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Kinetics of MHC-CD8 Interaction at the T Cell Membrane

Jun Huang,* Lindsay J. Edwards, † Brian D. Evavold, † and Cheng Zhu2*‡

CD8 plays an important role in facilitating TCR-MHC interaction, promoting Ag recognition, and initiating T cell activation. MHC-CD8 binding kinetics have been measured in three dimensions by surface plasmon resonance technique using purified molecules. However, CD8 is a membrane-anchored, signaling kinase-linked, and TCR-associated molecule whose function depends on the cell membrane environment. Purified molecules lack their linkage to the membrane, which precludes interactions with other structures of the cell as well as signaling. Furthermore, three-dimensional binding in the fluid phase is biologically and physically distinct from two-dimensional binding across apposing cell membranes. As a first step toward characterizing the molecular interactions between T cells and APCs, we used a micropipette adhesion frequency assay to measure the adhesion kinetics of single mouse T cells interacting with single human RBCs coated with MHC. Using anti-TCR mAb we isolated and characterized the specific two-dimensional MHC-CD8 binding from the trimolecular TCR-MHC-CD8 interaction. The TCR-independent MHC-CD8 interaction has a very low affinity that depends on the MHC alleles, but not on the peptide complexed to the MHC and whether CD8 is an αα homodimer or an αβ heterodimer. Surprisingly, MHC-CD8 binding affinity varies with T cells from different TCR transgenic mice and these affinity differences were abolished by treatment with cholesterol oxidase to disrupt membrane rafts. These data highlight the relevance and importance of two-dimensional analysis of T cells and APCs and indicate that membrane rafts play an important role in modulating the affinity of cell-cell interactions. The Journal of Immunology, 2007, 179: 7653–7662.


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3 Abbreviations used in this paper: Lck, Src kinase p56\(^{Lck}\); EAS45, experimental additive solutions 45; pMHC, peptide MHC; SPR, surface plasmon resonance; 2D, two dimension(a); 3D, three dimension(a).

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Indeed, the former is referred to as two-dimensional (2D) interaction and the latter as three-dimensional (3D) interaction, which are physically distinct (31). The equilibrium dissociation constant \(K_d\) is expressed as the number of molecules per unit space based on the mass action law. In the fluid phase, molecules are brought together by diffusion and/or flow in a volume, the concentration has a unit of M, and the 3D binding affinity is measured in M\(^{-1}\). By comparison, molecules anchored on the cell membrane diffuse along a surface, their site density has a unit of \(\mu m^{-2}\), and the 2D binding affinity is measured in \(\mu m^2\). More importantly, the two interacting cells have to be brought together because membranes separated by a distance greater than the molecular size would physically preclude binding. It has been demonstrated that the molecular length and orientation (32) as well as cell surface microtopology and stiffness can significantly affect 2D (but not 3D) affinity (33, 34). Although the reverse-rate \(k_r\) has the same unit (in s\(^{-1}\)) for 2D and 3D interaction, only in the 2D (but not 3D) case can \(k_r\) be regulated by force applied to the molecular bonds—they would be disrupted by the separation of the cells (31).

As a first step toward characterizing the 2D molecular interactions between T cells and APCs, we used a micropipette adhesion frequency assay (35) to measure the TCR-independent 2D kinetics of CD8 on the T cell membrane interacting with MHC coated on the surface of RBCs. Consistent with published SPR results, the 2D MHC-CD8 interaction was of very low affinity and depend on the surface of RBCs. Consistent with published SPR results, the affinity was disrupted by the separation of the cells (31).

For every micropipette experiment, site densities of pMHC on RBCs and CD8 on T cells were measured by flow cytometry (Fig. 1). To measure the pMHC site density, RBCs were incubated with PE-conjugated mAb for 30 min at 4°C, washed three times with PBS, incubated with titrated Biotin-X-NHS (Calbiochem) according to the manufacturer’s instruction. In brief, RBCs were washed three times with PBS, incubated with titrated Biotin-X-NHS at pH 7.2 for 30 min at room temperature, and washed five times with PBS/1% BSA to remove Biotin-X-NHS and stop the reaction. The biotinylation efficiency was checked via flow cytometry by using PE-conjugated anti-biotin mAb. After three washes with EAS45/1% BSA, biotylated RBCs were stored in EAS45 solution for further use within 3–4 wk.

