Inflammation-Associated Nitrotyrosination Affects TCR Recognition through Reduced Stability and Alteration of the Molecular Surface of the MHC Complex

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Abstract
Nitrotyrosination of proteins, a hallmark of inflammation, may result in the production of MHC-restricted neoantigens that can be recognized by T cells and bypass the constraints of immunological self-tolerance. Here we biochemically and structurally assessed how nitrotyrosination of the lymphocytic choriomeningitis virus (LCMV)-associated immunodominant MHC class I-restricted epitopes gp33 and gp34 alters T cell recognition in the context of both H-2D^b and H-2K^b. Comparative analysis of the crystal structures of H-2K^b/gp34 and H-2K^b/NY-gp34 demonstrated that nitrotyrosination of p3Y in gp34 abrogates a hydrogen bond interaction formed with the H-2K^b residue E152. As a consequence the conformation of the TCR-interacting E152 was profoundly altered in H-2K^b/NY-gp34 when compared to H-2K^b/gp34, thereby modifying the surface of the nitrotyrosinated MHC complex. Furthermore, nitrotyrosination of gp34 resulted in structural over-packing, straining the overall conformation and considerably reducing the stability of the H-2K^b/NY-gp34 MHC complex when compared to H-2K^b/gp34. Our structural analysis also indicates that nitrotyrosination of the main TCR-interacting residue p4Y in gp33 abrogates recognition of H-2D^d/gp33-NY complexes by H-2D^d/gp33-specific T cells through steric hindrance. In conclusion, this study provides the first structural and biochemical evidence for how MHC class I-restricted nitrotyrosinated neoantigens may enable viral escape and break immune tolerance.

Introduction
The biological function of proteins and cellular metabolism can be significantly modulated through post-translational modifications (PTMs) [1,2]. Naturally occurring PTMs can either result from enzyme modifications or arise spontaneously [3,4]. However, PTMs can also be used by bacterial and viral pathogens to compromise critical immune responses and host factors acting against infection [5]. The breakdown of immunological tolerance resulting from such modifications may render endogenous antigens immunogenic [3,6]. A large array of PTM peptides presented by major histocompatibility complex (MHC) class I and class II molecules have now been described [7,8], and it has been demonstrated that modifications such as cysteinylation [9,10], glycosylation [11,12] or phosphorylation [13,14] may affect T cell immunoreactivity, resulting in immune escape and/or initiation of autoimmunity.

Protein tyrosine nitration is a hallmark of inflammation, and is associated with up-regulated expression of inducible nitric oxide synthase (iNOS) [15,16]. Nitrotyrosinated proteins (NT-proteins) are generated in vivo by nitration with peroxynitrite, itself derived from nitric oxide and superoxide which are both released from activated inflammatory cells [17]. NT-proteins may accumulate in apoptotic or inflamed tissues [18]. Indeed, accumulation of the amino acid homolog 3'-nitrotyrosine (NY) and of NT-proteins has been observed at inflammatory sites in Alzheimer’s disease [19], arthritis [20,21,22,23], atherosclerosis [22,23], autoimmune diabetes [24], autoimmune uveitis [25], celiac disease [26], ischemia-reperfusion injury [27], multiple sclerosis [28], Parkinson’s disease [29], respiratory disease [30] and transplant rejection [31], as well as in various cancers and infectious diseases [32]. NT self-proteins may thus be highly immunogenic, eliciting both humoral and cellular responses [17,33].

The potential immunogenicity of MHC class II-restricted NT autologous peptides has been previously investigated using the I-A^b-restricted T cell pigeon/moth cytochrome c (PCC/MCC) [34] as well as the I-A^b-restricted hen egg-white lysozyme (HEL) epitopes [35]. Both studies demonstrated unequivocally that conversion of tyrosine to nitrotyrosine resulted in dramatic consequences for T cell recognition. Indeed, processing of native proteins by activated antigen presenting cells (APCs) resulted in the presentation of MHC class II-restricted NT-epitopes in lymphoid tissues, significantly altering specific T cell responses and eliciting NT-specific CD4+ T cells that evaded negative selection in the thymus and thus central tolerance [33]. Likewise
loss of tolerance by NT-specific CD4+ T cells has recently been shown to be critical for the production of autoantibodies [36]. These studies indicate that NT-proteins generated during inflammation might constitute an important class of neoantigens that could promote autoimmune T cell responses.

