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Dual Molecular Mechanisms Govern Escape at Immunodominant HLA A2-Restricted HIV Epitope

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Serial accumulation of mutations to fixation in the SLYNTVATL (SL9) immunodominant, HIV p17 Gag-derived, HLA A2-restricted cytotoxic T lymphocyte epitope produce the SLEFTIAVL triple mutant “ultimate” escape variant. These mutations in solvent-exposed residues are believed to interfere with TCR recognition, although confirmation has awaited structural verification. Here, we solved a TCR co-complex structure with SL9 and the triple escape mutant to determine the mechanism of immune escape in this eminent system. We show that, in contrast to prevailing hypotheses, the main TCR contact residue is 4N and the dominant mechanism of escape is not via lack of TCR engagement. Instead, mutation of solvent-exposed residues in the peptide destabilise the peptide–HLA and reduce peptide density at the cell surface. These results highlight the extraordinary lengths that HIV employs to evade detection by high-affinity TCRs with a broad peptide-binding footprint and necessitate re-evaluation of this exemplar model of HIV TCR escape.

Keywords: T-cell, T-cell receptor, HIV, immune escape, MHC

INTRODUCTION

CD8+ cytotoxic T lymphocytes (CTLs) form a critical component in immune control of HIV infection. However, the virus has adopted numerous strategies to evade detection by these important immune cells (1, 2). Notably, the high rate and poor fidelity of replication produces a viral quasispecies to allow rapid selection of sequence variants that evade CTLs. This immune escape mutation is often associated with loss of viral fitness (3). The effects of immune escape from some HLA class I alleles, particularly HLAs B27 and B57 (3, 4), can be significant and delay progression to AIDS. There has been considerable work on immune escape from HIV epitopes presented by the most frequent HLA worldwide, HLA A2 (5). The majority of these studies have focussed on the immunodominant response through the p17 HIV Gag-derived epitope SL9 (amino acids 77–85, sequence SLYNTVATL) (6–23). Responses to SL9 are uncommon in acute infection but are dominant during the long, asymptomatic, chronic infection period where over 70% of HLA A2+ HIV-infected individuals mount a strong response to this epitope (13, 18, 19). Immune escape from SL9-specific CTL during this period has been studied extensively and is known to follow well-described patterns (6, 7). The leucine residues at positions 2 and 9 in SL9 are located within an α-helix (21) and are critical for trimerisation of the P17 Gag protein (24).
Mutations within these residues are believed to produce highly compromised virus (11, 24). These residues also act as the principal anchors for securing SL9 into the HLA A2-binding groove (22) and ensure that the virus does not use the archetypical escape strategy of mutating primary MHC anchors to block epitope presentation at the infected cell surface (25–27). Instead, in the face of SL9-specific CTL pressure, HIV has been observed to undergo preferred mutation at positions 3, 6, and 8 in the epitope (6, 7, 9). The dominant mutations seen at these positions (Y3F, V6I, and T8V) correlate with a lower viral load escape mutant in SL9 (6).

Here we use structure and protein biophysics to deconstruct how HIV escapes from the well-studied (9–11) SL9-specific 868 TCR. We add to the existing data by providing the first ever TCR–pMHC structures in this exemplar model of how HIV evades engagement by host TCRs. We find that the 868 TCR interacts with HLA A2–SLYNTVATL (A2–peptide from here on in) with the expected binding topography but exhibits an unusual degree of structural rearrangement upon interaction with A2–SLYNTVATL. The 868 TCR also contacts all five central residues in SL9 (positions 3–8) and binds with an unprecedentedly high affinity for a natural TCR. Surprisingly, 4N is the major contact residue—not the 3Y, 6I, or 8T that mutate to precipitate immune escape. This enabled the generation of a high-resolution structure of the 868 TCR in complex with the SLFNTIAYL ultimate escape variant. These results challenge current thinking on the mechanism by which HIV escapes at SL9 and demonstrate the extraordinary tactics that the virus can employ to escape from CTL responses that can involve TCRs with a very broad footprint and a very high natural binding affinity.

**MATERIALS AND METHODS**

**Construct Design**

The 868 TCR α and β chains, HLA A2 heavy chain, and β2m chain were generated by PCR mutagenesis and cloning. All sequences were confirmed by automated DNA sequencing (Lark Genetics). TCR expression constructs were designed with a disulfide linked construct to produce the soluble domains (variable and constant) for both the α (residues 1–207) and β chains (residues 1–247) (28, 29). The HLA A2 heavy chain (residues 1–248) (α1, α2, and α3 domains), tagged, or not tagged with a biotinylation sequence, and β2m (residues 1–100) were also cloned and used to make the pMHCI complexes. The TCR α and β chains, the HLA A2 α chain, and β2m sequences were inserted into separate pGMT7 expression plasmids under the control of the T7 promoter (30).

**Protein Expression, Refolding, and Purification**

Competent Rosetta DE3 E. coli cells were used to produce the TCR α and β chains, HLA A2 heavy chain, and β2m in the form of inclusion bodies using 0.5 mM IPTG to induce expression and protein were chemically refolded as described previously (31).

**pMHCI Biotinylation**

Biotinylated pMHCI was prepared as previously described (32).

**Surface Plasmon Resonance (SPR) Experiments**

Surface plasmon resonance equilibrium binding analysis was performed using a BIAcore T100™ equipped with a CM5 sensor chip as previously reported (32, 33). HLA DR1, generated as in Ref. (34), was used as a negative control on flow cell 1. SPR kinetic analyses were carried out to determine the KD values for the TCR, at 25°C. For all kinetic experiments, approximately 300 RU of pMHC was coupled to the CM5 sensor chip surface. The TCR was then injected at concentrations ranging from 10 times above and 10 times below the known KD of the interaction at 45 µl/min. The KD values were calculated assuming 1:1 Langmuir binding [AB = B × ABMAX/(KD + B)], and the data were analysed using a global fit algorithm (BIAevaluation™ 3.1). The koff, koff/KD and KD values were calculated by global fitting of the data using BIAevaluation™ 3.1 software. For the thermodynamics experiments we used the KD determined by SPR at different temperatures with the standard thermodynamic equation \( ΔG = RT \ln K_D \) and the standard non-linear van’t Hoff equation \( \Delta G^° = \Delta H^° - T\Delta S^° + \Delta C_p^°(T^° - T)^{-} - T\Delta C_p^° \ln (T/T^°) \) with \( T_0 = 298 \) K. For stability experiments, single injections of 50 µM 868 TCR over the SL9 variants were performed after different incubation times at 37°C, and the RU values were recorded.

