Immune recovery in HIV-infected patients after 
*Candida* esophagitis is impaired despite long-term antiretroviral therapy

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**Objective:** *Candida* esophagitis belongs to the most common AIDS-defining diseases; however, a comprehensive immune pathogenic concept is lacking.

**Design:** We investigated the immune status of 37 HIV-1-infected patients from the Swiss HIV cohort study at diagnosis of *Candida* esophagitis, 1 year before, 1 year later and after 2 years of suppressed HIV RNA. We compared these patients with three groups: 37 HIV-1-infected patients without *Candida* esophagitis but similar CD4\textsuperscript{+} cell counts as the patients at diagnosis (advanced HIV group), 15 HIV-1-infected patients with CD4\textsuperscript{+} cell counts higher than 500 cells/\mu l, CD4\textsuperscript{+} cell nadirs higher than 350 cells/\mu l and suppressed HIV RNA under combination antiretroviral therapy (cART) (early cART group) and 20 healthy individuals.

**Methods:** We investigated phenotype, cytokine production and proliferative capacity of different immune cells by flow cytometry and enzyme-linked immunosorbent spot.

**Results:** We found that patients with *Candida* esophagitis had nearly abolished CD4\textsuperscript{+} cell proliferation in response to *Candida albicans*, significantly increased percentages of dysfunctional CD4\textsuperscript{+} cells, significantly decreased cytotoxic natural killer cell counts and peripheral innate lymphoid cell counts and significantly reduced IFN-\gamma and IL-17 production compared with the early cART group and healthy individuals. Most of these defects remained for more than 2 years despite viral suppression. The advanced HIV group without opportunistic infection showed partly improved immune recovery.

**Conclusion:** Our data indicate that *Candida* esophagitis in HIV-1-infected patients is caused by an accumulation of multiple, partly *Candida*-specific immunological defects. Long-term immune recovery is impaired, illustrating that specific immunological gaps persist despite cART. These data also support the rationale for early cART initiation to prevent irreversible immune defects.

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**Keywords:** *Candida* esophagitis, early combination antiretroviral therapy, HIV, IL-17 response, long-term immune recovery, proliferative impairment

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Introduction

The risk of opportunistic infections in patients with HIV infection has markedly declined since 1996 because of the widespread use of combination antiretroviral therapy (cART) [1]. Nevertheless, opportunistic infections still remain a leading complication with an incidence of 16% in late presenting patients [2]. Absolute CD4$^+$ cell counts less than 200 cells/µl and uncontrolled HIV RNA replication are well described major risk factors for the development of opportunistic infection, yet they also occur in patients with CD4$^+$ cell counts higher than 200 cells/µl with an incidence of 10.5 per 1000 patient-years follow-up, highlighting that apart from the absolute CD4$^+$ cell counts, additional risk factors for opportunistic infection must be present [3]. This is further supported by recent studies documenting that early initiation of cART at CD4$^+$ cell counts higher than 500 cells/µl is beneficial as it significantly reduces the risk for opportunistic infection and malignancies [4,5], yet opportunistic infections are not completely eliminated. It remains uncertain why certain HIV-infected patients are susceptible to specific opportunistic infections and how the infection influences long-term immune recovery.

*Candida* esophagitis is one of the most common AIDS-defining diseases, occurring in up to 10–15% of HIV-infected patients before introduction of cART [1,6,7]. Importantly, *Candida* esophagitis is often the first opportunistic infection and also develops in patients with rather high CD4$^+$ cell counts suggesting that the functionality of immune responses is diminished [7].

Earlier studies considered that susceptibility to *Candida* esophagitis is enhanced by a lack of protective Th1 responses and/or a shift to Th2 responses [8]. However, recent studies show that individuals with impaired IL-17 responses exhibit enhanced susceptibility to chronic mucocutaneous candidiasis [9]. In the context of HIV, progressive infection is accompanied by continuous loss of Th17 cells [10] and a decrease in the ratio of Th17 to Th1 cells in peripheral blood [11]. Recently, it has been demonstrated in a mouse model of oropharyngeal candidiasis that IL-17 secreting ROR$\gamma^+$ type 3 innate lymphoid cells (ILCs) also contribute to fungal clearance [12]. Moreover, natural killer (NK) cells are increasingly considered as part of the host defense against fungi [13], and their function was shown to be impaired against *Cryptococcus neoformans* in HIV-infected patients [14].