The well-established lymphocytic choriomeningitis virus (LCMV) glycoprotein system was also used by Hardy et al. to demonstrate that conversion of tyrosine to NY also profoundly affected T cell recognition of MHC class I-restricted epitopes [37]. A significant amount of the overall CD8+ T cell response to LCMV is dominated by very few viral epitopes, comprising the H-2Db-restricted epitope gp33–41 (hereafter referred to as gp33; KAVYNFATC) and the H-2Kb-restricted peptide gp34–41 (gp34; AVNYNATC) [38,39]. Both gp33 and gp34 contain a single tyrosine residue at positions 4 (p4Y) and 3 (p3Y), respectively. T cell populations, exclusively specific to the single tyrosine residue at positions 4 (p4Y) and 3 (p3Y), respectively. T cell populations, exclusively specific to the nitrotyrosinated MHC complexes H-2Db/NY-gp33, H-2Kb/NY-gp34 and/or H-2Kb/NY-gp34 [37], were elicited in C57BL/6 mice (H-2Db/H-2Kb) following immunization with the nitrated peptide NY-gp33. Importantly, CD8+ T cell hybridomas specific for NY-gp33 comprised two distinct subsets recognizing either H-2Db/NY-gp33 or H-2Kb/NY-gp33. While the T cell hybridoma 2H1 responded to stimulation with both H-2Kb/NY-gp34 and H-2Db/NY-gp33, it did not recognize the unmodified wild-type MHC complex. The possible implications for the role of nitrotyrosination in the creation of modified neoantigens that possibly results in autoimmune disorders are discussed.

Materials and Methods

Preparation and crystallization of H-2Kb-restricted gp34 and NY-gp34

Peptides gp34 (AVNYNATM) and NY-gp34 (AV-p3NY-NFATM) were purchased from Genscript (Piscataway, NJ, USA). Refolding and purification of MHC/peptide complexes were followed according to previously published protocols [40,41,42,43]. The best crystals for H-2Kb/gp34 and H-2Kb/NY-gp34 were obtained in hanging drops by vapor diffusion in 2.1 M NaH2PO4/K2HPO4 (pH 6.4), 1.5% MPD and 1.8 M NaH2PO4/K2HPO4 (pH 6.7), 1.5% MPD, respectively. Typically, 2 μl of 6 mg/ml protein were equilibrated against 2 μl of crystallization reservoir solution at 20°C.

Data collection and processing

Data collection was performed under cryogenic conditions (temperature 100 K) at beam lines ID14-2 and ID14-4 at the synchrotron radiation facility at ESRF (Grenoble, France) to a resolution of 2.0 and 2.6 Å for H-2Kb/gp34 and H-2Kb/NY-gp34, respectively. Crystals were soaked in a cryoprotectant solution containing 25% glycerol before data collection. A total of 180 and 360 images were collected with 0.5° oscillation per frame for both H-2Kb/gp34 and H-2Kb/NY-gp34. Data were processed with MOSFLM [44] and SCALa from the CCP4 suite [45]. The space group was determined to be P212121 for H-2Kb/gp34 with unit-cell parameters a = 80.4, b = 92.6, c = 128.8 Å. The space group for H-2Kb/NY-gp34 was P21 with unit-cell parameters a = 50.5, b = 88.5, c = 119 Å, α = γ = 90°0′ and β = 94.7°. Data collection statistics are provided in Table 1.

Crystal structure determination and refinement

The crystal structure of H-2Kb/gp34 was solved by molecular replacement (MR) using Phaser [46] and the H-2Kb/GNYSFYAL MHC complex (PDB code 1LK2) as a search model. Five percent of the total reflections were set aside for monitoring refinement by Rfree. The crystal structure of H-2Kb/NY-gp34 was solved thereafter by MR using H-2Kb/gp34 as a search model. Refinement of the two crystal structures was performed using REFMAC5 [47]. After each round of refinement, missing residues were added in successive cycles of manual building followed by restrained refinement cycles in REFMAC5 [47]. The final refinement parameters are presented in Table 1. Figures were created using the program PyMOL [48].

Peptide-MHC binding affinity assays

Peptide-MHC binding affinity assays were performed using transporter associated with antigen processing (TAP)-deficient RMA-S cells as described previously [49], by assessing the capacity of the different peptides to stabilize cell surface expression of H-2Kb complexes. Briefly, 5 × 10⁴ RMA-S cells were pulsed with different concentrations of indicated peptides in serum free AIM-V medium (Invitrogen, Carlsbad, CA, USA) at 26°C overnight in 5% CO2. Cells were subsequently washed and incubated in AIM-V medium at 37°C for 60 min in the absence of peptides. Cells were then washed twice with PBS before staining with anti-H-2Kb AF6-88.5 (BD Pharmingen/BD Biosciences, MountainView, CA, USA). Following washing and fixation in 1% PFA, H-2Kb cell surface expression was detected by flow cytometry on BD FACSCalibur (BD Biosciences). Flow cytometry data was analyzed using Cell Quest Pro (BD Biosciences). The mean fluorescence intensity (MFI) of H-2Kb expression for the indicated peptide concentrations was divided by the observed MFI on cells without peptide as an estimate of peptide binding. The HIV-derived H-2Dd-restricted epitope P16 (RGPGRAFVT) was used as a negative control while the H-2Kb-restricted Moloney murine leukemia virus peptide MULV (SSWDTFV) was used as a positive control.