For each experiment, biotinylated RBCs were incubated with streptavidin (Pierce) at 0.5 mg/ml for 30 min at 4°C, washed three times with EAS45/1% BSA to remove unbound streptavidin, and incubated with biotinylated pMHC monomers for 30 min at 4°C. After washing three times with EAS45/1% BSA, the pMHC coated RBCs were ready for site density determination and micropipette adhesion assay.

Determination of molecular density on cell surface

For every micropipette experiment, site densities of pMHC on RBCs and CD8 on T cells were measured by flow cytometry (Fig. 1). To measure the pMHC site density, RBCs were incubated with PE-conjugated mAb for 30 min at 4°C, washed three times with PBS/1% BSA to remove unbound streptavidin, and incubated with biotinylated pMHC monomers for 30 min at 4°C. After washing three times with EAS45/1% BSA, the pMHC coated RBCs were ready for site density determination and micropipette adhesion assay.

Micropipette adhesion frequency assay

2D kinetics of MHC-CD8 interactions were measured using a micropipette adhesion frequency assay modified from that described previously (35). In brief, a pMHC-coated RBC and a T cell were aspirated by two apposing micropipettes with respective diameters of 1.5 and 3 \(\mu m\) to observe initial contact. Surprisingly, the 2D affinity for the same MHC varied with T cells from different TCR transgenic mice on which the CD8 density was expressed as a log transformation of Equation 1 at

\[ P_a = 1 - \exp[-m \cdot t \cdot K_0^2 \cdot (1 - \exp[-k_d \cdot t])], \]

1 to estimate a pair of parameters: a zero-force reverse-rate, \(k_d^0\), and an effective binding affinity, \(A_s \cdot K_0^2\), where \(A_s\) is the contact area (which was kept constant in all experiments). Means and SDs of \(k_d^0\) and \(A_s \cdot K_0^2\) were calculated from their individual values estimated from different \(P_a\) vs. \(t\) curves corresponding to different \(m\) and \(m\) for each pMHC.

For each pMHC, 30 pairs of cells were used to obtain several \(P_a\) vs. \(t\) curves that correspond to different CD8 and pMHC densities, \(m\) and \(m\). Each binding curve was fitted using nonlinear regression to the following probabilistic kinetic model for simple single monovalent receptor-ligand interaction (35),

\[ \ln[1 - P_a(\tau)] = A_s \cdot K_0^2 \cdot m \cdot \tau \]

2
which predicts that the transformed adhesion frequency, \( \ln[1 - P_e(t)]^{-1} \), is proportional to the product of the CD8 and pMHC densities, \( m_p m_L \), with the effective binding affinity, \( A_i K_i^0 \), as the constant of proportionality. The second form is to normalize the adhesion frequency by dividing the molecular site densities after the log transformation, but keep the time dependence:

\[
\ln(1 - P_e(t))^{-1} m_p m_L = A_i K_i^0 \left[1 - \exp(-K_i^0 t)\right].
\]

Applying the log transformation and normalization according to Equation 3 is predicted to collapse a family of \( P_e \) vs \( t \) curves corresponding to different \( m_p \) and \( m_L \) into a single curve, provided that they correspond to the same set of kinetic reverse-rate and effective affinity.