In this study, we took the advantage of prospectively stored patient samples within the Swiss HIV Cohort Study (SHCS) and investigated the numbers and functions of different immune cell subsets in patients with *Candida* esophagitis over a longitudinal follow-up, including samples before disease development and after long-term suppression of HIV RNA and compared them with three groups of individuals, including HIV-infected patients with similarly advanced HIV infection without opportunistic infection, HIV-infected patients that initiated cART at CD4$^+$ cell nadirs higher than 350 cells/µl and were HIV RNA suppressed and healthy individuals.

Methods

Patients and healthy blood donors

The Swiss HIV Cohort Study is a large prospective observational cohort study with continuous enrolment of adult HIV-infected individuals initiated in 1988 and approved by the local institutional review boards [15]. Basic socio-demographic characteristics, data on clinical course, antiretroviral therapy, immunologic and virologic parameters are collected at enrolment and every 6 months thereafter. Viable peripheral blood mononuclear cells (PBMC) and plasma are stored every 6–12 months. Ethical approval and written informed consent from all patients enrolled in the SHCS have been obtained.

The diagnosis *Candida* esophagitis was based on clinical findings defined according to Centers for Disease Control and Prevention (CDC) criteria [16]. From January 2000 until December 2013, 465 HIV-1 infected patients were diagnosed with *Candida* esophagitis. Of these, 277 patients had *Candida* esophagitis as first and only AIDS-defining disease. Of these, 37 patients with available longitudinal PBMC were included. We analyzed cryopreserved PBMC from three time points: 6–18 months before diagnosis, at diagnosis (±6 months) and 6–18 months after diagnosis. For patients with suppressed HIV RNA (<50 copies/ml) over 2 years, an additional time point was included. These patients were compared with three groups. First, HIV-1-infected patients with similarly advanced disease but without opportunistic infection. Patients were matched to *Candida* esophagitis patients according to CD4$^+$ cell counts (±25 cells/µl), date of diagnosis of *Candida* esophagitis, use of cART, sex, age and absence of other opportunistic infection within 6 months prior to sample collection [17]. As for the *Candida* esophagitis patients, four time points were analyzed. Second, 15 SHCS patients with well controlled HIV-1 infection from outpatients of the HIV clinic at the University Hospital of Basel. Patients had the following criteria: HIV CDC A1 or A2 classification with suppressed HIV RNA (<50 copies/ml) and stable cART therapy for at least 6 months, CD4$^+$ cell counts higher than 500 cells/µl and CD4$^+$ cell nadirs higher than 350 cells/µl. Third, 20 healthy individuals after receipt of informed consent according to the ethic approval from the Ethikkommission Nordwest und Zentralschweiz (EKBB 242/11). For both latter groups, only one time point was analyzed. Baseline characteristics of patients and healthy individuals are included in the Supplementary table, http://links.lww.com/QAD/A947.
**Generation of heat-inactivated C. albicans**  
A mixture of *Candida albicans* strain SC5314 yeast and hyphae was cultured and heat-inactivated as previously described [18,19].

**Phenotypic characterization**  
T-cell activation/exhaustion was analyzed by staining with anti-CD3-PerCP, anti-CD4-PacificBlue, anti-CD8-APC, anti-CD25-PE/Cy7 and anti-CD279-PE (PD-1); ILCs by staining with anti-Lineage-Cocktail-APC (anti-CD3/CD14/CD16/CD19/CD20/CD56) and anti-CD127-PE; NK-cell subsets by staining with anti-CD3-PerCP, anti-CD14-FITC, anti-CD19-PE/Cy7, anti-CD56-APC/Cy7, anti-CD16-PacificBlue (all BioLegend, Fell, Germany), anti-NKG2A-APC (clone 131411) and anti-NKG2C-PE (both R&D Systems, Abingdon, UK) [20]. Samples were acquired on a CyAn ADP Analyzer (Beckman Coulter, Nyon, Switzerland) and data analyzed with FlowJo software vX.0.7 (FlowJo, Ashland, Oregon, USA).

