Upregulation of IL-21 Receptor on B Cells and IL-21 Secretion Distinguishes Novel 2009 H1N1 Vaccine Responders from Nonresponders among HIV-Infected Persons on Combination Antiretroviral Therapy

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Upregulation of IL-21 Receptor on B Cells and IL-21 Secretion Distinguishes Novel 2009 H1N1 Vaccine Responders from Nonresponders among HIV-Infected Persons on Combination Antiretroviral Therapy

Suresh Pallikkuth,* Sudheesh Pilakka Kanthikeel,* Sandra Y. Silva,* Margaret Fischl,† Rajendra Pahwa,‡ and Savita Pahwa*

Mechanisms underlying failure of novel 2009 H1N1 influenza vaccine-induced Ab responses in HIV-infected persons are poorly understood. This study prospectively evaluated 16 HIV-infected patients on combination antiretroviral therapy and eight healthy controls (HC) who received a single 15 μg dose of nonadjuvanted novel 2009 H1N1 influenza vaccine during the 2009 H1N1 epidemic. Peripheral blood was collected at baseline (T0) and at 7 d (T1) and 28 d (T2) postvaccination for evaluation of immune responses. Prevaccination hemagglutination inhibition Ab titer was <1:20 in all except one study participant. At T2, all HC and 8 out of 16 patients (50%) developed a vaccine-induced Ab titer of ≥1:40. Vaccine responder (R) and vaccine nonresponder patients were comparable at T0 in age, CD4 counts, virus load, and B cell immunophenotypic characteristics. At T2, HC and R patients developed an expansion of phenotypic and functional memory B cells and ex vivo H1N1-stimulated IgG Ab-secreting cells in an ELISPOT assay. The memory B cell response was preceded by a significant expansion of plasmablasts and spontaneous H1N1-specific Ab-secreting cells at T1. At T2, HC and R patients also exhibited significant increases in serum IL-21 levels and in the frequency and mean fluorescence intensity of IL-21R–expressing B cells, which correlated with serum H1N1 Ab titers. Vaccine nonresponder patients failed to develop the above-described vaccine-induced immunologic responses. The novel association of novel 2009 H1N1 vaccine-induced Ab responses with IL-21/IL-21R upregulation and with development of memory B cells and plasmablasts has implications for future research in vaccine design. The Journal of Immunology, 2011, 186: 000–000.

Vaccination is the most effective way to reduce the morbidity and mortality of influenza. In June of 2009, the World Health Organization declared a pandemic of a novel influenza A “novel 2009 H1N1” virus of swine origin, which resulted in an estimated 57 million cases in the United States from April 2009 to January 2010 (1). The 2009 influenza pandemic was considered to be especially dangerous because the influenza A novel 2009 H1N1 virus represented a novel strain caused by antigenic drift, unlike the yearly changes in the seasonal influenza virus resulting from antigenic drift. As this H1N1 strain was last associated with the 1918 influenza pandemic, people born after 1930 (i.e., in the age group <65 y) were most likely to lack pre-existing immunity (2–6) and to be more susceptible to the virus than people older than 65 y. The Centers for Disease Control recommended priority novel 2009 H1N1 influenza virus vaccination for selected populations that were considered to be at greatest risk of infection, including those with immunocompromised immune systems such as HIV-infected people (7).

Humoral immunity against influenza viruses is a good predictor of protection against infection (8). For the novel 2009 H1N1 influenza vaccine, an Ab titer of >1:40 hemagglutination inhibition units or a 4-fold increase from baseline is considered as being protective (9). Following the recommended single 15 μg dose of nonadjuvanted novel 2009 H1N1 vaccine, seroconversion with development of protective Ab titers has been reported to occur in 95% of the general population (6). Surprisingly, among HIV-infected individuals without pre-existing immunity to H1N1 virus, only 60% developed protective Ab titers following vaccination, even though they were well controlled for their HIV status (10). In this context, although greater expansion of memory B cells has been noted with adjuvanted H1N1 vaccines, their success in improving seroconversion rates has been variable (11–13). The mechanism underlying the poor serologic response to novel 2009 H1N1 vaccines in HIV-infected patients is not well understood.

In the generation of a primary Ab response to a vaccine, naive (N) B cells activated by the vaccine Ag are required to undergo a process of maturation, differentiation, and proliferation to gen-

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The online version of this article contains supplemental material.