### Results

**Isolation of MHC-CD8 binding from the trimolecular TCR-MHC-CD8 interaction**

The micropipette adhesion frequency assay is illustrated in Fig. 2. A mouse T cell and a human RBC were respectively aspirated by two apposing micropipettes (Fig. 2A). Cells were then brought into controlled contact (Fig. 2B), and subsequently moved apart to determine whether an adhesion was present (Fig. 2C) or not (Fig. 2D) at the end of the contact period. To determine binding kinetics in this mechanical manner, the receptor-ligand bond involved has to be able to sustain a minimum force detectable by the RBC pico-force sensor (35). This is indeed the case for the MHC-CD8 interaction in question. An adhesion event between a T cell and RBC was unambiguously observed from the RBC membrane elongation, which resulted from the molecular force anchoring the RBC apex to the T cell (Fig. 2C). This was distinctly different from a nonadhesion event in which the RBC membrane readily separated from the T cell as the micropipette retracted (Fig. 2D). Although in any particular contact test both positive (i.e., adhesion) as well as negative (i.e., no adhesion) outcomes were possible and random, the probability of adhesion \( P_e \) was determined by the contact area \( A_i \), and time \( t \), the densities of receptors \( m_p \) and ligands \( m_L \), as well as the reverse-rate \( K_i^0 \) and binding affinity \( K_i^0 \), as predicted by Equation 1. Rather than adhesion force, the adhesion frequency assay measures this probability by counting adhesion frequency in repeated contact tests (35).

CTLs express both TCR and CD8, which may bind different sites on the same or different pMHCs (12, 18, 37). A given TCR is capable of binding different peptides complexed with MHC of the same allele with different kinetic rates and affinities, which can differentially trigger T cell activation (10, 11). To isolate MHC-CD8 binding from trimolecular TCR-MHC-CD8 interactions, an anti-TCR Vα2 mAb (B20.1) was used to block TCR-MHC binding. Also, an anti-CD8α blocking mAb (CT-CD8α) was used to confirm that, apart from a low level background, the measured binding was predominately due to specific MHC-CD8 interaction.

T cells expressing a monoclonal TCR from transgenic mice were preincubated with 50 μg/ml anti-TCR Vα2 or 10 μg/ml anti-CD8α for 30 min at 4°C and micropipette assays were performed in the continuous presence of the same concentrations of mAbs. In this study, TCR-independent, MHC-CD8 specific interactions were isolated for the three types of T cells (OTI, P14, and F5) used in this study, and for all pMHCs tested, including the peptides listed in Tables I and II bound to either H-2Kb or H-2Db MHC alleles. Some of the results are exemplified in Fig. 3 using H-2Kb complexed with three peptides with various properties for OTI T cells—VSV (null), R4 (antagonist), and OVA (agonist) (10, 37).

For the null peptide VSV (Fig. 3A), micropipette adhesion tests in the absence of Ab yielded an adhesion frequency that increased with contact duration initially then reached a steady state (△). Addition of the anti-CD8α mAb completely abolishing binding at

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4 The online version of this article contains supplemental material.
all contact durations tested (○), suggesting that the measured adhesion between T cells and RBCs was solely mediated by the specific MHC-CD8 interaction. This was further supported by the lack of effect of the anti-TCR Vα2 mAb which yielded a binding curve (△) indistinguishable from that without Ab (□). Similar results were obtained for another null peptide K4 (data not shown).

The inability to detect TCR binding indicated that OTI TCR did not recognize the null peptides. This agreed with results from other assays including SPR measurement (10, 11), synapse formation, and fluorescence resonance energy transfer measurement (37). The results also indicated that anti-TCR Vα2 mAb did not sterically hinder MHC-CD8 interactions. Thus, binding of RBCs bearing MHC complexed with null peptides to T cells is measurable, TCR-independent, and MHC-CD8 specific. We also tested the effect of anti-TCR Vα2 mAb on binding between P14 T cells and RBCs coated with H-2Kβ VSV pMHC and obtained similar results (data not shown). These data allowed us to use null pMHCs to measure the specific MHC-CD8 interaction in future experiments without using anti-TCR mAbs.