**Isothermal Titration Calorimetry (ITC)**

Isothermal titration calorimetry experiments were performed using a Microcal VP-ITC (GE Healthcare) as previously described (35), with 30 µM phHLA-I in the calorimeter cell and 210 µM soluble 868 TCR in the syringe. Buffer conditions were 20 mM Heps (pH 7.4) containing 150 mM NaCl and 20 injections of 2 µl volume each were performed. Results were processed and integrated with the Origin 6.0™ software distributed with the instrument. ITC experiments were performed in duplicate.

**Adhesion Frequency Assay**

We used an adhesion frequency assay to measure the 2D affinity of TCR–pMHC interactions at the cell membrane as previously described (36). Briefly, human T-cells transfected with the 868 TCR were mounted onto one micro-pipette and, on the other pipette, human red blood cells coated with pMHC by biotin–streptavidin coupling served as both a surrogate APC and an adhesion sensor for detecting the TCR–pMHC interaction. Site
densities of TCR and pMHC were measured by flow cytometry as previously described (37). All assays were performed using at least 5 cell pairs, and calculated as an average of 100 cell–cell contacts.

Crystallisation, Diffraction Data Collection, and Model Refinement

All protein crystals were grown at 18°C by vapour diffusion via the sitting drop technique. 200 nl of 1:1 molar ratio TCR and pMHC (10 mg/ml) in crystallisation buffer (10 mM Tris pH 8.1 and 10 mM NaCl) was added to 200 nl of reservoir solution. 868 crystals were grown in TOPS (38) in 0.1 M sodium cacodylate pH 6.5, 20% PEG 8000, and 0.2 M ammonium sulphate (38). 868-A2–SLYNTVATL crystals were grown in TOPS in 0.1 M sodium cacodylate pH 6.0, 15% PEG 4000 and 0.2 M ammonium sulphate (38). 868-A2–SLYNTI ATL co-crystals were grown in TOPS in 0.1 M sodium cacodylate pH 6.0, 15% PEG 4000 and 0.2 M ammonium sulphate (38). 868-A2–SLYNTI ATL crystals were grown in TOPS1 in 0.1 M sodium cacodylate pH 5.5, 15% PEG 4000 and 0.2 M ammonium sulphate (38). A2–SLYNTI ATL crystals were grown in TOPS in 0.1 M sodium cacodylate pH 6.0, 25% PEG 4000 and 0.2 M ammonium sulphate (38). A2–SLYNTI ATL crystals were grown in TOPS in 0.1 M sodium cacodylate pH 6.0, 25% PEG 4000 and 0.2 M ammonium sulphate (38). All crystals were soaked in 30% ethylene glycol before cryo-cooling. All crystallisation screens and optimisation experiments were completed using an Art-Robbins Phoenix dispensing robot (Alpha Biotech Ltd, UK). Data were collected at 100 K at the Diamond Light Source, Oxfordshire. All datasets were collected at a wavelength of ~0.98 Å using an ADSC Q315 CCD or PILATUS Pixel detectors. Reflection intensities were estimated with the XIA2 package (39) and the data were scaled, reduced and analysed with SCALA and the CCP4 package (40). Structures were solved with molecular replacement using PHASER (41). Sequences were adjusted with COOT (42) and the models refined with REFMAC5. Graphical representations were prepared with PYMOL (43). The reflection data and final model coordinates were deposited with the PDB database (868, PDB: 5NMD; 868-A2–SLYNTVATL, PDB: 5NME; 868-A2–SLYNTI ATL, PDB: 5NMF; 868-A2–SLYNTI ATL, PDB: 5NMG; A2–SLYNTI ATL, PDB: 5NMH; A2–SLYNTI ATL, PDB: 5NMK).

Thermal Stability CD Analysis of HLA A2 Complexes

Thermal stability of HLA A2/β2m/peptide complexes was determined by circular dichroism spectroscopy following the change of ellipticities θ at 218 nm using an Aviv 215 instrument (Aviv Biomedical Inc., Lakewood, NJ, USA). Proteins were dissolved in 137 mM NaCl, 3 mM KCl, 8 mM Na2HPO4 and 1 mM KH2PO4, pH 7.4, at concentrations of ~3 µM as determined spectroscopically using calculated extinction coefficients (44). Melting curves were recorded from 4°C up to a maximum temperature when protein aggregation was observed using a heating rate of ~0.5°C/min. Melting curves were analysed assuming a two-state trimer-to-monomer transition from the native (N) to unfolded (U) conformation N ↔ 3U with an equilibrium constant K = [U]/[N] = F/[c 2 (1 – F)], where F and c are the degree of folding and protein concentration, respectively. Data were fitted as described (45) using the non-linear least-squares routine of Origin V7.5 (OriginLab, Northampton, MA, USA). Fitted parameters were the melting temperature Tm, van’t Hoff’s enthalpy ΔHnf, at the transition midpoint, and the slope and intercept of the native baseline assumed as a linear function of the temperature. As protein complexes aggregated at various degrees of unfolding, the ellipticity of the unfolded state was set as a constant of ~4,500 deg cm2 dmol−1 (46).

Generation of 868 TCR Transgenic T-Cells

868 TCR construct (codon-optimised for human expression) was cloned into the third-generation lentiviral expression vector pELNS (kindly provided by James R. Riley, University of Pennsylvania, PA, USA). pELNS vector was engineered to contain rat CD2 (rCD2) gene to be used as a marker of lentiviral transduction of target cells. 868 TCR-α and TCR-β, and rCD2 were separated by a pair of “self-cleaving” 2A sequences (11). Lentiviral particles were prepared by calcium phosphate transfection of packaging plasmids (pRSV-Rev, pMDLg/pRRE, pMD2.G) together with 868-encoding pELNS into HEK293T cells. Supernatant was harvested 48- and 72-h post transfection and was subsequently concentrated by ultracentrifugation. Primary CD8+ T-cells were obtained from healthy donor blood bags (Welsh Blood Service) by density gradient centrifugation and positive selection using CD8 MicroBeads (Miltenyi Biotech, Bergisch Gladbach, Germany). Isolated T-cells were stimulated overnight with Gibco human T-activator CD3/CD28 Dynabeads (ThermoFisher Scientific, Waltham, MA, USA) at 3:1 bead to T-cell ratio, and transduced with concentrated lentiviral stock in presence of 5 µg/ml polybrene (Santa Cruz Biotechnology, Santa Cruz, CA, USA). After 72 h rCD2+ cells were enriched by magnetic bead selection to yield >95% purity. For tetramer staining experiments, CD8+ T-cells were used without magnetic enrichment to leave a non-transduced internal control cell population. Similar methodology was used to transduce TCRβ chain negative Jurkat cells to allow the examination of tetramer binding in the absence of the CD8 glycoprotein.