Abbreviations used in this article: AM, activated mature differentiated; ASC, Ab-secreting cell; cART, combination antiretroviral therapy; Cy7, cyanin-7; ET, early transitional; ETL, exhausted tissue-like; HC, healthy control; LT, late transitional; MFI, median fluorescence intensity; N, naive; NR, vaccine nonresponder; R, vaccine responder; RM, resting memory; SAC, Staphylococcus aureus Cowan strain; SPB, short-lived plasmablast; T0, baseline; T1, 7 d postvaccination; T2, 28 d postvaccination.
erate Ab-secreting cells (ASC), memory B cells (14), and long-lived plasma cells (15). This process is regulated by cooperative interactions of B cell-intrinsic and extrinsic factors that are dependent upon a variety of accessory ligand–receptor interactions (16–18). IL-21 is a pleiotropic cytokine secreted almost exclusively by CD4 T cells and is known to play a major role in development of B cells and to promote Ab production (19–23). N human B cells are efficiently induced to secrete Ig via IL-21 following cognate T–B interaction (24). Recently, expression of the IL-21R on B cells was shown to be critical for development of memory B cells (25). HIV-infected persons are known to exhibit phenotypic and functional B cell defects, which are only partially corrected with potent combination antiretroviral therapy (26–28). Memory B cell deficiency may persist in these patients, resulting in poor responses to influenza virus vaccination (29–31), but the underlying mechanisms have not been elucidated. There is also accumulating evidence for impaired Ag-specific IL-21 secretion by CD4 T cells in progressive HIV infection (32–34). Based on the above observations and the high failure rates of the novel 2009 H1N1 vaccine in HIV-infected people, we hypothesized that defects in the IL-21/IL-21R pathway could prevent a successful serologic response to the vaccine. In the current study, we examined the relationship of the cytokine IL-21 and of its receptor expression on B cells to novel 2009 H1N1 vaccine responses in HIV-infected patients and healthy controls (HC). We observed that the HIV-infected vaccine responder (R) patients and HC were clearly distinguishable from vaccine nonresponder (NR) patients by their cellular immune responses and upregulation of IL-21/IL-21R expression. Notably, the findings imply that the IL-21/IL-21R pathway is likely to be playing an important role in generating serologic responses to the novel 2009 H1N1 vaccine.

Materials and Methods

Patient characteristics and response to vaccination

During the 2009–2010 influenza season, HIV-infected persons in the special immunology clinic at University of Miami who agreed to be vaccinated against novel 2009 H1N1 influenza virus as standard of care were invited to participate in a prospective study to evaluate immune responses to the vaccine. A single i.m. dose (15 μg; 0.5 ml) of inactivated monovalent A/California/07/2009 H1N1 vaccine (Novartis Vaccines and Diagnostics) was given to vaccine recipients. Data were presented for 16 patients who were on potent combination antiretroviral therapy (cART), had a titer of 1:40. Postvaccination, H1N1 Ab titers of ≤1:40 on day 28 postvaccination. Eight HIV uninfected persons who were given the same vaccine served as HC. Characteristics of the study population are shown in Table I. Prevaccine H1N1 Ab titers were ≤1:20 in all study participants except one HC (identification number 027) who had a titer of 1:40. Postvaccination, H1N1 Ab titers of >1:40 on day 28 or a 4-fold increase from baseline were considered as protective (9). At 4 wk postvaccination, 8 of 16 (50%) patients developed Ab titers of >1:40 (median, 480; range, 80–2560) and were designated as H1N1 R patients. Eight patients had H1N1 Ab titers of ≤1:20 U at T2 and were designated as H1N1 vaccine nonresponders (NR). At T0, mean age and frequencies of CD20+ B cells were statistically equivalent among the patient groups and HC; CD4 and CD8 T cell counts and plasma HIV RNA were not different between the patient groups. All HC developed protective H1N1 Ab titers (median, 720; range, 80–20,480). The study was approved by the institutional review board of the University of Miami.

mAbs

CD3 amcyan, CD20 allopocytocyanin cyanin-7 (Cy7), CD20 allopocytocyanin, CD10 PE-Cy7, CD21 PE-Cy5, CD27 Verso, 450, CD27 Alexa Fluor 700, K67 FITC; and IL-21R allopocytocyanin were purchased from BD Biosciences (San Jose, CA). Isotype Abs used were IgG1–Aleya 700, IgG1–PE-Cy7, IgG1–PE-Cy5, and IgG1 allopocytocyanin.

