CD26/dipeptidyl peptidase IV (CD26/DPPIV) is highly expressed in peripheral blood of HIV-1 exposed uninfected Female sex workers

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Abstract

Background: Design of effective vaccines against the human immunodeficiency virus (HIV-1) continues to present formidable challenges. However, individuals who are exposed HIV-1 but do not get infected may reveal correlates of protection that may inform on effective vaccine design. A preliminary gene expression analysis of HIV resistant female sex workers (HIV-R) suggested a high expression CD26/DPPIV gene. Previous studies have indicated an anti-HIV effect of high CD26/DPPIV expressing cells in vitro. Similarly, high CD26/DPPIV protein levels in vivo have been shown to be a risk factor for type 2 diabetes. We carried out a study to confirm if the high CD26/DPPIV gene expression among the HIV-R were concordant with high blood protein levels and its correlation with clinical type 2 diabetes and other perturbations in the insulin signaling pathway.

Results: A quantitative CD26/DPPIV plasma analysis from 100 HIV-R, 100 HIV infected (HIV +) and 100 HIV negative controls (HIV Neg) showed a significantly elevated CD26/DPPIV concentration among the HIV-R group (mean 1315 ng/ml) than the HIV Neg (910 ng/ml) and HIV + (870 ng/ml, p < 0.001). Similarly a FACs analysis of cell associated DPPIV (CD26) revealed a higher CD26/DPPIV expression on CD4+ T-cells derived from HIV-R than from the HIV+ (90.30% vs 80.90 p = 0.002) and HIV Neg controls (90.30% vs 82.30 p < 0.001) respectively. A further comparison of the mean fluorescent intensity (MFI) of CD26/DPPIV expression showed a higher DPP4 MFI on HIV-R CD4+ T cells (median 118 vs 91 for HIV-Neg, p = 0.0003). An evaluation for hyperglycemia, did not confirm Type 2 diabetes but an impaired fasting glucose condition (5.775 mmol/L). A follow-up quantitative PCR analysis of the insulin signaling pathway genes showed a down expression of NFκB, a central mediator of the immune response and activator of HIV-1 transcription.

Conclusion: HIV resistant sex workers have a high expression of CD26/DPPIV in tandem with lowered immune activation markers. This may suggest a novel role for CD26/DPPIV in protection against HIV infection in vivo.

Background

The disease AIDS ranks as one of the most devastating scourges of mankind. Since it was identified in 1983 more than 30 million people have died and an estimated 33 million currently live with the virus [1]. The majority of HIV infections occur in Sub-Saharan Africa, where in some countries prevalence rates of more than 40% have been documented among antenatal clinic attendees [2].

Despite progress made with antiretroviral therapy, less than a half of those requiring treatment receive it and new infections far outpace those on therapy [3]. Similar to previous viral epidemics, prevention through vaccination is believed to be the best approach. However several factors including HIV's ability to integrate itself to the host genome and to constantly mutate challenge the design of a safe and effective vaccine. More critically, a sufficient understanding of the immune correlates of protection from HIV infection and disease remains unresolved.
Individuals who though highly exposed to HIV but do not get infected (HIV R) provide an opportunity to better understand what mediates HIV protection. HIV R populations have been identified among intravenous drug users,[4], children born to HIV infected mothers [5,6], discordant couples [7], and commercial sex workers [8]. Of the various investigations conducted on HIV R, the Nairobi commercial sex worker cohort has provided vital clues for the understanding of correlates for protection against HIV infection. With more than 25 years of follow-up, this population has provided compelling evidence that HIV resistance is biologically mediated. The 32 base pair deletion in the CCR5 gene (CCR5 Δ32), which has been reported in Caucasian populations to be responsible for cellular resistance to HIV, has not been observed in this cohort nor have polymorphisms in other cytokine regions reported elsewhere to be connected to HIV resistance [9]. Although cytotoxic T-cell responses and HIV specific mucosal IgA have been noted in the Nairobi cohort [10], controversy surrounds whether they exert a protective effect or reflect a prior exposure to HIV [11]. A genetic basis for HIV resistance has been implied by the finding that among sex worker relatives of the HIV-1 resistant women, there has been a high degree of interfamily correlation of HIV infection independent of the HLA status [12]. This suggests that other genes in pathways beyond the HLA cluster may be key in elucidating the HIV resistance phenomenon.