Assessment of MHC complex stability using circular dichroism (CD) analysis

CD measurements were performed in 20 mM K2HPO4/KH2PO4 (pH 7.5) using protein concentrations from 0.15 to 0.25 mg/ml. Spectra were recorded with a Jasco J-810 spectropolarimeter (Jasco Analytical Instruments, Great Dunmow, UK) equipped with a thermoelectric temperature controller in a 2 mm cell. pMHC denaturation was measured between 30°C and 75°C at 218 nm with a gradient of 4°C/hour at 0.1°C increments and an averaging time of 8 seconds. The melting curves were scaled from 0% to 100% and the Tm values extracted as the temperature at 50% denaturation. Curves and Tm-values are an average of at least two measurements from two independent refolding assays per MHC complex. Spectra were analyzed using GraphPad Prism 5 and Tm values compared using an unpaired, two-tailed T test.
Assessment of MHC complex stability using Thermofluor analysis

Differential scanning fluorimetry experiments were carried out using an iQ5 real-time PCR detection system (Bio-Rad). 96-well PCR plates were filled with 23 μl of MHC solution (0.25 mg/mL) in 10 mM Na⁺ HEPES buffer (pH 7.3), 150 mM NaCl per well. Finally, the fluorophore (2 μl of 125μl SYPRO Orange (Sigma S5692)) was added to each well and the plate was sealed with optical sealing tape (Bio-Rad). An iQ5 Real-Time PCR Detection System, calibrated with External Well factor solution was used to monitor the changes in fluorescence intensity of the fluorophore. The temperature of the samples was changed from 20 to 95°C at a heating rate of 1°C/min and the fluorescence was recorded every 0.2°C. The melting temperature (Tm) of each melting curve was calculated as the maximum of the first derivative of the curve. The average Tm value of each protein was determined from three distinct melting curves.

Molecular modeling of TCR in complex with wild-type and nitro-tyrosinated MHC complexes

Molecular models of the TCR P14 in complex with H-2Db/gp33 or H-2Db/NY-gp33 were created using previously described protocols [50,51]. Interactions between P14 and each MHC complex were analyzed visually for improper sterical contacts. Docking of P14 on its MHC/peptide ligand was also assessed according to contacts between evolutionary conserved amino acids suggested to be important for TCR/MHC interaction [52]. The molecular model was subjected to several rounds of energy minimization using the CNS suite of programs [53]. The coordinates of the TCR/MHC/peptide complexes will be provided upon request.

Results and Discussion

Overall structures of H-2Kb in complex with wild-type gp34 and nitrotyrosinated NY-gp34

The three dimensional structures of H-2Kb in complex with the wild-type gp34 (AVYNFATM) and with the nitrotyrosinated NY-gp34 (AV-p3NY-NFATM) were determined to 2.0 and 2.6 Å resolutions, respectively (Table 1). Both crystal structures displayed good stereochemistry. The final electron density maps are of good quality with well-defined polypeptide chains (Figures 1A and 1B). In particular, the electron densities for all H-2Kb residues in contact with the peptides gp34 and NY-gp34 are clearly defined. The overall three-dimensional structures of H-2Kb/gp34 and H-2Kb/NY-gp34 are remarkably similar with a root mean square deviation (rmsd) of 0.37 Å for the entire MHC complexes (heavy chain).
chains in complex with peptides and β2-microglobulin (β2m) and of 0.27 Å² for heavy chain residues 1–176 corresponding to the α1 and α2 domains.

Protrusion of residue p1K in H-2Kb/gp33 does not alter the conformation of MHC residues within the N-terminal part of the peptide-binding cleft

The previously determined crystal structure of H-2Kb in complex with gp33 (KAVYNFATM) revealed that the nonameric peptide binds to H-2Kb as an octamer with the stretch of residues AVYNFATM filling the peptide-binding cleft, while the highly flexible residue p1K extended out of the binding groove [50] (Figure 2A). As for p2A in H-2Kb/gp33, the first peptide residue p1A in both H-2Kb/gp34 (AVYNFATM) and H-2Kb/NY-gp34 (AV-p3NY-NFATM) is surrounded by a cluster of tyrosine residues (Y7, Y159 and Y171) that forms a network of hydrogen bond interactions with the nitrogen atom of p1A. However, although the protrusion of p1K out of the peptide-binding cleft in H-2Kb/gp33 results in a significant modification of the position of the nitrogen atom of residue p2A when compared to p1A in H-2Kb/gp34 or H-2Kb/NY-gp34, this does not affect the conformation of the cluster of tyrosine residues that surround the N-terminal part of the peptide (Figure 2B). Furthermore, the conformation of the stretch of peptide residues AVYNFATM bound within the peptide-binding cleft is very similar in both H-2Kb/gp33 and H-2Kb/gp34 (Figure 2A). Thus although the N- and C- peptide termini play crucial roles in binding of epitopes within the peptide-binding cleft formed by the α1/α2 groove [54], these are not always necessarily required. It has been previously demonstrated that longer peptides may protrude out of the groove and that modified and/or shorter peptides can still bind efficiently within the groove through the use of a network of water molecules within the peptide binding cleft [50,55,56,57].