Reagents

Reagents used were 1× PBS, penicillin (100 U/ml), and streptomycin (100 mg/ml) (Cellgro, Manassas, VA); RPMI 1640 and l-glutamine (200 mM) (Life Technologies, Grand Island, NY); heat-inactivated FBS (HyClone, Logan, UT); Ficol Paque (GE Healthcare Biosciences, Uppsala, Sweden); BSA, DMSO, and Staphylococcus aureus Cowan strain (SAC) (Sigma-Aldrich, St. Louis, MO); Cytofix/Cytoperm, Perm Wash buffer, and 3-aminio-9-ethylcarbazole (BD Biosciences, San Jose, CA); goat anti-human IgG, HRP-conjugated streptavidin, and biotinylated goat anti-human IgG; Fcy fragment specific (Jackson ImmunoResearch Laboratories, West Grove, PA); multiscreen HA 96-well plate for ELISPOT (Millipore, Billerica, MA); violet amine reactive viability dye (Molecular Probes, Invitrogen, Eugene, OR); paraformaldehyde (Electron Microscopy Sciences, Hatfield, PA); cell-culture plates (Costar, Corning, NY); recombinant human IL-21 (R&D Systems, Minneapolis, MN); and IL-21 ELISA kit (eBioscience, San Diego, CA).

Processing of blood samples

Peripheral venous blood, 30 ml, was collected immediately before vacci-

nation (T0) and on days 7 (T1) and 28 (T2) postvaccination in heparinized vacutainer tubes after obtaining written informed consent from the participants. PBMC were isolated by density gradient isolation (35) using Ficoll Paque and processed fresh for immunohistotyping; remaining PBMC were cryopreserved in 10% DMSO and 90% FCS in liquid nitrogen. Serum samples were stored at −80°C. Hemagglutination inhibition Ab titers against H1N1 were performed at Bioqual (Rockville, MD) in T0 and T2 serum samples using virus and turkey RBC (10).

B cell characterization by flow cytometry

Total and maturational subsets of B cells were determined in HC and patients based on phenotypic characteristics described by Moir and Fauci (reviewed in Ref. 26). Freshly isolated PBMC (1 × 10⁶ cells from blood samples at T0, T1, and T2 were stained with Abs against surface Ags using CD3, CD20, CD10, CD21, CD27, or isotype control Abs. Intracellular staining with Ki67 was performed in fixed permeabilized cells for evaluating cell proliferation. Samples were acquired on an LSRII flow cytometer (BD Biosciences) with collection of 200,000–500,000 events and were analyzed using FlowJo software (version 8.8.6; Tree Star).

Flow cytometry analysis was performed after proper instrument setting, calibration, and compensation (36, 37). The gating scheme for identification of the B cell maturation subsets is shown in Fig. 1. Appropriate isotype control Abs were used at each step in gating of different B cell subsets to minimize effects due to nonspecific staining. The live cells (violet amine reactive viability dye negative) were first gated for CD3 CD20+ and CD3 CD20− subsets and sequentially on CD21, CD27, and CD20 to identify the B cell maturation subsets. The CD3+ CD20+ B cells were gated into CD21hi and CD21lo subsets, each subset divided into CD27− and CD27+ subpopulation, and further characterized on the basis of CD10 expression. The CD3+ CD20+CD10+ B cells were designated as resting memory (RM; CD27+CD10+), late transitional (LT; CD27+CD10−), and N (CD27+CD10−) subsets. The CD3+ CD20+CD10+ B cells were considered as activated mature differentiated (AM; CD27+CD10+), early transitional (ET; CD27+CD10+), and exhausted tissue-like (ETL; CD27+CD10−) subsets. The CD3+ CD20− subset was divided into CD21+ or CD21− B cells based on isotype Ab staining. The CD21hi cells were gated for CD27+CD10− cells and further characterized based on expression of Ki67 as short-lived plasmablasts (SPB; Ki67+) as described (26, 38).

Frequencies of B cell subpopulations were calculated as percentages of the total B cells identified in this study as CD3+ CD20+ plus CD3− CD20− lymphocytes. The various B cell populations were further investigated for frequency and median fluorescence intensity (MFI) for IL-21R based on isotype Ab staining (39).