A recent preliminary gene expression analysis of HIV resistant female sex workers who had been in active sex work for periods of more than seven years and remained HIV uninfected suggested a high expression of CD26/dipeptidyl peptidase IV (CD26/DPPIV) a gene coding for a multifunctional enzyme [13]. Previous studies have shown however that difference in genetic expression might not always correspond to similar protein levels [14]. As high expression of CD26/DPPIV has been shown previously to inhibit HIV-1 infection in vitro [15] validating gene expression reports among the HIV resistant female sex workers through protein analysis may support a quest for further functional studies on CD26/DPPIV gene as a novel contributor to the HIV resistant phenotype. In addition, high CD26/DPPIV protein levels in vivo have been shown to be a risk factor for type 2 diabetes [16]. Incretin hormones involved in insulin signaling have been shown to be degraded by high CD26/DPPIV levels leading to insulin resistance and hyperglycemia.

The present study therefore was to confirm if the high expression of CD26/DPPIV reported at the genetic level among the HIV resistant sex workers were concordant with higher peripheral blood protein levels, and whether this corresponded to clinical type 2 diabetes and perturbation of other genetic events related to insulin signaling.

Results
To investigate the level of expression of CD26/dipeptidyl peptidase IV among HIV highly exposed uninfected sex workers, we conducted a quantitative DPPIV ELISA on plasma samples obtained from 100 HIV resistant sex workers (HIV-R), 100 HIV infected (HIV+) and 100 HIV negative controls (HIV Neg). CD26/DPPIV plasma concentration was significantly elevated among the HIV-R group (mean 1315 ng/ml) than the HIV Neg (910 ng/ml) and HIV + (870 ng/ml, p < 0.001, Figure 1).

To determine whether the observed differences in soluble plasma expression would correspond to differences CD26/DPPIV expression on T-cells we carried out a FACs analysis on peripheral mononuclear cells derived from the HIV-R, HIV-Neg and HIV+ women. These data revealed higher CD26/DPPIV expression on CD4+ T-cells from the HIV-R group when compared to HIV+ (90.30% vs 80.90 p = 0.002) and HIV Neg controls (90.30% vs 82.30 p < 0.001) respectively (Figure 2a). No difference in CD26/DPPIV expression on CD8+ T cells was observed between the groups (Data not shown). To further characterise CD26/DPPIV expression on CD4+ T-cells, we compared the mean fluorescent intensity (MFI) of CD26/DPPIV expression between HIV-R, HIV Neg and HIV+. This analysis showed a higher CD26/DPPIV MFI on resistant CD4+ T cells

Figure 1 Plasma concentrations of Dipeptidyl peptidase IV enzyme among HIV resistant (HIV-R) HIV positive (HIV+) and HIV negative (HIV neg) controls (n = 100, each).ELISA (R&D Systems, Minneapolis MN), was done in duplicate: dilution factor 1:1000.

p < 0.0001
n, 100; p value calculated by Mann Whitney test
compared to HIV-Neg (median 118 vs 91, p = 0.0003, Figure 2b) suggesting that more CD26/DPPIV molecules are expressed per CD4+T cell in HIV-R subjects.