In the prior study by Hardy et al., T cells could be readily elicited in C57/B16 mice following immunization with NY-gp33 [37].
Interestingly, while the H-2K\textsuperscript{b}/NY-gp33-specific CD8\textsuperscript{+} T cell hybridoma 24H1 recognized H-2K\textsuperscript{b} target cells coated with both NY-gp33 and NY-gp34, it did not recognize H-2K\textsuperscript{b}/gp34 MHC complexes. Furthermore, it should also be noted that 24H1 T cells recognized H-2K\textsuperscript{b}/NY-gp34 to a higher level when compared to H-2K\textsuperscript{b}/NY-gp33. The relatively lower recognition of H-2K\textsuperscript{b}/NY-gp33 by 24H1 could be explained by the protrusion of the side chain of p1K from the N-terminal part of the peptide-binding cleft of H-2K\textsuperscript{b}/NY-gp33, which could potentially limit optimal recognition by the TCR. However, the question remained as to why the H-2K\textsuperscript{b}/NY-gp33 and H-2K\textsuperscript{b}/NY-gp34-specific TCR of 24H1 did not recognize H-2K\textsuperscript{b}/gp34?

Nitrotyrosination of peptide residue p3Y induces a profound conformational change of the H-2K\textsuperscript{b} residue E152, altering TCR recognition

The conformations of the peptides gp34 and NY-gp34 are very similar when bound to H-2K\textsuperscript{b}, besides a lateral shift of 0.7 Å of the side chain of the nitrotyrosine p3NY in H-2K\textsuperscript{b}/NY-gp34 when compared to the conformation of p3Y in H-2K\textsuperscript{b}/gp34 (Figure 3A). In H-2K\textsuperscript{b}/gp34 the side-chain of p3Y projects down into the D-pocket formed by residues Q114, E152, R155, L156 and Y159 (Figures 3A and 3B). Three hydrogen bond interactions are formed between the side chain of p3Y and the side chains of the H-2K\textsuperscript{b} residues E152 and R155 (Figure 3B). Nitrotyrosination of p3Y alters only the conformation of the H-2K\textsuperscript{b} heavy chain residue E152 which, in order to accommodate the side-chain of p3NY, undergoes a large conformational modification in H-2K\textsuperscript{b}/NY-gp34 when compared to H-2K\textsuperscript{b}/gp34 (Figure 3C). In contrast, the rest of the H-2K\textsuperscript{b} peptide-binding groove is not affected by the nitrotyrosination of the peptide in H-2K\textsuperscript{b}/NY-gp34 when compared to H-2K\textsuperscript{b}/gp34.

In conclusion, substitution of the tyrosine residue at position 3 of the peptide gp34 to a nitrotyrosine exclusively affects the conformation of the side-chain of the H-2K\textsuperscript{b} residue E152. The specificity of the T cell hybridoma 24H1 to H-2K\textsuperscript{b}/NY-gp34 thus seems to be dependent on the conformation of the MHC region surrounding the side-chains of p3NY and the H-2K\textsuperscript{b} residue E152. It should be noted that previous studies have demonstrated the importance of this region for recognition by H-2K\textsuperscript{b}-specific TCRs [58,59]. One may thus speculate that nitrotyrosination of the immunodominant peptide which alters considerably T cell recognition may be advantageous for the virus in order to escape immune recognition.

Nitrotyrosination of gp34 strains the overall conformation of the H-2K\textsuperscript{b} complex and significantly decreases its stability

Two additional interactions are formed between peptide residue p3NY and the heavy chain residues Q114 and Y116 in the H-2K\textsuperscript{b}/NY-gp34 peptide when compared to H-2K\textsuperscript{b}/gp34 (Figures 3B and 3C), suggesting that nitrotyrosination of the peptide may result in higher stabilization capacity of H-2K\textsuperscript{b} complexes by NY-gp34 when compared to gp34. However, binding affinity assays using TAP-deficient cells demonstrated that while both gp33 and gp34 displayed similar binding affinities to H-2K\textsuperscript{b} complexes, nitrotyrosination of peptide residue 3 significantly reduced binding affinity of NY-gp34 to H-2K\textsuperscript{b} (Figure 4A).