ELISPOT assays for memory and effector H1N1 IgG ASC

Cryopreserved PBMC from different time points were thawed, restimulated overnight, and analyzed for ASC by ELISPOT using the method described by Kyu et al. (40). For analysis of memory B cells, 1 × 10⁶ PBMC/ml from T0 and T2 were cultured for 5 d at 37°C in the presence of 5 μg/ml H1N1 vaccine Ag. SAC, 1/10,000 dilution, was used as a positive control. In some experiments, exogenous cytokine IL-21, 50 ng/ml, was added to cultures together with H1N1 Ag at culture initiation. On day 5 of culture, cells were washed, and 100,000 cells/well were plated in triplicate for 5 h at 37°C in anti-IgG-coated ELISPOT plates. Thereafter, the plates were sequentially washed and incubated at room temperature for 1 h first with 100 μl/well biotinylated goat anti-human IgG (1:5,000 dilution) followed by 100 μl/well streptavidin-horseradish peroxidase (1:5,000 dilution). Plates were washed and developed using 3-amino-9-ethylcarbazole substrate (41, 42). Spots representing H1N1 Ag-stimulated IgG-secreting ASC were enumerated using an automated ELISPOT reader (Immunoscan version 2.08;
Cellular Technologies), and values in unstimulated wells were subtracted from those in test wells (41). IgG-secreting cells were expressed as ASC/10^6 B cells.

For analysis of effector cell responses, spontaneous H1N1-specific ASC (43) were determined in rested cryopreserved PBMC samples from T0, T1, and T2 without ex vivo Ag stimulation. Cells were washed and plated in triplicate for 6 h at 37˚C at 100,000 cells/well in H1N1 Ag-coated (5 µg/ml) ELISPOT plates. Plates were washed and processed as described above for determining frequency of ASC/10^6 B cells.

**ELISA assays for serum levels IL-21**

The concentrations of IL-21 were determined in serum samples using a commercial ELISA kit (eBioscience) as previously described (44). Plates were coated with 100 µl capture Ab (1:250 dilution) for 12 h at 4˚C, washed five times with wash buffer (PBS plus 0.05% Tween 20), and blocked with assay diluents (1:5 dilution) for 1 h. Thereafter, standards, 100 µl/well, and serum, 100 µl/well, were added in duplicate and plates left overnight at 4˚C. Plates were washed five times, and 100 µl/well detection Ab (biotin-labeled, 1:250 dilution in assay diluent) was added at room temperature for 1 h. Plates were washed again followed by addition of 100 µl enzyme conjugate (avidin-GRP, 1:250 dilution in assay diluent) at room temperature for 30 min and 100 µl substrate for 15 min. The reaction was stopped by adding 50 µl stop solution (2 N H2SO4), and plates were read at 450 nm. Enzyme activities were determined at an OD of 450 nm. Mean values of serum IL-21 were reported as picograms per milliliter.

### Table I. Characteristics of study population and Ab responses

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ID, identification number.

HIV-infected persons are known to manifest alterations in B cell maturation subsets, which are partially reversed when they are treated with potent cART (38, 45). Frequencies of B cell maturation subsets were determined in patients and HC in PBMC based on the gating scheme (Fig. 1) described in Materials and Methods and are depicted in Fig. 2 and Supplemental Fig. 1. A composite of all B cell maturation subsets at T0 and T2 is shown in Fig. 2A. Frequencies of B cell maturation subsets in patients differed from HC as has been previously described (38, 45), but there was no obvious difference between vaccine responder and nonresponder patients. At T0, frequencies of RM B cells were reduced in patients. At T2, however, the R and NR patients had widely divergent cellular responses, with expansions of RM and SPB in R patients but not in NR patients. A significant expansion of RM B cells from baseline was noted at T2, and this occurred in HC and R patients but not in NR patients. Expansion of the RM B cells was correlated with the H1N1Ab titer in patients (Fig. 2C) and also in HC at T2 (r = 0.53, p = 0.02; not shown). Responses of R patients resembled HC but did not always exhibit the same magnitude of response. Frequencies of AM B cells were not significantly different among
study groups at T0, and all showed expansions at T2 (Fig. 2D). A notable characteristic of B cells in influenza vaccinees is the rapid expansion of plasmablasts within 1 wk, and this has been primarily associated with the response of memory B cells (43, 46, 47). In the current study, frequencies of the SPB, which were equivalent at T0 among study groups, showed an increase of $3$-fold at T1 (day 7 postvaccination) in HC and in R patients but not in NR patients, followed by a downward trend toward baseline by T2 in all (Fig. 2E). The H1N1 Ab titers at T2 correlated with frequency SPB at T1 in patients ($r = 0.78$, $p = 0.004$; data not shown). As expected, LT B cells (Supplemental Fig. 1C) were decreased at T2 postvaccination in HC and R patients, and N B cells (Supplemental Fig. 1D) showed a decrease in HC, as both these subsets constitute developmental intermediates for generation of human mature B cells (48). Patients had greater frequencies of ET (Supplemental Fig. 1B) and ETL B cells (Supplemental Fig. 1G) than HC at T0, and these subsets did not change following vaccination. Except for SPB, no other B cell subset was altered at T1 in relation to T0 (not shown).