**Enzyme-linked immunosorbent spot assay**  
IFN-γ and IL-17 enzyme-linked immunosorbent spot (ELISPOT; Mabtech; Abingdon, UK) were performed according to manufacturer's instructions as previously published [21]. Briefly, 3–5 × 10⁵ cells/well were stimulated in duplicates with *C. albicans* [multiplicity of infection (MOI) 0.05], 0.05 μg/ml cytomegalovirus (CMV) pp65 (JPT Peptide Technologies, Berlin, Germany) or 0.5 μg/ml staphylococcal enterotoxin B (SEB; Sigma-Aldrich, Buchs, Switzerland) for 72 h. The number of spot forming units per well was counted by ELISPOT reader (Cellular Technologies Ltd., Bonn, Germany). Data are shown after subtraction of unstimulated controls.

**Proliferation assay**  
PBM of patients were labeled with carboxyfluorescein diacetate succinimidyl ester (CFSE, Invitrogen, Fisher Scientific, Reinach, Switzerland) according to manufacturer's instructions and stimulated with *C. albicans* (MOI 0.05) or 0.5 μg/ml SEB for 7 days in RPMI 1640 (Gibco, Fisher Scientific, Reinach, Germany) with 5% pooled human serum. Medium was replenished as needed. Cells were stained with anti-CD3-PerCP, anti-CD4-PacificBlue, anti-CD8-APC and anti-CD56-APC/Cy7 (all Biolegend) and acquired on a CyAn ADP Analyzer (Beckman Coulter) and data analyzed with FlowJo software vX.0.7.

**Statistical analysis**  
Comparisons between two groups were performed with the two-sided Mann–Whitney U test. P values of 0.05 or less were considered statistically significant. Statistical analyses were done using GraphPad Prism 6.0f and Stata 13.1 software (StataCorp LP, College Station, Texas, USA). Shown are median values ± interquartile ranges (Tukey plots).

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**Results**

**Patients with *Candida* esophagitis have low and dysfunctional CD4⁺ cell counts and decreased *Candida*-specific cytokine responses and proliferative capacity**  
We first analyzed T-cell phenotype and function of patients diagnosed with *Candida* esophagitis and compared them with HIV-1-infected individuals with early initiation of cART and sustained viral suppression (<50 copies/ml) and healthy individuals (for baseline characteristics of patients and healthy individuals see the Supplementary table, http://links.lww.com/QAD/A947).

As expected from earlier studies, absolute CD4⁺ cell counts were significantly lower in *Candida* esophagitis patients compared with patients with early initiated cART and healthy individuals (Fig. 1a) and the frequencies of exhausted PD-1⁺ and activated/regulatory CD25⁺CD4⁺ cell counts were significantly increased compared with healthy individuals (Fig. 1b). The median percentage of CD25⁺ CD4⁺ cell counts was 6-fold higher in patients with *Candida* esophagitis compared with healthy individuals. In accordance with the significantly reduced number of functional CD4⁺ cell counts, the IFN-γ and IL-17 responses of PBMC to the superantigen SEB or *C. albicans* were significantly lower in *Candida* esophagitis patients compared with patients with early initiated cART and healthy individuals. The IFN-γ response to CMV pp65 in CMV seropositive donors was not affected, showing that viral reactivation was still able to trigger functional immune responses independent of the CD4⁺ cell counts (Fig. 1c and d).

In line, the proliferative capacity of CD4⁺ and CD8⁺ cells to *C. albicans* was significantly lower in patients with *Candida* esophagitis compared with healthy individuals (Fig. 1e and f). The median percentage of proliferating CD4⁺ cell counts was 7-fold and of proliferating CD8⁺ cell counts 12-fold decreased compared with healthy individuals. Interestingly, CD4⁺ cell proliferation to SEB was comparable to healthy individuals. To examine whether the inability to proliferate was due to a lack of IL-2 production, we supplemented some cultures with 50 U/ml recombinant IL-2 on day 2. However, T-cell proliferation was not improved (Supplemental Fig. 1, http://links.lww.com/QAD/A920).

Thus, development of *Candida* esophagitis was associated with reduced and dysfunctional CD4⁺ cell counts that showed significant impairments in cytokine production and proliferation to specific antigens.

**Patients with *Candida* esophagitis have decreased peripheral natural killer cells and innate lymphoid cells**  
NK cells and ILC are increasingly considered as part of the host defense against fungi. We therefore investigated...
whether these cells and their functionality are also impaired in patients with *Candida* esophagitis.