Tissue Culture

T2 cells were maintained as suspension cells in R10 (RPMI 1640 media, 10% FBS, penicillin, streptomycin, and l-glutamine, all Life Technologies, Paisley, UK) and passaged weekly or when required. Post enrichment, primary CD8+ T-cells were routinely expanded with allogeneic feeder PBMCs and phytohaemagglutinin (PHA) in T-cell media as previously described (47). 868 TCR expression was checked prior to assay using anti-rCD2 antibody and A2-SLYNTVATL tetramer, with the protein kinase inhibitor Dasatinib (48) (Axon Medchem, Reston, VA, USA), and unconjugated anti-PE antibody (PE001, BioLegend, London, UK) used to enhance tetramer staining, as previously described (47, 49).
T-Cell Activation Assays

For peptide pulsing assays, T2 cells were incubated with peptide for 1 h in 0.5 ml of AIM-V serum free media (ThermoFisher Scientific) in 5 ml FACS tubes at 37°C and 5% CO₂, followed by 4 washes with 4 ml of RPMI 1640 media. In assays using primary CD4⁺ T-cells as targets, these cells were purified from PBMC using magnetic microbead separation (Miltenyi Biotech). Peptides were synthesised to >95% purity, stored as 20 mM DMSO stocks at −80°C and 1 mM RPMI dilutions made on the day of assay. DMSO controls were treated the same as peptide pulsed. Post pulsing, T2 cells or CD4⁺ T-cells were incubated in 4.5 ml of AIM-V media at 37°C and 5% CO₂ for 3–24 h or used immediately (0 h), such that cells were ready at the same time. 868 TCR transduced and non-transduced T-cells (30,000) were rested overnight in R5 (as for R10 but with 5% fetal bovine serum) then co-incubated in 96 U-well plates with T2 or CD4⁺ T-cells (60,000) in the presence of GolgiStop and GolgiPlug according to the manufacturers’ recommendations (BD Biosciences, Oxford, UK), and anti-CD107a PE Antibody (H483, BD Biosciences). Controls included T-cells alone and T-cells with unpulsed T2 cells (negative control) or PHA. After 3.5–5 h of incubation, cells were labelled with the violet LIVE/DEAD fixable dead cell stain Vivid, anti-CD8 APC (BW135/80; Miltenyi Biotech) and anti-CD19 pacific-blue (T2 cells) (BioLegend), and rCD2 (CD4 assay) and anti-CD4 APC-Vio770 (M-T466, Miltenyi Biotech; CD4 assay) antibodies. Cells were then stained with anti-TNFα PE-Vio770 (BD Biosciences) for 10 min at RT, washed then stained with anti-HLA-A2 antibody (BB7.2 (ThermoFisher Scientific) in 5 ml FACS tubes at 37°C and 5% CO₂, followed by 1 h in 0.5 ml of AIM-V serum free media then co-incubated in 96 U-well plates with T2 or CD4⁺ T-cells (60,000) and incubated in 4.5 ml of AIM-V for the specified times (h) at 37°C and 5% CO₂, dilutions made on the day of assay, to allow peptide to dissociate, then stained and fixed as before. Flow cytometry was performed on a FACSCanto II (BD Biosciences) and data analysed with FlowJo software (TreeStar).

Peptides Used in This Study

This study made use of the HIV SL9 peptide and variants thereof. The sequences and origin of these peptides, and other control peptides utilised, are listed in Table 1.

RESULTS

868 TCR Binding to A2–SLYNTVATL Involves Extensive Structural Reorganisation

Previous studies have demonstrated that over 1.7% of CD8⁺ T-cells from HIV-infected patient 868, stained with A2–SLYNTVATL tetramer (10). 75% of these tetramer⁺ cells expressed a TRBV5-6-β chain was observed in 10/18 transcripts sequenced in this study (CDR3 sequence CASSLSAVQNNEQF) (10). This same β-chain was later shown to be present in all A2–SLYNTVATL tetramer⁺ cells in a T-cell line grown from patient 868 in 1996 (9). TCR chain antibody staining showed that all of these A2–SLYNTVATL tetramer⁺ T-cells co-expressed a TRAV12-2 chain (11). Our previous study confirmed that CD8⁺ T-cells expressing this TCR were unable to control HIV virus in vitro (11). Nevertheless, we were able to use phage display to select an artificially enhanced version of the 868 TCR that could recognise the SLEN'TIAYL ultimate escape variant; thereby engineering foresight into an immune receptor (11). The detection of common SL9 escape mutants on the surface of HIV-infected HLA A2⁺ T-cells by CD8⁺ T-cells expressing engineered, but not wild-type, 868 TCR reinforced earlier studies suggesting that these mutants are presented by HLA A2 and that escape from the wild-type epitope is mediated by loss of TCR binding (so-called “TCR escape”) (6, 7, 11, 18). To understand how positions 3, 6 and 8 in SL9 impinge on TCR binding, we solved the structure of the wild-type 868 TCR in complex with A2–SLYNTVATL to 2.9 Å resolution (Table 2). The 868 TCR bound to A2–SLYNTVATL with a conventional, centrally-located, diagonal orientation with the TCR α-chain positioned over the α helix of MHC class I and the TCR β-chain positioned over the α1 helix (Figure 1A). The TCR crossing angle of 51.6° lies within the normal range for human TCR–pMHC complexes (50), enabling positioning of the TCR CDR3 loops centrally over the peptide, and the CDR1 and CDR2 loops predominantly over the MHC helices (Figure 1B). To gain extra insight into the mechanism by which the 868 TCR adopted this

| Peptide sequence | Abbreviation | HLA-restriction | Parent protein |
|------------------|--------------|----------------|---------------|
| SLYNTVATL        | SL9          | HLA A*0201     | HIV p17 Gag   |
| SLNTVATL         | 3F           | HLA A*0201     | HIV p17 Gag   |
| SLYNTVATL        | 6V           | HLA A*0201     | HIV p17 Gag   |
| SLNIVTVATL       | 8V           | HLA A*0201     | HIV p17 Gag   |
| SLYNTVVTL        | 3F688V       | HLA A*0201     | HIV p17 Gag   |
| GILGTVSTL        | A2-4u        | HLA A*0201     | Influenza matrix |
| HPVGEADYFEY      | B*3501       |                | EBV EBNA-1    |

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November 2017 | Volume 8 | Article 1503