Furthermore, the capacity of each peptide to stabilize H-2K\textsuperscript{b} complexes was also investigated using circular dichroism (CD) by assessing the thermostability of soluble H-2K\textsuperscript{b} molecules in complex with gp34 or NY-gp34. Melting temperatures (T\textsubscript{m}) were derived from changes in ellipticity at 218 nm, corresponding to loss of secondary structure during denaturation. Our analysis demonstrates that the T\textsubscript{m} value for H-2K\textsuperscript{b}/NY-gp34 (47.3°C) was nearly 10°C lower than for H-2K\textsuperscript{b}/gp34 (56.7°C), reflecting the significantly reduced capacity of NY-gp34 to stabilize H-2K\textsuperscript{b} complexes (Figure 4B). Similar very significant shifts in thermal stability were also obtained for the two MHC class I complexes upon analyzing their stability with differential scanning fluorimetry.

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Furthermore, the capacity of each peptide to stabilize H-2K\textsuperscript{b} complexes was also investigated using circular dichroism (CD) by assessing the thermostability of soluble H-2K\textsuperscript{b} molecules in complex with gp34 or NY-gp34. Melting temperatures (T\textsubscript{m}) were derived from changes in ellipticity at 218 nm, corresponding to loss of secondary structure during denaturation. Our analysis demonstrates that the T\textsubscript{m} value for H-2K\textsuperscript{b}/NY-gp34 (47.3°C) was nearly 10°C lower than for H-2K\textsuperscript{b}/gp34 (56.7°C), reflecting the significantly reduced capacity of NY-gp34 to stabilize H-2K\textsuperscript{b} complexes (Figure 4B). Similar very significant shifts in thermal stability were also obtained for the two MHC class I complexes upon analyzing their stability with differential scanning fluorimetry.
The melting temperatures $T_m$ corresponding to the mid-point of the sharp transition from folded to unfolded state were 54.4 $\pm$ 0.1 and 44.5 $\pm$ 0.2 for H-2Kb/gp34 and H-2Kb/NY-gp34 (Figure 4C). These significant differences in stabilization capacity were at first surprising when considering that additional hydrogen bonds and long ionic interactions are formed between NY-gp34 and H-2Kb when compared to H-2Kb/gp34 (Figure 3C). However, further comparative analysis of the two crystal structures provided an explanation to these unexpected results.

In H-2Kb/NY-gp34, the side-chain of the nitrotyrosine p3NY projects within the D-pocket of H-2Kb, resulting in structural over-packing (Figures 3A and 3C) and a 'strained' overall conformation of the MHC complex. Although this has not, to our knowledge, yet been reported for MHC class I molecules, strained conformations are relatively common and have been described for several proteins including e.g. native serine protease inhibitors [61] and spectrin SH3 core-domains [62]. Induction of strained conformations is also important within several processes including catalysis in thymidylate synthase [63] and nuclear transport of exportins [64]. Proteins with strained ‘spring-loaded’ conformations are usually intrinsically unstable. For example, the release of the course of the strained conformation in circularly permutated barnases resulted in a 20°C shift in melting temperature [65] or in a 25-fold increase in stability for mutated serpins [66]. In the crystal structure of the H-2Kb/NY-gp34 complex, introduction of the side chain of p3NY resulted in close contacts between the nitrotyrosinated peptide and several H-2Kb residues, forcing them away from the relaxed low-energy conformation observed in H-2Kb/gp34 in order to accommodate the side chain of p3NY, thus resulting in unfavorable strained conformations. Besides straining the backbone of the heavy chain residues 124–126 and 156–157, the side chain of p3NY also incurred strained higher internal energy conformations for the side-chains of residues V97, I98 and Q114, when compared to H-2Kb/gp34.

In conclusion, the potential gain in enthalpy resulting from the formation of two additional interactions between p3NY and the H-2Kb residues Q114 and Y116 may be outweighed by the loss of a hydrogen bond interaction formed with the side chain of E152 combined with the free energy cost resulting from the strained conformations of several MHC residues. Our results indicate that formation of additional interactions between peptides and MHC residues does not always necessarily increase the overall stabilization of MHC complexes. Furthermore, alteration of MHC complex stabilization through nitrotyrosination of the presented epitope may represent an additional alternative strategy used by the virus to prevent adequate presentation of the peptide at the surface of the infected cell.

Nitrotyrosination of p4Y in H-2D$^b$/NY-gp33 directly affects recognition by H-2D$^b$/gp33-specific TCRs