As high CD26/DPPIV expression has been associated with type 2 diabetes [16], we next determined the blood glucose level of HIV-R women (mean age 38.5 ± 5). Their mean fasting blood sugar level was 5.775 mMol/L (103 mg/dL, Figure 3). Although this was significantly higher than the HIV-Neg (4.886 mmol/L or 87.94 mg/dL p = 0.0061) this was not indicative of clinical Type 2 diabetes mellitus but suggested a prediabetic or impaired fasting glucose state. The 103 mg/dL of the mean fasting blood sugar level of HIV-R subjects was lower than the WHO criteria (110 mg/dL) and slightly higher than the ADA criteria (100 mg/dL) for the impaired fasting glucose state [17,18].

To elucidate further the relationship between HIV-R female sex workers with the diabetic or prediabetic phenotype, we investigated the expression patterns of genes in the insulin signaling pathway which are significantly perturbed during Type 2 diabetes state. To achieve this we randomly selected six HIV-R and an equal number of HIV-Neg controls from the study population.
took their blood samples extracted total RNA and prepared cDNA for gene expression using quantitative real time PCR analysis. We selected a number of genes from the insulin signaling pathway and some that had previously been shown to be related with type 2 diabetes [19]). Relative expression levels of Insulin receptor substrate 1 (INSR-1), PI4K, AKt, TSC2, NFκB, ICOS, LOC6 and Calg were analysed with real time PCR. Figure 4 shows that the differential expression patterns of these genes are consistent with a type 2 diabetes state. INSR, located upstream of the insulin pathway and a target of insulin, was downregulated hence possibly affecting the downstream events such as, PI4K, AKt and TSC2. Similarly immune regulators previously associated with insulin dysregulation such as NFκB, the proinflammatory cytokine mediator S100/Calgranulin and the inducible T-cell costimulator ICOS were down expressed in the HIV-R group compared to the HIV-Neg.

**Discussion**

CD26/DPPIV is a multifunctional protein ubiquitously expressed in both soluble and cell surface forms in various endothelial and epithelial cells including T-cells and exert it different functions depending on cell type and conditions it is expressed [20]. It acts as a proteolytic enzyme, receptor and co-stimulatory protein and its substrates have been reported to be involved in various physiological process including immunomodulation and homeostasis. Dysregulation of CD26/DPPIV has been suggested to result in various pathophysiological process including rheumatoid arthritis, melanoma and Crohn’s disease among others [20,21]. Its association with Type 2 diabetes is however well confirmed with DPPIV inhibitors being used worldwide as approved regimens for anti-type 2 diabetes treatment [22]. Our observation of high CD26/DPPIV expression, significant down expression of genes in the insulin signaling pathway and impaired fasting glucose state (IFG) implied an evolving type 2 diabetes condition among the HIV resistant commercial sex workers. The duration for evolution to full type 2 diabetes from an IFG state is however controversial. While some studies indicate that IFG conditions often lead to diabetes within 3 years, others have suggested a 50% risk over a ten year period [23]. The Nairobi female sex worker cohort has been followed up biannually for periods longer than ten years, and though blood sugar analysis has not been a routine feature, cases of overt Type 2 diabetes has rarely been observed. We postulate that this population, despite its high CD26/DPPIV levels may not be at risk of developing type 2 diabetes. There will however be a need to incorporate type 2 diabetes evaluations in future follow-up to determine the validity of this assertion taking into account specific DPPIV enzyme activity which we did not evaluate in this study.

![Figure 4](http://www.virologyj.com/content/7/1/343)