For antagonist peptide R4 (Fig. 3B), micropipette adhesion tests in the absence of Ab yielded much higher adhesion frequencies at all contact durations compared with VSV at equivalent site densities (□, compare Fig. 3, A and B). These adhesions, especially those at contact durations >2 s, were also stronger, as it was more difficult to separate the RBC from the T cell. When the anti-TCR Vα2 mAb was added, however, the adhesion frequency dropped to levels that matched those mediated by the binding of CD8 to VSV pMHC at corresponding contact durations (△, compare Fig. 3, A and B). Thus, there was a significant contribution from the TCR-MHC interaction to the measured adhesion when the antagonist R4 was used. This also ruled out the possibility that, when the peptide was VSV, the lack of blocking was due to problems with the anti-TCR Vα2 mAb (Fig. 3A). Addition of anti-TCR Vα2 and anti-CD8α mAbs to block both TCR and CD8 completely abrogated adhesion (○), suggesting that binding in the presence of anti-TCR Vα2 alone (△) was mediated solely by specific MHC-CD8 interaction. Similar data were obtained for another antagonist V-OVA (data not shown). Thus, both TCR and CD8 bind MHC complexed with antagonist peptides R4 or V-OVA, and the specific MHC-CD8 interaction can be isolated by blocking the TCR-pMHC interaction using the anti-TCR Vα2 mAb.

For the agonist peptide OVA (Fig. 3C), micropipette adhesion tests in the absence of Ab yielded adhesion in every single test (i.e., 100% adhesion frequency) at all contact durations (□). These adhesions were even stronger than those mediated by R4 pMHC, as it was sometimes impossible to separate the RBC from the T cell, despite the fact that the pMHC densities on RBCs were matched for all three peptides (VSV, R4, and OVA). Also, addition of the anti-CD8α mAb alone did not lower the frequency of adhesions (○), although they appeared somewhat weaker. This indicates that OTI TCR interacted with agonist (OVA) pMHC much more strongly than antagonist (R4 or V-OVA) pMHC. When the anti-TCR Vα2 mAb was added, however, the adhesion frequency dropped to levels similar to those mediated by binding of CD8 to VSV-loaded MHC at corresponding contact durations (△, compare Fig. 3, A and C). Again, addition of mAbs to block both TCR and CD8 completely abrogated adhesion (○), suggesting that binding in the presence of anti-TCR Vα2 alone (△) was mediated by specific MHC-CD8 interaction. Thus, both TCR and CD8 bind MHC complexed with agonist pMHC, and the specific MHC-CD8 interaction can be isolated by blocking the TCR-pMHC interaction using the anti-TCR Vα2 mAb.

Table I. Kinetic parameters for OTI TCR-independent, CD8-mediated binding to H-2Kβ MHC

<table>
<thead>
<tr>
<th>H-2Kβ pMHC</th>
<th>m (μm⁻²)</th>
<th>k⁺ (s⁻¹)</th>
<th>A_rk⁺ (10⁻⁶ μm⁻²)</th>
</tr>
</thead>
<tbody>
<tr>
<td>VSV</td>
<td>305</td>
<td>820</td>
<td>1.53 ± 0.37</td>
</tr>
<tr>
<td>VSV</td>
<td>297</td>
<td>715</td>
<td>0.88 ± 0.48</td>
</tr>
<tr>
<td>VSV</td>
<td>297</td>
<td>126</td>
<td>1.63 ± 1.01</td>
</tr>
<tr>
<td>VSV</td>
<td>233</td>
<td>1661</td>
<td>1.58 ± 0.30</td>
</tr>
<tr>
<td>VSV</td>
<td>326</td>
<td>1254</td>
<td>2.52 ± 0.61</td>
</tr>
<tr>
<td>VSV</td>
<td>370</td>
<td>434</td>
<td>1.53 ± 0.24</td>
</tr>
<tr>
<td>VSV</td>
<td>404</td>
<td>544</td>
<td>2.22 ± 0.24</td>
</tr>
<tr>
<td>VSV</td>
<td>Combined data</td>
<td>1.53 ± 0.22</td>
<td>5.75 ± 0.31</td>
</tr>
<tr>
<td>V-0VA</td>
<td>404</td>
<td>158</td>
<td>0.87 ± 0.18</td>
</tr>
<tr>
<td>V-0VA</td>
<td>338</td>
<td>499</td>
<td>1.08 ± 0.34</td>
</tr>
<tr>
<td>V-0VA</td>
<td>233</td>
<td>1529</td>
<td>2.00 ± 0.17</td>
</tr>
<tr>
<td>V-SVV</td>
<td>Combined data</td>
<td>1.12 ± 0.15</td>
<td>6.66 ± 0.36</td>
</tr>
<tr>
<td>OVA</td>
<td>281</td>
<td>1659</td>
<td>0.68 ± 0.14</td>
</tr>
<tr>
<td>OVA</td>
<td>198</td>
<td>844</td>
<td>1.14 ± 0.25</td>
</tr>
<tr>
<td>OVA</td>
<td>Combined data</td>
<td>0.83 ± 0.12</td>
<td>3.51 ± 0.22</td>
</tr>
<tr>
<td>K4</td>
<td>290</td>
<td>648</td>
<td>1.04 ± 0.17</td>
</tr>
<tr>
<td>R4</td>
<td>276</td>
<td>761</td>
<td>1.55 ± 0.15</td>
</tr>
</tbody>
</table>