4
TABLE 2 | Data collection and refinement statistics for unligated 868 TCR and complex structures.

|                      | 868 TCR | 868-A2–SLYNTVATL | 868-A2–SLYNTIAVL | 868-A2–SLFNIAVL | A2–SLYNTIAVL | A2–SLFNIAVL |
|----------------------|---------|------------------|------------------|------------------|--------------|--------------|
| **Data collection**  |         |                  |                  |                  |              |              |
| Space group          | P 1 21 1| P 21 2 1        | P 21 2 1        | P 21 2 1        | P 1 21 1     | P 1 21 1     |
| **Cell dimensions**  |         |                  |                  |                  |              |              |
| a, b, c (Å)          | 87.5, 50.6, 114.3 | 211.1, 85.1, 113.2 | 207.6, 84.7, 112.1 | 209.36, 85.11, 113.15 | 56.1, 80.1, 57.6 | 55.84, 79.03, 58.37 |
| α, β, γ (°)          | 90, 90.2, 90 | 90, 90, 90      | 90, 90, 90      | 90, 90, 90      | 90, 90, 90   | 90, 90, 90, 90 |
| Resolution (Å)       | 43.8–2.1 (2.13–2.07) | 64.8–2.9 (3.01–2.94) | 34.0–2.9 (2.96–2.89) | 99.5–2.8 (2.82–2.75) | 31.7–1.6 (1.59–1.55) | 48.4–1.7 (1.70–1.66) |
| Beam line            | I24     | 103              | 103              | 104–1           | 103          | 103          |
| Beam time code       | 6232–9  | 4532–1           | 6232–1           | 6232–10         | 4532–1       | 3262–9       |
| Rmerge (%)           | 5.0 (72.3) | 12.5 (95.9)     | 15.1 (84.0)     | 11.7 (86.7)    | 6.6 (73.3)   | 7.4 (103.3)  |
| Rfree (%)            | 98.6 (99.6) | 99.1 (99.8)     | 100 (100)       | 99.9 (100)     | 99.4 (97.7)  | 98.9 (99.8)  |
| Completeness (%)     | 3.5 (3.8) | 7.5 (7.1)       | 7.9 (8.2)       | 7.4 (7.8)      | 4.1 (4.0)    | 3.8 (3.7)    |
| No. reflections      | 60,507 (4,492) | 44,013 (3,213) | 44,013 (3,284) | 53,560 (3,895) | 66,024 (4,752) | 53,691 (4,000) |
| **Refinement**       |         |                  |                  |                  |              |              |
| No free reflections  | 3,051   | 2,212            | 2,223            | 2,717           | 3,346        | 2,724        |
| Rfree/Rmerge (%)     | 18.9/22.3 | 19.4/28.9       | 18.8/27.5        | 19.2/27.0       | 14.4/19.4    | 17.5/20.8    |
| **R.m.s. deviations**|         |                  |                  |                  |              |              |
| Bond lengths (Å)     | 0.0189  | 0.015           | 0.012           | 0.017           | 0.025        | 0.017        |
| Bond angles (°)      | 1.970   | 1.665           | 1.483           | 1.954           | 2.107        | 1.832        |
| Mean B value (Å²)    | 57.0    | 69.0            | 53.7            | 31.2            | 16.8         | 24.3         |
| Wilson B-fac (Å²)    | 40.7    | 72.2            | 60.3            | 58.9            | 17.2         | 22.5         |
| ML estimated         | 0.144   | 0.387           | 0.385           | 0.300           | 0.049        | 0.070        |
| coordinate error (Å) | 0.144   | 0.387           | 0.385           | 0.300           | 0.049        | 0.070        |

One crystal was used for solving each structure. Figures in brackets refer to the highest resolution bin.

In summary, the 868 TCR utilised an induced fit, or conformation selection, binding mode to form a pincer-like interaction around the central solvent-exposed portion of the peptide. Additional contacts with distal peptide residues enabled a relatively extensive and broad binding interface between the TCR and pMHC.

TCR Structure with the Ultimate SL9 Escape Variant

The 868–A2–SLYNTVATL complex structure demonstrated that the dominant contacts with the SL9 peptide were with the 4N residue rather than with the 3Y, 6V, or 8T residues that result in immune escape when substituted to create the SLFNIAVL ultimate escape variant (6, 7). Alanine scan mutation across the SL9 peptide backbone showed an almost complete loss of recognition with the L2A and Y3A changes. Alanine substitution at positions 4, 5, and 6 had the next greatest effect whereas peptides substituted at positions 1, 8, and 9 exhibited recognition equivalent to the wild-type peptide (Figure 2E). In order to gain insight as to how natural escape mutations impinge on TCR binding, we also solved the structure of the unbound 868 TCR with the Ultimate sl9 escape variant present in the patient at the time the blood sample was used to grow the 868T-cell clone (9), and the SLFNIAVL ultimate escape variant (Table 2). Comparison of these structures with the 868 TCR in complex with the A2–SLYNTVATL index sequence showed that the 868 TCR engaged all the antigens in
Figure 1 | Structural analysis of the 868-A2–SL9-binding mode. (A) The overall binding mode of the 868 TCR (red and magenta cartoon) interacting with HLA A2 (grey surface) and the SLYNTVATL peptide (red sticks and surface). (B) Surface representation of the A2–SLYNTVATL complex looking down at the peptide. Positions of the 868 TCR CDR loops (1α—red, 2α—green, 3α—blue, 1β—yellow, 2β—cyan, 3β—orange) and the crossing angle of the TCR with the MHC groove is shown. (C) The positions of the 868 TCR CDR3 loops differ considerably between the ligated (red cartoon) and unligated (grey cartoon) structures.

Previous studies comparing SLYNTVATL and the SLFN'TIAVL mutated peptide showed that they adopted different conformations within the HLA A2 binding groove (7, 22). The T-cell recognition and on-rate kinetics with the G10 SL9-specific TCR in this study were very similar with A2–SLYNTVATL and A2–SLFN'TIAVL, leading the authors to conclude that one conformation, the 868 TCR could tolerate mutations at residues 3, 6, and 8 in SL9 (Table 3). Thus, because our structural data were inconsistent with the hypothesis that SLFN'TIAVL primarily affects TCR binding, and because sum contact numbers do not necessarily represent the accurate binding energy between two molecules, we undertook further structural and biophysical experimentation to determine the mechanism that results in poor T-cell recognition of SLFN'TIAVL.