Serum IL-21 and frequency of IL-21R$^+$ B cells are selectively increased at T2 in vaccine responders

In support of the hypothesis that the IL-21R/IL-21 axis was involved in the development of Ab responses to the novel 2009 H1N1 vaccine, we noted that serum IL-21 levels increased from T0 to T2 in HC and R but not in NR patients (Fig. 3A). To further ascertain the role of IL-21R$^+$ B cells in HC and R patients with increases in MFI (156.9 ± 35.5 and 103.1 ± 32.5, respectively), but not in NR patients (MFI 36.9 ± 11.4). A direct correlation was observed between H1N1 Ab titer and IL21R$^+$ B cells at T2 (Fig. 3D). Among B cell subsets at T0 (Fig. 3E), frequencies of IL-21R$^+$ cells were highest in the LT subset and lowest in AM B cells and were significantly higher in HC in LT and RM subsets than both R and NR patients and also in N B cells of HC in comparison with NR patients. At T2, however, the IL-21R$^+$-expressing cells increased (Supplemental Fig. 2), and H1N1 Ab titer was correlated with IL-21R$^+$ LT, N, and memory B cells in patients as well as HC (not shown). No differences were noted in IL-21R$^+$ AM B cells among study groups, and no change occurred in frequencies of IL-21R$^+$ B cells or their subsets at T1.

H1N1 vaccine responders mount ASC responses at T1 and T2

Next, we aimed to determine if the serum Ab responses were correlated with functional responses of memory B cells and plasmablasts in ELISPOT assays for ASC in cryopreserved PBMC. As a measure of functional plasmablasts, spontaneous (unstimulated ex vivo) H1N1-specific ASC responses were detected at T1 (Fig. 4A), in concurrence with increase in phenotypic SPB and were noted to occur only in HC and R patients, with 1.5-fold higher ASC in HC. These responses were short-lived and were close to baseline at T2. An expansion of spontaneous H1N1-specific ASC at T1 did not occur in NR patients.

Functional memory B cells were analyzed by determining IgG ASC after ex vivo stimulation of PBMC with H1N1 Ag for 5 d. The H1N1 Ag-stimulated IgG ASC response occurred at T2 in R
patients and in HC but not in NR patients (Fig. 4B). The IgG ASC response at T2 in HC was greater than R patients. At T2, serum H1N1 Ab titers correlated with H1N1-stimulated IgG ASC in patients (Fig. 4C) and in HC ($r = 0.53$, $p = 0.02$, data not shown). ASC responses to SAC were equivalent at T0 in R and NR patients but significantly lower than in HC (Fig. 4D) and did not change from T0 at T1 or T2 (not shown).

**Discussion**

Influenza is a serious threat to the HIV-infected population and can cause significant morbidity. During the 2009 H1N1 influenza pandemic, the HIV-infected populations in our clinics were offered priority vaccination with a nonadjuvanted inactive novel 2009 H1N1 vaccine. The 2009 H1N1 vaccine Ag was novel for the age bracket of our study population, and a majority of recipients were seronegative for this vaccine at the time of vaccination. In agreement with published data (10), a large proportion of patients on cART failed to seroconvert despite good virologic control and CD4 reconstitution. This study was conducted to understand why 2009 H1N1 vaccination did not always engage the immune system efficiently in this patient population. We made the novel observation that R patients uniquely upregulated IL-21R on B cells and increased their serum IL-21 levels following vaccination. Notably, serum H1N1 Ab titers were correlated with vaccine-induced changes in phenotypic and functional memory B cells, plasma blasts, and IL-21R expression on B cells.

