Immunosuppression Induced by HIV Infection

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Abstract

Immunosuppression due to HIV infection is a resultant of several factors including defective innate signaling pathways, increased viral replication and virus load, gradual loss of peripheral CD4+ T cells and depletion of T lymphocytes at mucosal sites that collectively lead to progressive immune deficiency and AIDS development. T cells derived from HIV patients were anergic and failed to mount a robust adaptive immune response. The host, on the other hand, becomes vulnerable to opportunistic infections such as tuberculosis and AIDS-related cancer such as Kaposi sarcoma, non-Hodgkin lymphoma, and cervical cancer with remarkable level of immunosuppression. Elucidation of the various mechanisms involved in these immunologic perturbations is necessary to understand HIV pathogenesis for an effective immunotherapy.

Keywords: HIV pathogenesis; Immune activation

Background and Significance

Infection with human immunodeficiency virus (HIV) has been identified as a causal agent of AIDS, a pandemic disease that constitutes a global public health threat for the past 3 decades [1]. It is well established that HIV preferentially infects lymphocytes with consequent depletion of CD4+ lymphocytes at mucosal sites [2,3]. The progressive death of T lymphocytes is attributed to immune activation rather than to viral cytopathic effect. The resulting immune dysfunction in HIV patients is likely due to combinatorial effects resulting from infection of immune cells (CD4+ T cells, macrophages, dendritic cells) with HIV, uncontrolled viral replication that impairs antigen presentation, increased mutations in env protein gp120 that leads to virus tropism and survival, increased activation of T helper cells by alloantigens, increased apoptosis by activated CD4+ T helper cells, down-regulation of CD4+ synthesis with functional impairment, and perturbation of cytokine pathways [4-6]. The sequel of these effects renders the patients immunosuppressed and vulnerable to opportunistic infections with consequent death. Among the common opportunistic disorders encountered in HIV patients, tuberculosis and AIDS-related cancer such as Kaposi sarcoma, non-Hodgkin lymphoma and cervical cancer.

However, these immunologic defects can be partially restored in HIV patients treated with highly active antiretroviral therapy (HAART) [7]. Patients who accessed HAART treatment and adhered to its regimen had shown significant reduction in AIDS-related mortality and high prevalence of comorbidities due to cardiovascular disease, diabetes, and non-AIDS cancers [8]. Thus, HIV is a relentlessly progressive disease and difficult to cure with HAART due to the persistence of HIV-1 virus in latent reservoirs that may lead to viral remission and development of AIDS [9]. The progression of HIV disease in HAART-treated patients can be virtually monitored by evaluation of inflammatory markers (e.g. kynurenines, Kyn) and Th17/Treg balance in HIV progressors [10, 11]. Despite the functional improvement in HAART-treated patients, as evidenced by reduced viral load and improved CD4+ T cells count, a paradoxical response known as immune reconstitution inflammatory syndrome (IRIS) has evolved [12]. The induction of IRIS is worsened in HIV patients with preexisting opportunistic infections. In general, the severity of IRIS depends on CD4+ T cells count (< 100-200 cells/μl), degree of lymphocyte apoptosis or proliferation, and the degree of viral suppression and immune recovery after the initiation of treatment with HAART.

Furthermore, the status of immunosuppression is complicated by increased HIV mutations and genetic diversity during virus replication, ability of the virus to establish a pool of latently infected CD4+ T cells during early stage of infection, failure of neutralizing antibodies to eliminate HIV replication, and failure of viral antigens to induce an effective T cell memory [13-18]. All of these factors lead the virus to evade the immune system with consequent failure of immune control. Therefore, development of an effective HIV vaccine remains elusive [13] unless obstacles pertaining to the virus and the host are defined for successful discovery of vaccine targets. In the era of omics technologies [19], application of high-throughput methods for vaccine discovery is promising to resolve the above hurdles and might be helpful for better understanding of the immune response to HIV infection with consequent development of an effective vaccine for which conventional methods had failed. In light of these challenges, current efforts for vaccine discovery against HIV are focused on the identification of immunogens that elicit broadly neutralizing antibodies coupled with the induction of HIV-specific cytotoxic T lymphocytes [13].

Conclusion:

In conclusion, immunosuppression elicited by HIV infection is different from other microbial infections [20], and solely complicated to be unidirectionally manipulated. Elucidation of the mechanistic events involved in virus replication coupled with the incurred perturbations in the host innate and adaptive immune signaling pathways are of paramount importance to define HIV pathogenesis and immunity.
References