A2–SLYNTVATL Does Not Alter Conformation When Bound to 868 TCR

Previous studies comparing SLYNTVATL and the SLFN'TIAVL mutated peptide showed that they adopted different conformations within the HLA A2 binding groove (7, 22). The T-cell recognition and on-rate kinetics with the G10 SL9-specific TCR in this study were very similar with A2–SLYNTVATL and A2–SLFN'TIAVL, leading the authors to conclude that one conformation,
common to both peptides, was recognised by TCRs (22) and suggesting that TCR interactions with either A2–SLYNTVATL or A2–SLFNTVATL required an induced fit in the pMHC. Thus, SL9-derived peptides may bind to MHC dynamically and undergo conformational changes during TCR docking. Conformational peptide changes would likely elicit an entropic penalty during TCR ligation, not revealed by differences in contacts, and could contribute to weaker binding. However, based on previously solved structures [A2–SLYNTVATL (22), A2–SLFNTVATL (22), A2–SLANTVATL (21), A2–SLYNTIAL (21), A2–SLYNTVATL (21), A2–SLYNVATL (21), A2–SLYNTIAAL (21), and A2–SLFNTIAL (21)], some of which we replicated in this study, we show that A2–SLYNTVATL, A2–SLYNTIAL, and A2–SLFNTIAL all adopt a similar structure to every unbound SL9 peptide variant, apart from A2–SLFNTVATL (22), when ligated to the 868 TCR (Figure 4). These data make it possible to be more definitive about the direction of travel of the previously suggested induced fit model (22) (i.e., the odd one out is A2–SLFNTVATL and this peptide presumably adopts the more conventional structure for A2–SLYNTVATL type ligands prior to, or at the time

**FIGURE 2** The 868 TCR focuses on residues 4N and 5T in the peptide. Hydrogen bonds show as red dotted lines, van der Waal's contacts shown as black dotted lines in panels (A–D). (A) The 868 TCR (CDR3α in blue, CDR1α red and CDR3β orange sticks) binding to the solvent-exposed peptide residues 4N and 5T in the peptide (red sticks). (B) The 868 TCR (CDR3α in blue, CDR1α red sticks and surface) forms a tight pocket around peptide residue 4N (red sticks) during binding. (C,D) The restriction triad residues 72Q, 69A, and 155Q (grey sticks) play a central role in 868 TCR–MHC-mediated interactions (TCR CDR3α residues shown in blue, CDR2β in cyan, and CDR1α in green sticks). (E) 868 TCR transduced CD8+ T-cells from two HLA A2+ donors were co-incubated with T2 cells and alanine-substituted peptides of SLYNTVATL (alanine already at position 7) overnight and supernatants used for MIP-1β ELISA. Non-linear regression curves and $pEC_{50}$ for each peptide are displayed.
of TCR binding). Furthermore, these data do not support the notion that HIV escape is mediated by dynamic differences in ligand engagement during TCR binding to SL9 variants.

The 868 TCR Adopts a Similar Energetic Strategy during Binding to SL9 Escape Variants

We next explored the evidence for HIV escape being mediated by distinct TCR-binding parameters. It has been previously shown that the G10 SL9-specific TCR binds with a similar affinity to both A2–SLYNTVAL (K_D = 2.2 μM) and A2–SLẸNTYATŁ (K_D = 5.2 μM) by SPR (22). Another study demonstrated that half-maximal pMHC tetramer binding (D_Ro) to the SL9-specific D3 TCR was near identical for A2–SLYNTVAL, A2–SLĘNTYATŁ, A2–SLYNTVAL, A2–SLẸNTYATŁ, and A2–SLẸNTIŁAYL (21). Finally, we have previously shown that the 868 TCR binds to a range of SL9 variants (11) (A2–SLYNTVAL, A2–SLĘNTIŁATŁ, A2–SLĘNTIŁATŁ, A2–SLĘNTYATŁ, A2–SLĘNTYATŁ, and A2–SLĘNTIŁAYL). Consequently, the available data show that SL9-specific TCRs can engage SL9 variants within an affinity range that would be expected to easily activate T-cells. Here, we reproduced SPR-binding assays with the 868 TCR for the three ligands included in this study. These data demonstrated that the 868 TCR bound to A2–SLYNTVAL with a K_D of 82 nM (Figure 5A) confirming that this is, by far, the strongest binding natural TCR–peptide–MHC interaction ever recorded (51, 52). Indeed, the affinity was even greater with the A2–SLĘNTIŁATŁ variant (K_D = 38 nM; Figure 5B) that dominated in the patient at the time the blood sample was taken that was used to generate the 868 T-cell line (9, 10). 868 TCR interactions with A2–SLYNTVAL and the A2–SLĘNTIŁATŁ variant were characterised by a relatively slow off-rate (k_off = 1.6 × 10^{-2}–7.3 × 10^{-3} s^{-1}) compared to the majority of other natural TCR–pMHC interactions (average k_off ~ 2 × 10^{-1} s^{-1}) (51). Although the strength of the 868-A2–SLẸNTIŁAYL interaction was 21-fold and 45-fold weaker than for A2–SLYNTVAL or A2–SLĘNTIŁATŁ, respectively, due to a much faster off-rate (k_off = 2.1 × 10^{-1} s^{-1}), the affinity (K_D of 1.77 μM; Figure 5C) was still in the range of the very best natural anti-pathogen TCRs (52, 53). Thus, this weaker affinity alone was unlikely to explain the ability of the SLẸNTIŁAYL variant to evade T-cell detection. Indeed, we have described a functional HLA A2-restricted TCR that binds its cognate pMHC with over 100-fold weaker affinity than the 868 TCR and A2–SLẸNTIŁAYL (54, 55). To get a more complete picture of how the 868 TCR engages its cognate ligands we also undertook thermodynamic analyses by SPR (Figures 5D–F; Table 4) and directly by isothermal calorimetry (ITC; Figures 5G–I). The results of these independent techniques correlated well and showed that 868 TCR engagement was driven by energetically favourable enthalpy and entropy changes to all three ligands. Thus, for the 868 TCR, there was a net increase in the number of electrostatic interactions and a transition from order–disorder during binding. Considering that the 868 TCR CDR3 loops underwent similarly large conformational changes during engagement to all three SL9 variants studied here (shown for 868-A2–SLY interaction in Figure 1C), a movement that would probably be entropically unfavourable, the expulsion of ordered water molecules from the interface, and/or order–disorder transitions in other parts of the molecules presumably contributed to the net favourable entropic change of the interaction. The lower affinity of 868 TCR for A2–SLẸNTIŁAYL compared to A2–SLYNTVAL or A2–SLĘNTIŁATŁ was associated with a much reduced enthalpic component, but more favourable entropy. We used 2D binding analysis to confirm our observations using SPR (Figures 5J,K). In accordance with our previous findings with other TCRs (55, 56), we observed an identical relationship between the strength of interaction between 868 and the three SL9-derived ligands by 2D analysis and SPR. 868-A2–SLĘNTIŁATŁ was characterised by the strongest 2D effective affinity (1.6 × 10^{-3} μM), followed by 868-A2–SLYNTVAL (1.0 × 10^{-3} μM), with 868-A2–SLẸNTIŁAYL generating the weakest effective 2D affinity (5.9 × 10^{-4} μM). To further confirm our observations from 2D and 3D binding experiments, we next examined the binding of pMHC tetramers to the 868 TCR at the T-cell surface. Staining of 868 TCR-transduced CD8+ primary T-cells was similar for tetramers made with A2–SLYNTVAL, A2–SLĘNTIŁATŁ, and A2–SLẸNTIŁAYL (Figure 6A). The CD8 coreceptor is known to bind to a site on MHC class I distinct from the TCR docking platform and cooperates with the TCR to aid the engagement of pMHC tetramers (57) and the absence of CD8 binding is known to raise the TCR affinity threshold required for tetramer staining (58, 59). 868 TCR Jurkat T-cells, that do not express CD8, also stained equally well with A2–SLYNTVAL, A2–SLĘNTIŁATŁ, and A2–SLẸNTIŁAYL tetramers (Figure 6B). Given our observations that the TCR affinity threshold required for T-cell activation is considerably lower than that required for pMHC tetramer binding (58) and finding that these reagents can even fail to stain fully functional T-cells (47, 60), our results suggest that the SLẸNTIŁAYL ultimate escape variant would still be a very