Absolute NK-cell counts (CD3<sup>+</sup>CD56<sup>+</sup>) and especially the cytotoxic CD16<sup>+</sup> NK-cell subset were significantly lower in patients with *Candida* esophagitis compared with patients with early initiated cART and healthy individuals (Fig 2a). The absolute number of CD16<sup>+</sup> NK cells was with a median of 42 cells/μl nearly 5-fold lower than in healthy controls (201 cells/μl). The percentage of NK cells expressing the inhibitory receptor NKG2A was lower in patients with *Candida* esophagitis than in the other groups, whereas the percentage of cells expressing the activating receptor NKG2C was higher in *Candida* esophagitis cases and in HIV-1-infected virologically suppressed patients than in healthy individuals (Fig 2b). The proliferative capacity of NK cells to *C. albicans* was not significantly affected (Fig 2c).

Similar to NK cells, also peripheral ILC counts (lineage<sup>+</sup>CD127<sup>+</sup>) were significantly reduced in patients with *Candida* esophagitis compared with patients with early initiated cART and healthy individuals (Fig 2d).

In conclusion, additionally to defective CD4<sup>+</sup> cell responses, patients with *Candida* esophagitis had significantly reduced numbers of NK cells and ILC.

### The proliferative responses to *C. albicans* and NK-cell counts and function are impaired despite higher CD4<sup>+</sup> cell counts at diagnosis

Next, we examined whether higher CD4<sup>+</sup> cell counts at diagnosis of *Candida* esophagitis are associated with better functionality of the different cell subsets (Fig 3).

Patients with CD4<sup>+</sup> cell counts higher than 350 cells/μl (18 of 37 patients) showed decreased percentages of dysfunctional PD-1<sup>+</sup> and CD25<sup>+</sup> CD4<sup>+</sup> T cells and increased cytokine responses to SEB and *C. albicans* compared with patients with CD4<sup>+</sup> cell counts less than 200 cells/μl (12 of 37 patients; Fig 3a–c; the Supplementary table, http://links.lww.com/QAD/A947). By contrast, CMV-specific responses were comparable regardless of the absolute CD4<sup>+</sup> cell counts. The proliferative response to *C. albicans* was reduced even in patients with CD4<sup>+</sup> cell counts higher than 350 cells/μl (Fig 3d).

Although NK-cell counts and the percentage of cytotoxic CD16<sup>+</sup> NK cells increased with higher CD4<sup>+</sup> cell counts, the absolute numbers of cytotoxic NK cells in patients with CD4<sup>+</sup> cell counts higher than 350 cells/μl still remained more than 4-fold reduced compared with healthy individuals (Fig 3e). The number of ILC in peripheral blood also significantly increased with higher CD4<sup>+</sup> cell counts (Fig 3f).
Patients with *Candida* esophagitis show a significant drop in CD4$^+$ cell counts and nearly abolished T-cell proliferation to *C. albicans* at diagnosis and retain immunological impairments even after disease resolution and successful combination antiretroviral therapy

We further analyzed the immune status of patients with *Candida* esophagitis prior to disease development to identify immunological changes associated with disease development and after disease resolution and successful cART to identify possible long-term defects. We therefore additionally examined PBMC 6–18 months before diagnosis, after disease resolution (6–18 months after diagnosis) and after successful cART with stably suppressed HIV RNA (<50 copies/ml) for more than 2 years (Fig. 4).

Interestingly, the patients with *Candida* esophagitis showed a significant drop in absolute CD4$^+$ cell counts at diagnosis and nearly abolished proliferation of CD4$^+$ and CD8$^+$ cells in response to stimulation with *C. albicans*. Proliferation to SEB was not significantly affected. Also NK-cell proliferation in response to *C. albicans* dropped during development of *Candida* esophagitis (Fig. 4a,f,i).

In line with previous studies, *Candida* esophagitis patients with low CD4$^+$ cell nadirs (<320 cells/μl, median 87 cells/μl) restored CD4$^+$ cell counts after viral suppression to a lower level compared with patients with early cART and higher CD4$^+$ cell nadirs (>350 cells/μl, median 397 cells/μl) (Fig. 4a). The percentage of CD25$^+$ CD4$^+$ cell counts and the IFN-γ responses to SEB and *C. albicans* normalized after suppression of viral replication, whereas the IL-17 response remained impaired despite long-term viral suppression under cART (Fig. 4b–d). The proliferative capacity of CD4$^+$ cells to *C. albicans*
after suppression of HIV replication recovered but remained in median lower than in patients with early initiated cART and healthy individuals (Fig. 4e and f).