Table II. Kinetic parameters for OTI TCR-independent, CD8-mediated binding to H-2Dβ MHC

<table>
<thead>
<tr>
<th>H-2Dβ pMHC</th>
<th>m (μm⁻²)</th>
<th>k⁺ (s⁻¹)</th>
<th>A_rk⁺ (10⁻⁶ μm⁻²)</th>
</tr>
</thead>
<tbody>
<tr>
<td>gp33</td>
<td>253</td>
<td>1157</td>
<td>0.55 ± 0.17</td>
</tr>
<tr>
<td>HIV gag</td>
<td>326</td>
<td>1713</td>
<td>0.55 ± 0.12</td>
</tr>
</tbody>
</table>
As an additional control, biotinylated RBCs not coated with pMHC were tested against T cells, which produced <2% adhesion regardless of the T cell specificity (data not shown). Thus, no adhesion molecules other than pMHC on RBCs and TCR and CD8 on T cells contribute to the adhesion frequencies measured with the micropipette adhesion assay.

To rule out the possibility that anti-TCR Vα2 mAb might cause changes in CD8 expression, we incubated OTI T cells with anti-TCR Vα2 mAb (50 μg/ml) in chamber medium (L-15/5 mM HEPES/1% BSA) for 90 min, which was the time elapse of a typical micropipette experiment, then stained with PE-conjugated anti-CD8 mAbs to quantify the expressions of CD8α and CD8β by flow cytometry. We also stained T cells not preincubated with anti-TCR Vα2 mAb with the same anti-CD8α and anti-CD8β mAbs as a control. CD8 expression was indistinguishable (data not shown), indicating that anti-TCR Vα2 mAb did not alter CD8 expression. It must be stressed that to detect the low affinity specific MHC-CD8 interaction, we had to use very high MHC site densities (~1000 sites/μm²). In sharp contrast, ~10 sites/μm² of OVA pMHC were sufficient to produce a specific trimerolecular TCR-MHC-CD8 binding comparable to the TCR-independent MHC-CD8 binding (data not shown).

Stoichiometry of the MHC-CD8 interaction

The density m₀ of CD8 expressed on OTI T cells slightly varied from mouse to mouse. The density mᵢ of MHC coated on RBCs also varied depending on the coating conditions. How CD8 and MHC densities regulate the adhesion frequency Pₐ (through mass action) depends on the stoichiometry, or valency, of the MHC-CD8 interaction. To determine this valency, we took the natural log of the reciprocal of the frequency of no adhesion to yield ln(1 – Pₐ), then plotted vs the product of the CD8 and MHC site densities, m₀mᵢ, for H-2Kb MHC complexed with VSV (A), V-OVA (B), and OVA (C) peptides separately. The data from panels A–C were pooled in panel D on which data from two other peptides, R4 and K4, were also plotted. The error bars were computed from SEM of Pₐ, according to the Gaussian error propagation law. A straight line with zero y-intercept was fit to data in each panel. The goodness-of-fit was indicated by the R² values.