TABLE 3 | Summary of co-complex structures of 868-A2–SLYNTVAL, 868-A2–SLĘNTIŁATŁ, and 868-A2–SLẸNTIŁAYL

|          | 868-A2–SLYNTVAL | 868-A2–SLĘNTIŁATŁ | 868-A2–SLẸNTIŁAYL |
|----------|-----------------|------------------|------------------|
| H-bonds (≥3.4 Å) | 20              | 20               | 18               |
| vdWs (≤4 Å)      | 121             | 118              | 128              |
| Total contacts   | 141             | 138              | 146              |
| CDR1/CDR2/CDR3 contacts (≥4 Å) | 13/33/41        | 9/23/46          | 8/22/47          |
| α Chain         | 0/17/37         | 0/27/43          | 0/24/45          |
| β Chain         | 62              | 56               | 63               |
| Peptide contacts | 79              | 82               | 83               |
| Crossing angle (°) | 51.6           | 50.3             | 51.7             |
| Buried surface area (Å²) | 2,396.2        | 2,469            | 2,698            |
| Surface complementarity | 0.50           | 0.51             | 0.55             |
| TCR–MHC          | 0.72            | 0.70             | 0.71             |
| TCR–peptide      | 0.68            | 0.55             | 0.59             |
| TCR–pMHC         | 0.72            | 0.70             | 0.71             |

HB, hydrogen bond; SB, salt bridge; WB, water bridge; vdWs, van der Waals interactions.

A 3.4 Å cut-off was used for HBs, SBs, and WBs, and a 4 Å cut-off was used for vdWs.
efficient T-cell agonist. Indeed, the 868 TCR-transduced primary CD8+ T-cells exhibited similar reactivity to SLYNTVATL and SLFNTIAVL peptides when using standard T-cell activation assay whereby T-cells, antigen-presenting cells, and peptide are added directly to tissue culture wells and allowed to incubate (Figure 6C). We conclude that while there is a substantial
Figure 4 | 868 TCR binding alters the peptide conformation of the A2–SLYNTVATL escape mutants. Structural comparison of the A2–SLYNTVATL escape variants unligated and in complex with the 868 TCR. (A,B) A2–SLYNTVATL in complex with the 868 TCR (red sticks) vs. unligated A2–SLYNTVATL (cyan sticks). Small changes in the positions of the side chains that occur during TCR binding are circled. (C,D) A2–SLYNTIATL (orange sticks) in complex with the 868 TCR vs. unligated A2–SLYNTIATL (yellow sticks). Small changes in the positions of the side chains that occur during TCR binding are circled. (E,F) A2–SLFNTIAVL (green sticks) in complex with the 868 TCR vs. unligated A2–SLFNTIAVL (purple sticks). Small changes in the positions of the side chains that occur during TCR binding are circled.

reduction of TCR binding to the SLFNTIAVL triple mutant sequence, this reduction cannot explain the observed immune escape per se as assumed by all previous studies in this system (6, 7, 11, 23).

The other way by which peptides can escape from host CTL surveillance is by mutating MHC anchors to reduce binding and presentation at the infected cell surface. However, previous studies have shown that the SLFNTIAVL peptide binds to HLA A2 with 50–400% of the affinity the index SL9 sequence depending on the study (Table 5). Such minor relative differences were not thought to be able to explain how the triple mutant peptide escapes from CTL (6, 7, 9, 18). Our own studies with a modified version of the 868 TCR that recognises the escape variants well showed that this TCR could recognise CD4+ T-cells infected with primary HIV isolates carrying known escape variants (11). Thus, these variants must be presented at the cell surface in the context of HLA A2. Importantly, these variants were not recognised by T-cells expressing the wild-type 868 TCR, thus verifying that these viruses are indeed escape mutants (11). Consequently, the established literature indicates that the common variant peptides in this system bind sufficiently well to HLA A2 to be presented (6, 7, 9, 18) and sufficiently well to the 868 and D3 TCRs to be recognised by T-cells expressing these SL9-specific receptors. These apparently incongruous findings prompted us to next re-examine peptide binding in the SL9 system.

CTL Escape by the Ultimate SL9 Escape Variant Mediated by a Fast Peptide Off-Rate

Detailed analysis of the thermal melting of A2–SLYNTVATL, A2–SLYNTIATL, and A2–SLFNTIAVL by circular dichroism showed the melting temperature (T_m) of the wild-type peptide was higher than that of the triple mutant (57°C compared to 49°C; Figures 7A,B) suggesting that the triple mutant is less stable. These data are in accordance with two previous studies showing that SLFNTIAVL binds to HLA A2 with ~50% reduced affinity compared to SL9 (9, 18) but at odds with two studies indicating that SLFNTIAVL is the better ligand (6, 7). Peptide binding to HLA A2 has not previously been thought to result in the immune escape observed with this epitope, which instead has been attributed to a lack of recognition by SL9-specific TCRs (6, 7, 11, 23). Stabilisation of HLA A2 at the T2 cell surface in
the continuous presence of 100 µM peptide (steady-state binding; Figure 7C) showed that the SLFNTIAYL peptide stabilised HLA A2 with 89% of that seen with the wild-type SL9 peptide after 3 h. This relatively minor difference increased in an overnight assay where stabilisation with SLFNTIAYL was ~65% of that seen with the wild-type peptide. Our previous examinations of HLA A2 peptide stability by SPR have shown that good agonists produce pMHC with a half-life of >6 h in a cell free system (61). Similar experiments, performed by examining the relative binding of 868 TCR to A2–peptide by SPR over a time course at 37°C, showed that both A2–SLYNTVATL and A2–SLYNTIATL had a half-life of >7 h. By contrast, the half-life of A2–SLFNTIAYL on the same chip was <2 h (Figure 8A). We thus reasoned that the stabilisation of peptide–HLA measured in a short-term binding assay like those previously used in this system (9, 18) would be determined more by peptide on-rate than the peptide off-rate or dwell time. To further test this possibility, we examined the “on rate” of peptide in T2 cell stabilisation assays. Relative stabilisation was similar for the SLYNTVATL, SLFNTIAYL and the GILGFVFTL influenza matrix-derived peptide that is often used as a positive control for HLA A2 binding (Figure 8B: left panel).