**Figure 4** Quantitative RT PCR expression patterns of genes previously associated with Type 2 diabetes and in the insulin signaling pathway between HIV Resistant (HIV -R and HIV negative controls (HIV-Neg) (n = 6 each). Calg-Calgranulin, ICOS-Inducible T-cell costimulator, INSR-Insulin receptor substrate 1, LOC6-LOC6 gene, PI4K-phosphatidylinositol 4 kinase, AKt-serine/threonine protein kinase, NFκB-nuclear factor kappa B protein, TSC2-Tuberin.
Although an association of high CD26/DPPIV expression with HIV resistance is well supported by our data at the gene, soluble protein and cell surface level, the mechanism by which it protects against HIV acquisition remains unclear. One possibility is the key role it plays in cleaving off dipeptides from amino termini containing proline or alanine moieties at the penultimate position [24]. A number of chemokines have been shown to share this sequence at their termini including RANTES, an interleukin 8 superfamily chemokine has been shown to inhibit HIV infection by competing with the virus for its coreceptor CCR5. Previous data has shown that in contrast to intact RANTES, a DPPIV truncated RANTES inhibited in vitro HIV infection of mononuclear cells by M-tropic HIV strains fivefold efficiently [15,24]. Similarly several workers have shown that HIV infected subjects have defective CD4+ T cells with a poor ability to recognize and respond to antigens [25,26]. Antigen recognition is the preserve of CD4+ T-cells expressing CD26/DPPIV which proliferate in response to soluble antigens and activate MHC-restricted cytotoxic T cells resulting in annihilation of virally infected cells [20,27]. This has been supported by findings that antigen response in HIV-1 infected individuals could be restored by addition of soluble CD26/DPPIV in vitro, suggesting an important role for CD26/DPPIV in HIV-specific immunity [28].

The probable mechanism of CD26/DPPIV activity against HIV acquisition in the Nairobi cohort may however add a new dimension to the above observations. Our recent findings indicate a significantly lowered immune activation state among the HIV-R female sex workers as compared to HIV negative controls [13,29]. This lowered activation state or immune quiescence has also been observed among HIV exposed seronegative partners of HIV infected spouses [7] and HIV uninfected hemophiliacs transfused with HIV seropositive donor blood [30]. The contribution of CD26/DPPIV to a lowered immune activation state among the HIV-R female sex workers is puzzling.

There is substantial evidence that CD26/DPPIV can act as the trigger to immune activation [31-34]. This has been supported by studies showing that reversible DPPIV inhibitors suppress proliferation of human PBMCs s and enhance production of cytokines that inhibit antigen stimulation of T-cells [35]. Our findings of high CD26/DPPIV expression in vivo in an environment of lowered T-cell immune activity is hence intriguing. One possible explanation maybe due to the perturbations of the insulin signaling pathway in these women which induced the down expression of the nuclear factor kappa-light-chain-enhancer of activated B cells (NF-κB) and other pro-inflammatory enhancers (calgranulin and ICOS). NF-κB is a protein complex that regulates the expression of a multitude of immune response genes including cytokines, chemokines, antigen presenting cells and adhesion receptors [36]. In response to a variety of stimuli, including cytokines, viral and bacterial pathogens, a latent inactive NF-κB complex is activated in the cytoplasm through phosphorylation and translocates to the nucleus, where it stimulates transcription of genes. Activation of NFκB by the inflammatory cytokine alpha tumor necrosis factor (TNFα) has been shown to involve the phosphotidylinositol pathway together with its downstream targets-Akt, TSC1 and TSC2 through IRS 1 [37,38]. Quantitative real time PCR confirmed low expression in IRS1, PI4K TSC and, NFκB among the HIV resistant female sex workers (Figure 4). We hypothesize that in presence of high CD26/DPPIV, the reduced insulin signaling leads to a low activity of phosphotidylinositol cascade system which in turn resulted in a lower expression of TSC-2, a build-up of the MTORC1 and subsequent repression of NFκB. In-depth functional studies will be required to test this assertion. Our observations suggest that metabolic and signaling pathways that predispose to an impaired glucose fasting state and Type 2 diabetes may be new correlates of HIV resistance. In addition, it underscores the key role of a systems biology in discovery of novel candidate biological markers that may be crucial in the design of effective preventive strategies against HIV/AIDS.