Median NK-cell counts remained below 200 cells/µl despite viral suppression. Although the percentage of CD16<sup>+</sup> cells increased, the absolute number of cytotoxic CD16<sup>+</sup> NK-cell counts remained more than 3.5-fold lower compared with healthy individuals (Fig. 4g). By contrast, the percentages of NKG2A<sup>+</sup> and NKG2C<sup>+</sup> NK cells and the proliferative responses were comparable with healthy controls (Fig. 4h and i). Similarly to NK cells, also ILC counts remained 2.5-fold reduced for at least 2 years after viral suppression (Fig. 4j). NK-cell and ILC reconstitution seemed to correlate directly with CD4<sup>+</sup> cell reconstitution (Supplemental Fig. 2, http://links.lww.com/QAD/A920).

The data show that development of Candida esophagitis is associated with a drop in absolute CD4<sup>+</sup> cell counts and proliferative capacity of Candida-specific T cells and that despite successful cART, patients with previous Candida esophagitis have prolonged immune defects of CD4<sup>+</sup> cells, NK cells and ILC.

**Patients with similarly advanced HIV infection but without an opportunistic infection show overall better immune recovery after successful combination antiretroviral therapy**

The last comparator group consisted of patients with similarly advanced HIV infection but without opportunistic infection. These patients were matched to Candida esophagitis patients according to CD4<sup>+</sup> cell counts, use of cART, sex, age and absence of other opportunistic infection within 6 months prior to sample collection. Also the viral load was comparable between these two patient groups. Notably, patients with similarly advanced HIV infection but without opportunistic infection showed overall very similar immune cell impairments and reconstitution as patients with Candida esophagitis (Supplemental Fig. 3, http://links.lww.com/QAD/A920). However, they differed by a 2-fold lower percentage of CD25<sup>+</sup> CD4<sup>+</sup> cells prior to disease development and consistently higher CD4<sup>+</sup> cell counts at diagnosis (a) Percentage of PD1<sup>+</sup> and CD25<sup>+</sup> CD4<sup>+</sup> T cells, (b) IFN-γ response of peripheral blood mononuclear cell to staphylococcal enterotoxin B or heat-inactivated C. albicans, (c) IL-17 response of peripheral blood mononuclear cell to staphylococcal enterotoxin B or heat-inactivated C. albicans, (d) percentage of proliferating (CFSE<sup>dim</sup>) cells in the CD4<sup>+</sup> T-cell population after 7 days stimulation with staphylococcal enterotoxin B or heat-inactivated C. albicans, (e) absolute CD3<sup>+</sup> CD56<sup>+</sup> natural killer cell counts, percentage of CD16<sup>+</sup> natural killer cell counts, absolute CD16<sup>+</sup>CD3<sup>+</sup>CD56<sup>+</sup> natural killer cell counts, percentages of NKG2A<sup>+</sup> and NKG2C<sup>+</sup> natural killer cell counts and (f) absolute lineage CD127<sup>+</sup> CD4<sup>+</sup> T-cell counts less than 200 cells/µl or more than 350 cells/µl. Shown are median values + interquartile ranges (Tukey plot). Broken and dotted lines represent medians of healthy individuals (healthy donor) and HIV-1-infected patients with early initiated combination antiretroviral therapy and a viral load less than 50 copies/ml (viral load <50), respectively. Data (b)–(d) are shown after subtraction of unstimulated controls. Number of patients less than 200/more than 350 CD4<sup>+</sup> cell counts were n = 10/16 (a), n = 8/7 (b), n = 5/5 (c), n = 1/3 (d), n = 9/13 (e) and n = 11/13 (f). *P ≤ 0.05 and **P ≤ 0.01 (Mann–Whitney test).
proliferation to *C. albicans*. In addition, long-term recovery of the IL-17 response, total NK cells, CD16⁺ cytotoxic NK cells and ILCs was better in these patients without *Candida* esophagitis.

**Discussion**

The detailed pathogenesis of opportunistic infections is still unknown for many pathogens. *Candida esophagitis* is one of the most frequent opportunistic diseases in untreated HIV-infected individuals but also occurs in patients with other underlying conditions.