We also examined the stability of these A2–peptide complexes at the cell surface by monitoring the amount of HLA A2 on T2 cells incubated with each peptide prior to washing and further culture.

![Graphs showing peptide binding and stability](image-url)
The vast majority of HLA A2 stabilised with GILGFVFTL or SLYNTVATL peptide was still present at the cell surface 3 h after the peptide was removed from the assay by cell washing. In stark contrast, most of the HLA A2 stabilised by the SL\textsuperscript{ENTIAVL} peptide had decayed in half this time (Figure 8B: right panel).

Collectively, these binding data explain how previous studies, using short-term binding assays in the continuous presence of peptide, have shown minor differences in the binding of SL\textsuperscript{9} and SL\textsuperscript{ENTIAVL} escape variant (Table 5). They also suggest that the dominant mechanism by which the SL\textsuperscript{ENTIAVL} mutant allows immune escape is due to this peptide producing a much shorter-lived peptide–HLA complex. This reduced half-life of antigen at the infected cell surface would be expected to considerably reduce the antigen density on infected cells and thereby facilitate immune escape. To test this notion, we next examined how long peptide pulsed cells could remain T-cell targets for CD\textsuperscript{8\textsuperscript{+}} T-cells transduced with the wild-type 868 TCR. These revealing experiments allowed examination of the combined effects of differences in TCR binding and HLA stability in a relevant biological context. Cells pulsed with the SLYNTVATL remained effective targets for 868 TCR-expressing CD\textsuperscript{8\textsuperscript{+}} T-cells after 24 h of culture (Figure 8C; Figure S1 in Supplementary Material). By contrast, targets pulsed with the SL\textsuperscript{ENTIAVL}, SL\textsuperscript{ENTIAVL}, or SLYNTVATL variant peptides showed reduced levels of recognition after 24 h. The SL\textsuperscript{ENTIAVL} triple mutant sequence produced the largest effect and was recognised well if experiments were performed immediately but recognition dropped to less than 20% of that seen for the SLYNTVATL peptide at t\textsubscript{0}, after targets were cultured for 24 h prior to the T-cell recognition assay. Similar data were observed when primary CD\textsuperscript{4\textsuperscript{+}} T-cells, which form the major reservoir for HIV virus \textit{in vivo}, were used as targets (Figure 8D). Relative to SLYNTVATL, the SL\textsuperscript{ENTIAVL} pulsed cells made
FIGURE 6 | 868 TCR-expressing T-cells bind escape mutant tetramers efficiently without the need for CD8. (A) Primary CD8+ T-cells were co-transduced with the 868 TCR and rat (r)CD2 prior to staining with 0.5 µg (with respect to MHC) of irrelevant (A2–ALWGPDPAAA), A2–SLYNTVATL, A2–SLYNTIATL, and A2–SLFNTIAVL PE-conjugated peptide–MHC tetramers. Cells are gated on rCD2+CD8+ cells. Histograms show staining with indicated PE-conjugated tetramer with the mean fluorescence intensity (MFI) of this population displayed. (B) As for panel (A), but 868 TCR and rat (r)CD2 were transduced into TCRβ chain negative Jurkat cells. MFI of tetramer staining is shown for cells in the rCD2+ tet+ gate. (C) 868 TCR transduced were incubated with T2 cells and either SLYNTVATL or SLFNTIAVL peptide at the concentrations shown for 5 h followed by the detection of CD107a and TNFα by flow cytometry.

Table 5 | Relative binding of SLYNTVATL and SLFNTIAVL to HLA A*0201.

| SLYNTVATL | SLFNTIAVL | Method |
|------------|-----------|--------|
| 100%       | 48%       | S35 methionine pulse chase (9) |
| 100% (Kd = 50 nM) | 53% (Kd = 94 nM) | Direct competition with radio-labelled ligand (18) |
| 100% (Kd = 300 nM) | 300% (Kd = 100 nM) | Quantitative HLA A2 ELISA (6) |
| 100% (Kd – 3.5 nM) | 400% (Kd – 0.8 nM) | Quantitative HLA A2 ELISA (7) |

better targets at t₀ (125%) but their ability to activate the 868 TCR cells diminished to 18% when allowed to culture for 24 h prior to assay (Figure 8D; Figure S2 in Supplementary Material). These results suggest that the major mechanism by which SLFNTIAVL escapes from CTL is through loss of peptide–HLA stability, although reduced TCR interaction with this mutant is also likely to play a secondary role.
**Figure 7** | Stability of A2–SLYNVTAVL, A2–SLYNTIAVL, and A2–SLFNTiAVL. (A) CD thermal denaturation curves recorded at 218 nm are shown for selected samples as indicated. Dots represent measured values fitted assuming a two-state trimer-to-monomer transition (solid lines) as described in Section “Materials and Methods.” (B) Bar graphs of the thermal stability with respect to melting temperature (upper) and van’t Hoff’s enthalpy of unfolding (lower panel). Error bars in panels (A,B) represent SD resulting from the multivariable curve fitting of data from one experiment with each peptide. (C) T2 cells incubated for 3 or 22 h with 10⁻⁵ M of each peptide [Influenza (Flu) Matrix, GILGFVFTL, HIV Gag wild-type SLYNTAVL, and Gag triple mutant SLFNTiAVL or DMSO control were stained for HLA-A2 and fixed. The dashed line is set at the mean fluorescence intensity (MFI) of the negative control peptide (EBV HPVGEADYFEY that binds HLA B*3501).

**Figure 8** | Instability at the cell surface mediates viral escape by the triple mutant epitope, A2–SLFNTiAVL. (A) Relative binding of 868 TCR to A2–peptide complexes by SPR over time. The SPR chip was maintained at 37°C. Functional half-life was determined by injecting the 868 TCR at a fixed concentration of 100 µM over 720 min and recording the drop in total RUs over 8 different time points. A faster drop in RUs (converted into a T<sub>1/2</sub>) was interpreted to indicate greater peptide instability, detected by loss of binding by the antigen-specific 868 TCR. (B) Left panel shows relative HLA-A2 expression following incubation (0, 3, and 22 h) of T2 cells with indicated peptide. Results are normalised to the maximum mean fluorescence intensity seen for each peptide. The right panel shows T2 cells incubated with peptide for 22 h prior to being washed and cultured in an excess of AIM-V media for the indicated times prior to staining and fixing. Data displayed as HLA-A2 expression relative to maximum seen at 22 h. (C) T2 cells were pulsed for 1 h with the peptides shown, washed extensively, and then cultured in an excess of media for 3 h (black bar) or 24 h (open bar) before being co-incubated for 5 h with T-cells from two donors transduced with the 868 TCR. CD107a and TNFα were used to establish percentage reactivity by flow cytometry. Values were normalised relative to SLYNTAVL peptide. SEM is shown for two different donor T-cells. Untransduced CD8<sup>+</sup> T-cells did not respond to peptide (data not shown). Raw data for panel (C) is shown in Figure S1 in Supplementary Material. (D) As panel (C) but using primary HLA-A2<sup>−</sup> and HLA-A2<sup>+</sup> CD4<sup>+</sup> T-cells as targets and culturing in excess media for either 0 or 24 h prior to assay. Raw data for panel (D) is shown in Figure S2 in Supplementary Material.
DISCUSSION