**Methods**

**Study population**

The Nairobi (Pumwani) commercial sex worker cohort was established in 1985 and has provided vital data that there might be biological mediated resistance to HIV infection [11,39]. Despite repeated exposures to HIV-1, a number of women in the cohort have remained HIV uninfected for long periods and have epidemiologically been defined as resistant. The definition used in the current study were female sex workers actively followed in the cohort for more than seven years and persistently seronegative at biannual visits and negative by a sensitive PCR assay for proviral HIV-1 [8]. HIV infected commercial sex workers who were not on antiretroviral treatment (HIV+) and HIV negative antenatal clinic attendees (HIV Neg) were included as controls. Written Informed consent was sought from all volunteers and the study received ethical approval from ethical review boards of University of Nairobi and the Kenya Medical Research Institute.

**Soluble CD26/DPPIV analysis**

Peripheral blood samples were randomly obtained from HIV Resistant and HIV positive female sex workers. Samples were also obtained from a HIV negative
population of female antenatal clinic attendees matched for age (38.5 yrs) and race from the same part of Nairobi. All samples were separated to yield plasma and peripheral mononuclear cells. Soluble CD26/DPP IV plasma levels were quantified by Human DPP IV/CD26 Quantikine ELISA kit (R&D Systems, Minneapolis MN) following the manufacturer’s recommendation. Briefly, standards and samples were diluted to a 1:1000 and ran in triplicate. Optical densities of the standards were used to generate a standard curve to determine sample concentration. Mean values were calculated and tests for significance determined with Graphpad Prism using student t-tests. A total of 300 subjects (100 in each arm) were used for the study.

Cell surface CD26 expression analysis

To determine expression differences at cell surface level (CD26) a 3 color flow-cytometry (FACS) analysis was performed with the use of anti-CD26 antibody. PBMC were stained for CD4, CD8, and DPP4 washed and fixed using 1% paraformaldehyde. FACS data was analyzed using FlowJo (Tree Star, Ashland, OR). Lymphocytes were gated on forward versus side scatter, and then CD4 versus CD8. CD26/DPP IV expression levels were analyzed on CD4+ cells. Gates were set on the basis on single stained samples that did not contain CD26/ DPP IV and used to calculate expression and mean fluorescence intensity levels. Significant values were determined in Prism using the Mann Whitney test.

Investigation for clinical type 2 diabetes

To investigate clinical type 2 diabetes, commercial sex workers who had a significantly elevated soluble CD26/ DPP IV plasma concentration were re-assessed for blood sugar levels. Identified volunteers were requested to suspend sex work for at least 24 hours and undergo an overnight fast. Blood samples were obtained before breakfast and analyzed for blood sugar using the glucose oxidase test system (Accu-check, Roche, Basel). Volunteers with blood glucose levels above 125 mg/dl (6.9 mmol/l) were considered diabetic and those between 100-125 mg/dl considered as having impaired fasting glucose state as per the ADA definition criteria [18].

Expression patterns of genes in the insulin/type 2 diabetes signaling pathway

Quantitative PCR analysis of selected genes in the insulin signaling pathway from a representative sample of HIV-R and HIV-Neg controls were done using Biosystems 7500 Real-Time PCR System (ABI, Foster City, CA). Assay protocols followed the Quantitect SYBR Green RT PCR a kit (Qiagen GmbH, Hilden D2Z). Briefly 500 ng of total RNA extracted was reverse transcribed into cDNA using superscript II reverse transcript.

Aliquots of equal amounts of cDNA from each sample were ran in duplicate and normalized using an internal control gene 18s RNA. Relative expression of each gene was determined from a standard curve generated from a pool of cDNA from the same samples. P-values for significance were analyzed using student t-test.

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Authors’ contributions

EMS conceptualized, designed, supervised the study and did the manuscript write-up. BO, LM, MR, WA, EM and AV supported the laboratory analysis. JK and CW were project physicians involved in day to day sample collection, treatment and follow-ups. BTR, ML, FP and SM were project advisors and contributed to the manuscript write-up. All authors have read and approved the final manuscript.

Competing interests

The authors declare that they have no competing interests.

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