The previously determined crystal structures of H-2D$^b$ and H-2K$^b$ in complex with gp33 demonstrated that the peptide binds in two diametrically opposed manners to the two MHC class I molecules [50]. Thus in contrast to H-2K$^b$/gp33 and H-2K$^b$/ gp34, in which the side-chain of the tyrosine residue acts as a secondary anchor position, the side-chain of p4Y in H-2D$^b$/gp33 protrudes out of the peptide-binding cleft, playing a key role in TCR recognition [41,43,50]. Indeed, mutation of p4Y to a
Figure 4. Nitrotyrosination of gp34 significantly decreases both its binding affinity and stabilization capacity of the H-2Kb complex. A. The binding affinity of NY-gp34 to H-2Kb on the surface of TAP-deficient RMA-S cells was significantly reduced when compared to gp33 and gp34. Cells were incubated with the indicated concentrations of each peptide at 26°C for 12 hours in 5% CO₂. Cells were collected and stained with the H-2Kb-specific antibody AF6-88.5. The mean fluorescence intensities (MFI) observed for each timepoint were normalized as described in the peptide binding affinity assays outlined above. The H-2Dd and H-2Db-restricted peptides P18-I10 and MULV were used as negative and positive controls, respectively. The peptide binding assays were repeated three times. B. The capacity of NY-gp34 to stabilize soluble H-2Kb complexes is significantly reduced when compared to gp34. MHC complex melting temperatures (Tm) were measured using circular dichroism. Tm values were derived from normalized thermal denaturation curves for the MHC complexes H-2Kb/gp34 (in grey) and H-2Kb/NY-gp34 (in black) as the temperature corresponding to 50% denaturation (dashed line). The presented denaturation curves are an average from at least two independent measurements for each MHC complex. C. The denaturation curves of the two MHC complexes, using the differential scanning fluorimetry approach (thermofluor), are consistent with the two-state model of thermal unfolding. The melting temperatures, defined as the inflection points of the escalating parts of the curves, the curves of H-2Kb/gp34 and H-2Kb/NY-gp34 are in grey and black, respectively.
doi:10.1371/journal.pone.0032805.g004
Figure 5. Nitrotyrosination of p4Y in H-2D\(b\)/NY-gp33 directly affects recognition by H-2D\(b\)/gp33-specific TCR. Nitrotyrosination of the main TCR-interacting peptide residue p4Y will affect the structural conformation of both TCR interacting residues on H-2D\(b\) and of the TCR P14. The peptide binding cleft of H-2D\(b\) and the TCR, both colored in white, are annotated. Hydrogen bond interactions appear as dotted lines. A. In H-2D\(b\)/gp33, the side chain of p4Y protrudes out of the H-2D\(b\)peptide-binding cleft, positioning itself perfectly in the hot spot of the p14 TCR composed of the CDR3 loops from both V\(\alpha\) and V\(\beta\). It forms three hydrogen bonds, two of them directly with Y36(V\(\alpha\)) and G102(V\(\beta\)) on the TCR P14. The last hydrogen bond is formed with the side chain of the H-2D\(b\) histidine residue H155, linking this domain of the heavy chain to the TCR. B. The side chain of the nitrotyrosinated p4-NY can not be accommodated within the hot-spot of P14, resulting in sterical clashes with the side chain of the TCR residue Y36(V\(\alpha\)). Furthermore, the negatively charged side chain of the H-2D\(b\) residue E163, important for TCR recognition, would also be repelled by the introduced negatively charged nitrotyrosination. C. Similarly, the other rotamer of the nitrotyrosinated p4-NY would result in sterical clashes with both G102(V\(\beta\)) on the side chain of H155, abolishing all formed hydrogen bond interactions.

doi:10.1371/journal.pone.0032805.g005

phenylalanine (Y4F) reduced the affinity of the H-2D\(b\)/gp33 specific TCR P14 by 100-fold [51,67]. A molecular model of P14 in complex with the crystal structure of H-2D\(b\)/gp33 indicated that the side-chain of the protruding p4Y is perfectly positioned in the hot spot of the TCR composed of the CDR3 loops from both the V\(\alpha\) and V\(\beta\) domains. It forms three hydrogen bonds with the TCR residues Y36(V\(\alpha\)) and G102(V\(\beta\)), as well as with the side chain of the H-2D\(b\) histidine residue H155, linking this domain of the heavy chain to the TCR (Figure 5A). In contrast, insertion of the negatively charged NO\(_2\) group in p4Y would result in deleterious sterical clashes with Y36(V\(\alpha\)) or G102(V\(\beta\)) (Figures 5B and 5C). The side chain of one of the possible p4-NY rotamers would also repulse the negatively charged side-chain of the MHC residue E163, important for TCR recognition (Figure 5B).

In conclusion, nitrotyrosination of p4Y in H-2D\(b\)/NY-gp33 is clearly deleterious for recognition by P14 as previously demonstrated [37]. Thus, one may here speculate again about the possibility that nitrotyrosination of the H-2D\(b\)-restricted peptide gp33 may represent an important strategy used by LCMV in order to avoid CD8\(^+\) T cell recognition and escape immune recognition. Furthermore, and as for H-2K\(b\)/NY-gp34, our structural analysis also strongly indicates that NY-gp33 may act as an H-2D\(b\)-restricted neo-epitope with the capacity to break tolerance, selecting and inducing inadequate T cell responses that may result in auto-reactivity.