Typically, Ag-induced activation, proliferation, and differentiation of N B cells at the site of immunization and draining lymph nodes generates plasmablasts in the germinal centers followed by their migration to bone marrow, where they develop into plasma cells for secretion of serum Ab. Migration of B cells to sites such as the respiratory tract where they secrete mucosal Abs including IgA has been described for parenteral influenza vaccines (17, 49–51). Memory B cells are generated, poised to respond by activation and proliferation following re-exposure to the same Ag. Indeed, following H1N1 vaccination, we noted phenotypic and functional evidence for expansions of peripheral memory B cells at T2 in R patients and HC but not in NR patients. In the current study, all
patients were on cART, and the observed Ab or B cell responses did not correlate with CD4 T cell counts; thus, the deficiencies in memory B cells cannot be attributed to a quantitative loss of total CD4 T cells previously described in HIV-infected patients (30).

There was, however, evidence for qualitative T cell dysfunction in vaccine NR based on analysis of serum IL-21 levels, which may have contributed to the failure of maturation and expansion of ASC. Increases in serum IL-21 at T2 were noted selectively in the R patients and HC in this study and occurred concomitantly with increase in frequency and MFI of IL-21R+ B cells, suggesting that responsiveness to IL-21 was important for the Ab response. As IL-21 is produced mainly from CD4 cells, in particular from T follicular helper cells (52), reconstitution of this subset of CD4 T cells with cART in HIV-infected patients may be critically important for restoring B cell function.

Morita et al. (24) recently reported that specific CXCR5+CD4+ T cell subsets in peripheral blood represent T follicular cell counterparts and induce Ig production by B cells through secretion of IL-21. We found that frequency of peripheral CD4+CXCR5+ cells increased from T0 to T2 in R patients and HC (S.Y. Silva, S. Pallikkuth, S. Kantihekel, M. Fischl, R. Pahwa, and S. Pahwa, manuscript in preparation). Thus, it is quite possible that these CD4 T cells are involved in regulation of vaccine-induced Ab responses, and a deficiency in this subset is not readily apparent in total CD4 T cell analysis because of its low frequency in peripheral blood. A deficiency of HIV-specific IL-21–secreting CD4 T cells following ex vivo activation has in fact been described in HIV infection (32–34). The importance of IL-21 in HIV infection has been attributed mainly to its role in augmenting antiviral properties of CD8 T cells (34, 53), and recently, IL-21 was shown to be more highly expressed in CD4 T cells of HIV-infected elite controllers versus progressors (32, 33). Our data suggest that IL-21 may also play an important role in regulating B cell differentiation to generate novel 2009 H1N1 influenza vaccine-stimulated memory B cells and that failure to upregulate IL-21/IL-21R could constitute a critical defect, leading to impaired vaccine-induced Ab responses in HIV-infected patients.

Upregulation of IL-21R on B cells (including LT, N, and memory subsets) was strongly correlated with the ability to make a serologic response to novel 2009 H1N1 vaccine.Activation of B cells is known to upregulate IL-21/IL-21R could constitute a critical defect, leading to impaired vaccine-induced Ab responses in HIV-infected patients.

![Figure 3](http://www.jimmunol.org/)