Evaluation of kinetic parameters for MHC-CD8 interactions

Having confirmed the 1:1 stoichiometry of MHC-CD8 interaction, Equation 1 was fit to the TCR-independent, CD8-mediated binding data to evaluate the kinetic parameters, kₒ and AₒKₒ₀, for various MHC-CD8 interactions studied in this work, as exemplified in Fig. 3. Table I summarizes the kinetic parameters so evaluated for OTI T cell CD8 interacting with H-2Kb MHC complexed with five peptides for each pair of CD8 and pMHC densities tested (indicated). It is evident that, for each peptide, the kinetic parameters evaluated from individually fitting different Pₐ vs t data sets agree well, despite the fact that they correspond to different m₀ and mᵢ levels, as expected from the monovalency of the MHC-CD8 interaction, which also demonstrates reproducibility of our assay. To further test the reliability of these best-fit parameter values, the mean kₒ and AₒKₒ₀ values were calculated and used, along with the corresponding m₀ and mᵢ values measured from independent flow cytometry experiments, to predict each Pₐ vs t data set, which shows excellent agreement, as exemplified in Fig. 5A for V-OVA.

MHC-CD8 interaction is peptide independent

Although differing in amino acid sequences and molecular weights, the five peptides listed in Table I were complexed with MHC of the same allele, H-2Kb. Cocrystal structures of pMHC:CD8 complexes show no contact between CD8 and the peptide (18). However, it is still possible that a peptide could influence MHC-CD8 interaction should its binding induce a conformational change in the MHC that would alter the CD8 binding site. In Fig. 4D, the ln[1 – Pₐ(∞)]⁻¹ vs m₀mᵢ data from all five peptides appear to line up, suggesting that their binding affinities (equal to

As an additional control, biotinylated RBCs not coated with pMHC were tested against T cells, which produced <2% adhesion regardless of the T cell specificity (data not shown). Thus, no adhesion molecules other than pMHC on RBCs and TCR and CD8 on T cells contribute to the adhesion frequencies measured with the micropipette adhesion assay.
the slope of the line) have similar values. In Fig. 3, the TCR-independent, CD8-mediated $P_a$ vs $t$ data (Δ) appear to have similar shapes and plateau levels regardless of the peptide used, indicating similar kinetic parameters. It can be seen from the binding affinities and reverse-rates listed in Table I that these are indeed comparable for all five H-2Kb pMHCs.

To visualize the peptide independence of the kinetic parameters for the TCR-independent, CD8-mediated binding to MHC, we make use of Equation 3, which predicts that the $\ln(1 - P_a)$ vs $t$ data depend only on the kinetic parameters. Data for MHC complexed with all five different peptides were plotted in Fig. 5A (points). The collapse of data demonstrates graphically that these interactions have comparable kinetic parameters. Conversely, similar kinetic parameters predict similar $\ln(1 - P_a)$ vs $t$ curves, as shown by using the mean best-fit $k_0^c$ and $A_cK_a^c$ values for each peptide to plot the right-hand side of Equation 3 (Fig. 5B, curves), which overlay regardless of the peptide involved.

To further test our hypothesis that the MHC-CD8 interaction is peptide independent, kinetic parameters were measured for OTI T cells interacting with RBCs coated with H-2Db MHC complexed with gp33 or HIV gag, which are null peptides not recognized by OTI TCR. As can be seen from Table II where data are listed, the corresponding $A_cK_a^c$ and $k_0^c$ values are very similar for the two peptides tested. Taken together, the data suggest that loading different peptides on MHC does not induce enough change in the CD8 binding site to affect the measured kinetic parameters of the MHC-CD8 interactions. In other words, the interactions between CD8 and pMHC are peptide-independent.

**CD8 binds different MHC alleles with distinct kinetics**

Using MHC tetramer staining and SPR experiment with purified molecules, Moody et al. found that CD8 bound H-2Kb MHC with higher avidity/affinity than H-2Db MHC (25). Our 2D kinetic data support this finding. As shown in Fig. 6, CD8 from OTI T cells and P14 T cells respectively bound H-2Kb MHC with 7- and 20-fold higher 2D effective binding affinities than H-2Db MHC, respectively (Fig. 6, left column, solid bars or H-2Db (open bars) were compared. Data were presented as mean ± SEM. ND = not detectable.