In this study, we investigated in depth 37 HIV-1-infected *Candida esophagitis* patients and compared them with advanced HIV-1-infected patients without opportunistic infection, HIV-1-infected patients with early initiation of cART and healthy individuals. We found that patients with *Candida esophagitis* showed a significant drop in CD4⁺ cell counts at diagnosis, a nearly abolished proliferative capacity to *C. albicans*, an impaired IFN-γ and IL-17 production to *C. albicans* and a dysfunction of CD4⁺ cells with increased percentages of CD25⁺ and PD-1⁺ cells. In addition, these patients had significantly decreased peripheral ILC and cytotoxic NK-cell counts. Recovery of the proliferative capacity of CD4⁺ cc and IL-17 production to *C. albicans* and of ILCs and cytotoxic NK cells was impaired for years despite effective cART.

HIV infection is commonly associated with an inability to proliferate and produce IL-2. Even patients with normal
IFN-γ production often have a proliferative defect to different antigens, especially patients with previously low CD4+ cell counts and persistent HIV replication [22–24]. In line with these data, we observed an overall decreased proliferative capacity to Candida in patients with Candida esophagitis at diagnosis and even after recovery of the IFN-γ response. These findings suggest that the proliferative defect could contribute to disease development and that these patients might remain vulnerable despite effective cART.

CD4+ T cells from chronically infected HIV patients show diminished IFN-γ and IL-17 production [25], but data on cytokine production in response to different fungi in HIV-infected patients are scarce. This study demonstrates that all patients with advanced HIV infection including patients with and without Candida esophagitis had an overall impaired IFN-γ and IL-17 production to C. albicans, which probably is partly due to the significantly reduced CD4+ cell count. However, HIV-infected patients with early initiation of cART had comparable or even higher Candida-specific cytokine responses compared with healthy individuals despite significantly reduced CD4+ cell counts, showing that low CD4+ cell counts do not necessarily lead to reduced antigen-specific responses. Similarly, CMV pp65-specific IFN-γ production by CMV-seropositive patients was comparable in all groups, independent of CD4+ cell counts. It is however possible that patients with acute Candida esophagitis have additional impairments in antigen-presenting cells leading to reduced T-cell responses or that Candida-specific T cells in these patients are recruited to sites of infection and therefore disappear from the blood. Interestingly, the recovery of IL-17-producing cells was slower in patients with Candida esophagitis compared with patients with advanced HIV infection without opportunistic infection. These data fortify previous findings in chronic mucocutaneous candidiasis in humans and oropharyngeal candidiasis in mice [9–11,25,26]. Interestingly, CMV pp65-specific IFN-γ production by CMV-seropositive patients was comparable to healthy controls, showing that not all antigen-specific responses are impaired in these patients. [NB change g in IFN-γ to Greek gamma].

We further found that patients with Candida esophagitis had a significantly higher percentage of CD25+ CD4+ cells before disease development compared with healthy individuals. The patients with advanced HIV without Candida esophagitis showed a 2-fold lower percentage compared with the patients with Candida esophagitis. We cannot state if these cells were activated or regulatory T cells, as we did not include additional markers. However, previous studies showed that a higher percentage of regulatory T cells was associated with lower HIV-specific and Candida-specific responses [27]. Furthermore, in-vitro depletion of the Treg-containing CD25+ T-cell population greatly enhanced the response to HIV and CMV antigens [28–30]. Moreover, PD-1+ CD4+ cells significantly increased at diagnosis of Candida esophagitis, which further supports the assumption that a dysfunction of CD4+ cells might be one factor leading to susceptibility to Candida esophagitis.

Healthy individuals also show proliferation of CD8+ cells in response to C. albicans. Previous work in mouse models has shown that in the absence of CD4+ cells, CD8+ cells were able to confer protection to the fungal pathogens Blastomyces dermatitidis and Histoplasma capsulatum [31]. Interestingly, in our study we did not observe compensatory proliferation of CD8+ cells in patients with low CD4+ cell responses after stimulation with C. albicans. It is possible that the CD8+ cells proliferating to C. albicans in healthy individuals are mucosal-associated invariant T cells that respond to C. albicans and are depleted in the course of HIV infection [32–34].