**FIGURE 3.** Serum IL-21 levels and IL-21R expression in B lymphocyte subsets in HC and HIV-infected patients pre- and post-H1N1 vaccination: serum levels of IL-21 were measured by ELISA (A). Fresh PBMC (1 × 10^6 cells) were stained with surface markers for B cell subset characterization (CD20, CD21, CD10, CD27) and for IL-21R. Total B cells and various B cell subsets were analyzed at baseline (T0), 7 d (T1), and 28 d (T2) postvaccination for IL-21R by flow cytometry. Data are shown for T0 and T2 as no changes were noted at T1 compared with T0. B. Representative histograms showing frequencies and MFI (in parentheses) of IL-21R expression in total B cells of eight HC, eight HIV-infected H1N1 Ab responders, and eight HIV-infected H1N1 Ab nonresponders at T0 (gray) and T2 (black) compared with isotype controls (filled). C. Box plots are shown to illustrate frequency of IL21R expression in total B cells. D. Correlation of H1N1 Ab titer with percentage of IL21R+ B cells at T2 in HIV-infected patients. E. Frequency of IL21R expression B cell subsets at baseline. Box plots include median with 25th and 75th percentile borders, and error bars represent 10th and 90th percentiles. Stars indicate the level of significance. ★ p < 0.05, ★★ p < 0.01, ★★★ p < 0.001.
IL-21 was involved in upregulation of IL-21R on B cells is not known. In CD4 T cells, IL-21 is clearly known to exercise amplification of its receptor (55). If a similar scenario were true for B cells, failure of IL-21R amplification by IL-21 could be a cause for the impaired IL-21R upregulation in B cells of NR patients. Preliminary observations made with PBMC of healthy donors suggest that ex vivo culture with IL-21 can result in increased frequencies and MFI of IL-21R expression in total B cells and in RM, LT, and N B cell subsets. Intrinsic B cell defects in IL-21R expression cannot be ruled out, as baseline frequencies of IL-21R–expressing total B cells and IL-21R+ LT and RM B cell subsets were reduced in patients despite equivalent serum IL-21 levels among patients and HC at T0. Regardless of the underlying mechanism, failure to upregulate IL-21R was a distinguishing feature of NR vaccine recipients. Collectively, the data for IL-21/IL-21R upregulation and the associated expansion of memory B cells in R patients suggest that this ligand–receptor pair was contributing to the regulation of B cell differentiation to generate novel 2009 H1N1 influenza vaccine-stimulated memory B cells and that failure to upregulate IL-21/IL-21R could be an underlying mechanism for impaired vaccine-induced Ab responses in HIV-infected patients. This contention is supported by other studies in which upregulation of IL-21/IL-21R were shown to be critical for memory B cell generation (20, 25, 56, 57). IL-21 signals through the common \( \gamma \)-chain receptor subunit of the IL-21R to phosphorylate STAT3 signaling molecules in a sustained manner in human B cells. STAT3 activation induces B cell maturation with expression of plasma cell genes, phenotypic plasma cell formation, and Ab secretion (20, 58, 59). Defects in IL-21 signaling can impair B cell responses to protein Ags, resulting in poor germinal center persistence and function, impaired transition into memory B cells, poor affinity maturation, and absence of Ig class switching (56, 57). Recently, it was shown that IL-21R knockout mice fail to develop Ag-specific memory B cells and plasma cells and have severely compromised secondary immune responses (25). The addition of exogenous IL-21 to cultures of H1N1-stimulated memory B cells resulted in significant enhancement of ASC responses only in R patients and HC. The inability of IL-21 to enhance memory B cell responses ex vivo in NR patients could partially be a consequence of the decreased frequency of IL-21R–expressing memory B cells in addition to a reduced memory pool size. Moreover, the increased frequency of B cells with the exhausted phenotype in NR patients may render them unresponsive to helper signals.

An expansion of plasmablasts on day 7 postvaccination was another benchmark of successful vaccination in this study. Integrated signals from a wide spectrum of sources regulate the ability of B cells to differentiate into pathways resulting in generation of SPB and memory cells (18, 59). A rapid expansion SPB originating from memory cells is regulated largely by innate immune factors (60, 61). On day 7 post-seasonal influenza vaccination, healthy donors have been shown to manifest an expanded functional plasmablast population originating largely from Ag-responsive memory B cells (30, 43, 46, 47) that constitute \(~6\%\) of peripheral B cells (43). Surprisingly, in R patients in our study, we observed a similar kinetic and frequency of SPB, the presumed source of spontaneous H1N1 ASC detected at T1 in the ELISPOT assay. Although we cannot exclude the contribution of cross-

**FIGURE 4.** H1N1-specific ASC in healthy controls and HIV-infected patients pre- and post-H1N1 vaccination: ASC response was measured by ELISPOT in HC and HIV-infected patients. Box plots include median with 25th and 75th percentile borders, and error bars represent 10th and 90th percentiles. Spontaneous H1N1-specific ASC/10^6 B cells (A), IgG-producing ASC following ex vivo stimulation with H1N1 Ag (B), and IgG-producing ASC following ex vivo stimulation with SAC at baseline (C). D, Correlation between serum H1N1 Ab titer and H1N1-stimulated IgG ASC at T2 in PBMC of HIV-infected individuals. Stars indicate the level of significance. ★\( p < 0.05, ★★\( p < 0.01, ★★★\( p < 0.001.

