changes of FDC could be detected (Fig. 2). In contrast, GC in
lymph nodes with longstanding lymphadenopathy harbor many HIV. Hyperplasia (Fig. 3) and different degrees of degeneration of FDC (Fig. 4 and 5) accompany virus accumulation.

The intrafollicular accumulation of HIV-1 during the course of disease is probably due to continuous virus trapping and virus production inside the GC. Germinal centers contain a considerable
number of cells carrying the CD4 receptor. There are CD4+ T-cells, the average percentage of which can be as high as 25% of the total cell population of GC. Macrophages and FDC also express the CD4 molecule at a low density (37). Therefore, it is not surprising that budding profiles on FDC (11,12,14,38), lymphocytes and macrophages (14) have been demonstrated in the GC.

Retrovirus particles persist in GC of infected individuals for long periods of time. Examining repeated biopsy specimens, we detected HIV-1 ultrastructurally both in the first and the second lymph nodes removed at up to 25 month intervals (14,39).

Presence of structural proteins of HIV-1 in GC

Further evidence for infection of GC can be achieved by immunostaining of frozen sections with monoclonal antibodies against gag proteins, p18, p24 (13,32,40-42) or with anti-gp160 of HIV-1 (39). Each of these viral proteins can be visualized in GC. The reaction is highly sensitive and can be used for diagnostic purposes (13,32,40,42). In our material, delicate aggregates of gag proteins were seen as early as 2 weeks after seroconversion (Fig. 6). In longstanding lymphadenopathy the amount of structural proteins of HIV-1 is high and it can be detected as long as GC or remnants thereof are present (40,42). Double label studies demonstrate a close association of viral proteins with FDC (13,40,42). The FDC exhibit different degrees of destruction, and deposits of viral proteins can regularly be seen in the neighborhood of FDC destruction.

The mechanism responsible for the destruction of FDC in HIV-1 infection is probably complex. Virus replication in FDC can result in cytolysis. Cytotoxic T-cells could also have a role in this process. In PGL, CD8+ lymphocytes are often arranged in groups corresponding to the FDC destruction (43). When double immunolabeling is applied, cells reacting with gag proteins can be seen in the centers of these clusters of CD8+ T-cells (42).

Presence of cells expressing HIV-1 RNA in GC

A highly sensitive 35S-labeled RNA probe of HIV-1 has been developed by Harper et al. (44,45). Results of in situ hybridization with this probe revealed that the majority of cells expressing viral RNA were located in the GC (39,46). Performing in situ hybridization on serial
sections from lymph nodes, we noted that labeled cells often represented infected foci as judged by positivity in the same area of GC on subsequent cutting levels. In addition to hot spots, a relative high grain density diffusely distributed over the GC could be seen (Fig. 7). Because the background grain count over the mantle zone or lymphoid parenchyma was low and electron microscopically many free virions could be found, we assumed that the relatively high diffuse grain density is due or partly due to these virus particles.

**CONCLUSIONS**

The data presented indicate that the GC of lymph nodes, and probably those of other organized lymphoid tissues, are infected by HIV-1 and represent an important virus reservoir where the causative agent can persist indefinitely. It is reasonable to assume that FDC play a central role in this process. These antigen trapping, presenting and retaining cells in performing their physiologic function capture HIV-1 early in the course of disease. Deposited on FDC, the virus

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*Fig. 6. Delicate deposits of p24 of HIV-1 (arrow) in a germinal center 2 weeks after seroconversion. Alkaline phosphatase anti-alkaline phosphatase immunoreaction. Original magnification: x100.*

*Fig. 7. Demonstration of cell with RNA of HIV-1 (arrow) in a germinal center by in situ hybridization. Note the different diffuse grain density in germinal center (GC) and mantle zone (MZ). Original magnification: x400.*
finds a sufficient number of permissive cells to infect and destroy. As the disease progresses, virus accumulation and continuous disintegration of GC occur.

HIV-1 and viral proteins held on the antigen presenting cells could act immunologically in a similar fashion as in vitro and be responsible for the B-cell activation and its morphologic manifestation, the exuberant follicular hyperplasia in PGL. Continuous loss of FDC could contribute to the diminished ability of HIV-1 infected patients to develop specific humoral immunity to neoantigens.

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