Concluding remarks

Our study provides a structural platform underlying the effects of peptide nitrotyrosination on initiation of TCR responses. Such PTMs can directly affect TCR recognition by modifying the properties of key TCR-interacting residues on the presented peptide or by altering the conformations of other MHC residues that are of importance for TCR recognition. Nitrotyrosination can also indirectly affect TCR recognition by severely destabilizing the MHC complex. Although additional interactions were formed between the PTM peptide and H-2K\(b\), the conformation of a large amount of surrounding residues was strained, significantly reducing the overall stability of the MHC complex. We have previously demonstrated that subtle modifications in MHC-restricted peptides can result in significant alterations of MHC stabilization [51], a phenomenon that should be taken into account and measured upon trying to design altered peptide ligands for modulation of TCR response [49,68]. Several other amino acid modifications caused by oxidative stress could result in similar effects on TCR recognition [69,70]. In conclusion, our structural study indicates that the impact of post-translational modifications could be dual, possibly allowing for viral immune escape from TCR recognition but also potentially inducing the expansion of subset of T cells that could induce auto-reactivity. Likewise, nitrotyrosination of self-peptides during periods of inflammation or oxidative stress could lead to the formation of neo-epitopes that, through the same mechanisms described herein for gp33/34, could escape the constraints of central tolerance. The impact of these modified de-novo MHC complexes on the initiation of unwanted T cell responses that may result in auto-reactivity remains to be studied.

Acknowledgments

We gratefully acknowledge access to synchrotron beam lines 14.2 and 14.4 at ESRF (Grenoble, France), MX-41 at BESSY (Berlin, Germany) and I19 at MAX II (Lund, Sweden).

Author Contributions

Conceived and designed the experiments: CM ADD TS AA. Performed the experiments: CM ADD. Analyzed the data: CM ADD TS JRW AA. Contributed reagents/materials/analysis tools: JRW AA. Wrote the paper: CM TS JRW AA.

References

1. Deri\'e YL, Pavan T, Dikic I (2010) Post-translational modifications in signal integration. Nature structural & molecular biology 17: 666–672.
2. See BT, Dikic I, Zhou MM, Pavan T (2006) Reading protein modifications with interaction domains. Nature reviews Molecular cell biology 7: 473–483.
3. Doyle HA, Mamula MJ (2001) Post-translational protein modifications in antigen recognition and autoimmune. Trends Immunol 22: 443–449.
4. Uy R, Wold F (1977) Posttranslational modification of proteins. Science 198: 280–288.
5. Ribet D, Costantini P (2010) Pathogen-mediated posttranslational modifications: A re-emerging field. Cell 143: 694–702.
6. Utz PJ, Anderson P (1998) Posttranslational protein modifications, apoptosis, and the bypass of tolerance to autoantigens. Arthritis Rheum 41: 1152–1160.
7. Engelhard VH, Altrich-Vanlith M, Ostankovitch M, Zarghali AL (2006) Post-translational modifications of naturally processed MHC-binding epitopes. Current opinion in immunology 18: 92–97.
8. Petersen J, Purrell AW, Rosjohn J (2009) Post-translational modified T cell epitopes: immune recognition and immunotherapy. Journal of molecular medicine 87: 1045–1051.
9. Meadows L, Wang W, den Haan JM, Bökland E, Reinhardus C, et al. (1997) The HLA-A*0201-restricted HIV-antigen contains a posttranslationally modified cysteine that significantly affects T cell recognition. Immunity 6: 273–281.
10. Pierce RA, Field ED, den Haan JM, Caldwell JA, White FM, et al. (1999) Cutting edge: the HLA-A*0101-restricted HY minor histocompatibility antigen originates from DFFRY and contains a cysysteinated cysteine residue as identified by a novel mass spectrometric technique. J Immunol 163: 6360-6364.
11. Backlund J, Carlsten S, Hoger T, Holm B, Fugger L, et al. (2002) Predominant selection of T cells specific for the glycosylated collagen type II epitope (263-270) in humanized transgenic mice and in rheumatoid arthritis. Proc Natl Acad Sci U S A 99: 9960–9965.
12. Haurum JS, Hoier IB, Arequell G, Næs E, Valencia G, et al. (1999) Presentation of cytosolic glycosylated peptides by human class I major histocompatibility complex molecules in vivo. J Exp Med 190: 143–150.
13. Andersen MH, Bonfill JE, Neisig A, Arsequell G, Sondergaard I, et al. (1999) Phosphorylated peptides can be transported by TAP molecules, presented by class I MHC molecules, and recognized by phosphopeptide-specific CTL. J Immunol 163: 3812–3818.
14. Zargar AL, Ficarro SB, White FM, Shabanovice J, Hunt DF, et al. (2000) Phosphorylated peptides are naturally processed and presented by major
histocompatibility complex class I molecules in vivo. J Exp Med 192: 1755–1762.

15. Abello N, Kerstjens HA, Postma DS, Bischoff R (2009) Protein tyrosine nitration: selectivity, physiochemical and biological consequences, denitration, and proteomics methods for the identification of tyrosine-nitrate containing proteins. Journal of proteome research 8: 3222–3238.