Recently, it has been shown that not only Th17 cells, but also other cells such as ILCs can be a source of IL-17 and are involved in the host defense against fungal infections [12]. We found that the ILC counts in peripheral blood of Candida esophagitis patients were unable to recover even with suppressed viral replication. Remarkably, patients with CD4+ cell counts higher than 350 cells/μL had significantly higher ILC counts in the peripheral blood compared with patients with CD4+ cell counts less than 200 cells/μL, suggesting that the loss of ILCs occurs in parallel with the loss of CD4+ cells over time and may additionally increase the risk of developing Candida esophagitis. Furthermore, recovery of ILCs after suppressed viral replication seems to be correlated with CD4+ cell recovery. Nevertheless, these data should be interpreted with caution, as the number of patients is low, and we did not investigate the different subsets of ILCs and their involvement in the mucosa.

Also NK cells are increasingly considered as part of the host defense against fungi [13,14,35–37]. Impaired NK-cell activity was observed in patients with chronic mucocutaneous candidiasis [38]. Also HIV-infected individuals show quantitative and functional NK-cell impairments that continue during disease progression, such as a decrease of the CD3 CD56+ cell subset, a decreased cytotoxic capacity and aberrant expression of several surface receptors [39–42]. In fact, we found significantly lower NK-cell counts and a significantly lower percentage of CD16+ cytotoxic NK cells in patients with Candida esophagitis compared with healthy individuals and patients with early initiated cART. They also did not recover under stable, virologically successful cART. Furthermore, similar to ILCs, NK-cell recovery seemed to correlate with CD4+ cell recovery.

Recently, evidence has accumulated that early initiation of cART is beneficial for virological as well as immunological parameters. Early treatment decreases
cell–associated HIV RNA and DNA and limits the HIV reservoir [43–47], maintains numbers and function of the CD4\(^+\) cell compartment [4,48–50] and reduces the risk for disease transmission and the development of opportunistic viral and fungal infections and malignancies [4,51,52]. However, it was not clear, how early or late treatment affects T-cell responses to opportunistic pathogens and how other immune cell subsets such as NK cells or ILC are affected. In this study, we could confirm improved overall and Candida-specific CD4\(^+\) cell recovery in patients with early cART. We could further show that ILC and NK-cell reconstitution, immune cells with a likely role in antifungal defense, correlated with CD4\(^+\) cell recovery and was therefore superior in HIV-1-infected patients with early treatment, further arguing for early initiation of cART in HIV-1-infected individuals.

The strength of this study is the comprehensive longitudinal analysis of quantitative and qualitative immune responses in a large number of HIV-infected patients with Candida esophagitis. This allowed identifying significant immunological impairments compared with healthy individuals and HIV-infected patients with early initiation of cART. However, due to the limited availability of PBMC, we did not further characterize different cell subsets such as T\(_{\text{eq}}\) or T\(_{\text{H17}}\) cells, and functional analysis could not be performed in every sample.

In conclusion, this study demonstrates that HIV-1-infected patients with Candida esophagitis not only have deficient T-cell responses, but an accumulation of multiple, partly Candida-specific immunological defects. This may explain the fact that despite the high frequency of Candida esophagitis, only a part of AIDS patients develop this opportunistic infection. These defects are only partially reversible under cART, and long-term immune impairments remain. This is particularly apparent in patients with low CD4\(^+\) cell counts at initiation of cART showing greater general and Candida-specific immune impairments initially and under stable cART. Nevertheless, certain individuals even experienced Candida esophagitis at higher CD4\(^+\) cell counts. These patients showed similar immune defects as patients with low CD4\(^+\) cell counts highlighting that the presence of specific immunological gaps is relevant. We hypothesize that specific gaps due to underlying genetic and/or immunological predisposition may explain why certain individuals also develop opportunistic infection at higher CD4\(^+\) cell counts. In line with other current studies, our study similarly supports the rationale for early initiation of cART.

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Authors’ contributions: The study was conceived and designed by N.K., C.S., M.B. and C.B. Data acquisition and analysis was performed by C.B. and C.S. Sample recruitment and acquisition of clinical data was performed by C.B., M.S., S.Z., H.F and H.F.G. Statistical analysis was done by L.E. The article was written by N.K., C.S., M.B. and S.L., and reviewed by all coauthors.

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Conflicts of interest

There are no conflicts of interests.

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