**FIGURE 5.** H1N1 Ag-stimulated IgG responses in HC and HIV-infected patients pre- and post-H1N1 vaccination: H1N1 vaccine Ag-stimulated IgG ASC with and without exogenous IL-21 at T0 and T2. Box plots include median with 25th and 75th percentile borders, and error bars represent 10th and 90th percentiles. Stars indicate the level of significance. ★\( p < 0.05, ★★\( p < 0.01, ★★★\( p < 0.001.
reactive memory B cells (3, 46) to the SPB pool, it is more likely that SPB resulted from expansion of unprimed B cells (62), rather than pre-existing Ag-specific memory cells. It is interesting that the IL-21R knockout mice who manifested severe deficits in memory B cells and recall Ab responses also had a poor early primary immune response (25). We contend that the IL-21R–IL-21 interaction contributes in some measure to the expansion of SPB from IL-21R+ immature and N B cells in a primary immune response, in addition to the well-established role of IL-21/IL-21R signaling in the development of long-lived memory cells.

The results of our studies point to the need to define the molecular basis of the observed immunologic defects in efforts toward optimization of vaccination strategies for influenza viruses. Vaccine-induced transcriptional regulation of B cell survival, proliferation, differentiation, class switch, and somatic mutation need further study. Levels of activation-induced cytidine deaminase, important for regulating class switch, were recently found to be deficient in ex vivo influenza vaccine-stimulated B cells in aging humans (63) who are known to have poor Ab responses to influenza vaccines and have poor in vitro B cell differentiation responses (64). Given the prevailing notion that the immune system in HIV infection undergoes premature immune senescence (65–67), this line of investigation is worthwhile. We also recognize the need to conduct more precise studies in purified cell populations to address critical aspects of B cell maturation in the NR patients and to evaluate the role of innate factors such as BAFF and a proliferation-inducing ligand, which are important in B cell maturation and differentiation (16, 60, 68–70) (S. Pallikkuth, S. Pilaka Kanthikee, S.Y. Silva, M. Fischl, R. Pahwa, and S. Pahwa, manuscript in preparation). It is of interest that a combination of IL-21 and BAFF in the absence of further costimulation is capable of inducing differentiation of memory B cells that reside in the marginal zone region of the human spleen (71).

The major limitation of this pilot study is the small sample size and its effect on data interpretation. The small sample size may have masked greater differences between HC and patients and between the two patient groups. To maximize the impact of the study, we ensured that patients were virologically controlled on cART, randomly selected prior to vaccination before knowing the vaccination outcome, and studied prospectively. Despite the size and gender distribution limitations, differences between R and NR patients as indicated in Table I are quite remarkable. Not only did the proportion who failed to seroconvert closely match the published data (10, 12, 13), but also distinct and significant differences in immune responses were identified that set the NR apart from R patients and also revealed subtle differences between R patients and HC. The IL-21/IL-21R upregulation was associated with expansion of phenotypic and functional memory B cells, which, together with expansion of SPB with effector function, stood out as key indicators of successful vaccine-induced activation of the immune system. Future studies to evaluate IL-21/IL-21R screens to predict responsiveness to influenza vaccines at the level of individual patients may be worthwhile. These studies point to the need for continuing investigations to examine the cumulative signaling elements that control B cell differentiation in response to vaccines for insight into improving immunization strategies for immune deficient patients against influenza viruses.

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Disclosures

The authors have no financial conflicts of interest.

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Supplementary figure 1: Total B cells and B cell subsets in healthy controls and HIV-infected patients pre- and post H1N1 vaccination: Fresh PBMC (1x10^6 cells) were stained for surface markers for B cell subset characterization (CD20, CD21, CD10, CD27). Box plots showing frequency of (A), Total B cells; (B), ET; (C), LT; (D), Naive; (E), ETL subsets. Box plots include median with 25th and 75th percentile borders and error bars represent 10th and 90th percentiles. Stars indicate the level of significance (★ p<0.05, ★★★ P<0.01, ★★★★★ p<0.001).
Supplementary Figure 2: IL-21R expression in B lymphocyte subsets in healthy controls and HIV-infected patients pre and post H1N1 vaccination: Fresh PBMC (1x10^6 cells) were stained for surface markers for B cell subset characterization (CD20, CD21, CD10, CD27) and for IL-21R. Total B cells and various B cell subsets were analyzed at baseline (T0), 7 days (T1) and 28 days (T2) post vaccination for IL-21R by flow cytometry. Data is shown for T0 and T2 as no changes were noted at T1 compared to T0. Box plots are shown to illustrate frequency of IL21R expression in B cell subsets (A), LT; (B), Naïve; (C), RM; (D), AM B cell subsets. Box plots include median with 25th and 75th percentile boarders and error bars represent 10th and 90th percentiles. (★ p<0.05, ★★★ P<0.01, ★★★★★ p<0.001).