16. Aukal KS, Miyagi M, Yan L, West KA, Masillion D, et al. (2001) Proteomic method identifies proteins nitrosated in vivo during inflammatory challenge. Proc Natl Acad Sci USA 98: 12056–12061.

17. Ohnori H, Kanyamay N (2005) Immunogenicity of an inflammation-associated product, tyrosine nitro-derivatives of self-proteins. Autoimmunity 38: 224–229.

18. Burkhardt H, Kalden JR, Schultze-Koops H (2001) Chicken egg in autoimmunity and joint inflammation. Trends Immunol 22: 291–293.

19. Drew B, Liversidge J, Verwoerd C (2002) Aging and the role of reactive nitrogen species. Ann N Y Acad Sci 959: 46–81.

20. Cuzzocrea S, Chatterjee PK, Mazzeno E, McDonald MC, Dugo L, et al. (2002) Beneficial effects of GW517430, a novel, potent, and selective inhibitor of NOX activity, in a rodent model of collagen-induced arthritis. Eur J Pharmacol 453: 119–129.

21. Loeser RF, Carlson GS, Del Carlo M, Cole A (2002) Detection of nitrotyrosine in aging and osteoarthritic cartilage: Correlation of oxidative damage with the presence of interleukin-beta and with chondrocyte resistance to insulin-like growth factor 1. Arthritis Rheum 46: 2349–2357.

22. Beckmann JS, Ye YZ, Anderson PG, Chen J, Accavitti MA, et al. (1994) Extensive nitration of protein tyrosines in human atherosclerosis detected by polyclonal antibody. Circ Res 75: 81–88.

23. Shishibue MH, Haze SL (2004) Inflammatory and oxidative markers in atherosclerosis: relationship to outcome.Curr Atheroscler Rep 6: 243–250.

24. Cerillo A, Mercuri F, Quaglia L, Assaloni R, Motz E, et al. (2001) Detection of nitrotyrosine in the diabetic plasma: evidence of oxidative stress. Diabetologia 44: 834–838.

25. Liversidge J, Dick A, Gordon S (2002) Nitric oxide mediates apoptosis through a mechanism involving guanylyl cyclase and cGMP-dependent protein kinase. Am J Physiol 283: H597–H603.

26. ter Steege JC, Koster-Kamphuis L, Van Straten EA, Forre PT, Buurman WA (1998) Nitrotyrosine in plasma of celiac disease patients as detected by a new sandwich ELISA. Free Radic Biol Med 25: 953–963.

27. Noiri E, Nakao A, Uchida K, Tsukahara H, Ohno M, et al. (2001) Oxidative and nitrosative stress in acute renal ischemia. Am J Physiol Renal Physiol 281: F997–F1004.

28. Smith KJ, Kapoor R, Felts PA (1999) Demyelination: the role of reactive oxygen and nitrogen species and tyrosine nitration in the respiratory tract: epiphenomena or a damage linked to neurodegeneration by selective alpha-synuclein nitration in animal models. Brain Pathol 9: 69–92.

29. Giasson BI, Duda JE, Murray IV, Chen Q, Souza JM, et al. (2000) Oxidative stress and nitrosative stress in acute renal ischemia. Am J Physiol Renal Physiol 281: F1755–F1762.

30. Drew B, Leeuwenburgh C (2002) Aging and the role of reactive nitrogen species. Brain Pathol 9: 69–92.

31. Pieper GM, Khanna AK, Roza AM (2002) Prolonging organ allograft survival: current perspectives and future directions. Transplant Rev (Orlando) 16: 90–96.

32. Burkhardt H, Kalden JR, Schulze-Koops H (2001) Chicken egg in autoimmunity and joint inflammation. Trends Immunol 22: 291–293.

33. Liversidge J, Dick A, Gordon S (2002) Nitric oxide mediates apoptosis through a mechanism involving guanylyl cyclase and cGMP-dependent protein kinase. Am J Physiol 283: H597–H603.

34. Birnboim HC, Lemay AM, Lam DK, Goldstein R, Webb JR (2003) Cutting edge: MHC class II-restricted peptides containing the inflammation-associated terminal group substituted by a methyl group. Proteins 33: 97–106.

35. Achour A, Harris RA, Persson K, Sundborn J, Sentman CL, et al. (1999) Macrophage MHC-I-Restricted Nitrotyrosinated Neoantigen
Peptide and Major Histocompatibility Complex Protein Flexibility: IMPLICATIONS FOR VACCINE DESIGN. The Journal of biological chemistry 286: 40163–40173.

69. Ischiropoulos H, al-Mehdi AB (1995) Peroxynitrite-mediated oxidative protein modifications. FEBS Lett 364: 279–282.

70. Ischiropoulos H, Zhu L, Chen J, Tsai M, Martin JC, et al. (1992) Peroxynitrite-mediated tyrosine nitration catalyzed by superoxide dismutase. Arch Biochem Biophys 298: 431–437.