Recognition and escape from the HLA A2-restricted HIV epitope SL9 has been examined in well over 50 studies to date, but major questions are still outstanding in this system as there has been no TCR–A2–SL9 co-complex structure. Here, we solved the structure of the free 868 TCR, a TCR first described almost 20 years ago (10), and the structure of this TCR in complex with A2–SLYNTVATL. Comparison of the free and ligated TCR structures suggested an “induced fit” mechanism that involved major structural reorganisation of the antigen binding face of the TCR. The CDR3 loops made the biggest movement (almost 7 Å for CDR3α and >3 Å for CDR3β) to avoid steric clashes with the central portion of the peptide, while the CDR1 and CDR2 loops interacted mainly with MHC via a more “lock-and-key” interaction.

It has been known for over a decade that the SLYNTVATL and SLENTVATL bind to HLA A2 with very different conformations, but engage the G10 TCR with near identical on-rates, leading the authors to conclude that one conformation, common to both peptides, was recognised by TCRs (22). Since A2–SLYNTVATL does not change upon TCR binding, we can now be confident that the conformation adopted by this, and many other SL9 variants (21), dominates during TCR recognition. Thus, the evidence suggests that A2–SLENTVATL undergoes an induced fit upon TCR binding (22) although conclusive proof will await a TCR–A2–SLENTVATL structure. We further solved the co-complex structure of the 868 TCR with the A2–SLEYNTIAL (the variant that predominated in patient 868 in 1996 when the patient blood sample from which this T-cell clone was grown was taken), and the structure of the 868 TCR in complex with the A2–SLEYNTIAL, the ultimate escape variant in this system. Resolution of the latter structure came as a surprise as residues that mutate in positions 3, 6, and 8 to proline in SL9 escapes from CTL via a more “lock-and-key” interaction measured by BIAcore (52). The median affinity for human antiviral TCR–pMHC interactions is $K_D \sim 5 \mu M$ (52). 868 TCR binding was characterised by a relatively slow off-rate from the A2–SLYNTVATL and A2–SLEYNTIAL ligands. The other SL9-TCR that has been examined by SPR, G10, had an off-rate 3.75 times faster than 868, and a 4.5 times slower on-rate, resulting in a weaker affinity of $K_D$ of 2.2 µM for A2–SLYNTVATL compared to 868 (22). While binding of the 868 TCR was much reduced for A2–SLEYNTIAL compared to the other ligands, it still had a very respectable affinity ($K_D = 1.77 \mu M$) that falls at the higher end of the spectrum for TCR agonists (52).

The mode of 868 TCR engagement and its affinity with the A2–SLEYNTIAL ultimate escape variant in this system did not lend full support to the prevailing belief that loss of recognition of SL9 occurs mainly via interfering with TCR binding. The fact that CD8$^+$ T-cells transduced with the 868 TCR could recognise SLEYNTIAL pulsed target cells well and stained with A2–SLEYNTIAL tetramers was also a surprise, but in accordance with an earlier study that showed that A2–SLEYNTIAL bound to the D3 TCR as well as A2–SLYNTVATL in tetrameric form and that the SLEYNTIAL concentration required for half maximal lysis was a highly respectable $4 \times 10^{-10} M$ (21). In further accordance with positions 3, 6, and 8 not dominating A2–SL9-specific TCR binding, the position 4 asparagine sub-library was by far the most potent of the 180 sub-mixtures for activating T-cell clone 003 known to express a different SL9-specific TCR (62). These combined data do not support the notion that the SLEYNTIAL escapes from CTL via lack of TCR binding, prompting us to take a closer look at the MHC binding of ligands in this system. Data from three different studies using three different methodologies show that SLEYNTIAL binds to HLA A2 with 50–400% of the affinity of SLYNTVATL. Our previous SPR experiments have shown that good agonists produce pMHC with a half-life of >6 hours in a cell-free system (61). Indeed, our data here show that both A2–SLYNTVATL and A2–SLEYNTIAL had half-lives of over 7 hours on an SPR chip maintained at 37°C compared to <2 hours for A2–SLEYNTIAL. We reasoned that the peptide-binding assays that have been performed previously in this system, including by ourselves (9), would be most influenced by peptide on-rate but that the biology would be most linked to peptide off-rate at the target cell surface. HLA A2$^+$ target cells pulsed with SLYNTVATL peptide remained effective targets for SL9-specific CD8$^+$ T-cells after 24 hours of culture as has been observed with a different HIV Gag-derived epitope (27). The SLEYNTIAL, SLEYNTIAL and SLYNTVATL peptides [which bind to the 868 TCR with dissociation constants of 2.9 µM, <100 nM, and ~300 nM, respectively (52)] were seen well when targets presenting these ligands were used immediately for T-cell assays but exhibited only 67%, 60%, and 35% recognition of that observed with SLYNTVATL peptide.
when targets were cultured for 24 h prior to the T-cell assay. The biggest effect was observed with the SLFNTVIAVL where >80% of the activity was lost after 24 h of culture. These results were supported by other experiments looking the stability of the A2–SLFNTVIAVL complex in comparison to other ligands.

Collectively, our results suggest that the instability of the A2–SLFNTVIAVL molecule is likely to provide the major mechanism of escape from SL9-specific CTL. The accumulation of mutations at positions 3, 6, and 8 to fixation over a 10-year period (6, 7) demonstrates the extraordinary lengths that HIV will go to in order to avoid detection by these key antiviral cells. Our study further highlights that the results of short-term, steady-state peptide-binding assays can be misleading and suggests that future studies should employ more relevant biological readouts.

**AUTHOR CONTRIBUTIONS**

DC, AF, GD, EZ, ML, KM, ILB, FM, CH, AB, JSB, JM, AJAS, KB, BE, and PR performed and/or directed experiments, analysed data, and critiqued the manuscript. DC and AKS conceived, funded, and directed the project and wrote the manuscript.

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**SUPPLEMENTARY MATERIAL**

The Supplementary Material for this article can be found online at http://www.frontiersin.org/article/10.3389/fimmu.2017.01503/full#supplementary-material.

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