**FIGURE 5.** MHC-CD8 interaction is monovalent and peptide independent. A, Adhesion frequency was plotted against contact duration for each set of CD8 and V-OVA pMHC densities. Data (points) from three sets of site densities (indicated), each obtained from 15 to 29 pairs of OTI T cells and RBCs contacting 50 times each, were presented as mean ± SEM at each contact duration. Equation 1 was fit to each data set to evaluate a pair of kinetic parameters, $k_0^c$ and $A_cK_a^c$, the values of which are summarized in Table I along with the site densities. The mean values of $k_0^c$ and $A_cK_a^c$ (see Table I) were then used to predict theoretical binding curves for all three set of site densities (curves). The excellent agreement between experiment and theory supported the assumption on which Equation 1 was based, that the MHC-CD8 interaction is monovalent. B, Mean adhesion frequency data were transformed according to the left-hand side of Equation 3 and plotted vs the contact duration (points). The right-hand side of Equation 3 predicts that binding curves so plotted would depend only on the $k_0^c$ and $A_cK_a^c$ values of the interactions involved.

**FIGURE 6.** Dependence of MHC-CD8 binding on MHC alleles and T cell types. 2D effective binding affinity (left column) and kinetic reverse-rate (right column) of naive CTLs from three transgenic mice, OTI (A), P14 (B), and F5 (C), interacting with either H-2Kb (solid bars) or H-2Db (open bars) were compared. Data were presented as mean ± SEM. ND = not detectable.
The same MHC allele binds CD8 from different T cells with different affinities

Unexpectedly, CD8 on T cells from different TCR transgenic mice was found to bind the same allelic pMHC with different affinities; the H-2Kb MHC binding affinity for CD8 on OTI T cells was 50% and 100% more than those on P14 T cells and from F5 T cells, respectively (compare the three panels in Fig. 6, solid bars, left column). A similar trend was found for the H-2Db MHC, which bound OTI T cell CD8 with an affinity 2- (compare A and B in Fig. 6, open bars, left column) and >10-fold more than those of P14 and F5 T cell CD8, respectively. To exclude the possibility that CD8 expression was changed by repeated touches with a pMHC coated RBC during the micropipette assay, OTI T cells were first incubated with 50 μg/ml anti-TCR V2 mAb for 30 min, mixed with OVA, R4 or VSV pMHC coated RBCs at a ratio of 1:100 in the presence of anti-TCR V2 mAb, pelleted the cells and incubated at room temperature for 10 min. After the RBCs were lysed with lysis buffer (eBioscience), the T cells were stained with anti-CD8α or β mAb to compare the CD8 expression levels. Flow cytometry data showed the CD8 expression level did not alter (Fig. 7A).

CD8αβ is believed to be a more potent coreceptor than CD8αα. Therefore, we hypothesized that CD8αβ bound MHC better than CD8αα, and that CD8 expressed on OTI, P14, and F5 T cells with different CD8αα/CD8αβ ratios. However, flow cytometry results showed that the CD8αα/CD8αβ ratios were comparable for those three types of T cells (data not shown). To further test this hypothesis, we took advantage of the fact that the site densities of CD8αα and CD8αβ varied mildly among OTI T cells purified from different transgenic mice, with site density ratios of CD8αα/CD8αβ ranging from 0.9 – 2.5 (indicated in the upper-left corner of each panel in Fig. 7B). In the preceding section, we have shown that the MHC-CD8 binding affinity and reverse-rate are independent of the peptide. Here we plot the 2D effective binding affinity

FIGURE 7. Lack of effect on CD8αα and CD8αβ expression. A, CD8α (α) and CD8β (β) expression levels on T cell were compared before interaction (open histogram) or after interaction with OVA (dash histogram), R4 (dot histogram) or VSV (dash-dot histogram) pMHC coated RBCs in the presence of anti-Vα2 mAb (50 μg/ml). Isotype control for nonspecific binding is shown as shaded histogram. Part of some histograms are obscured due to overlapping. B, Flow cytometric analysis for the expression of CD8α (filled histograms) and CD8β (open histograms) on 11 batches of OTI T cells. The ratio value of CD8αα/CD8αβ was indicated on each small panel. C, 2D effective binding affinity (A, kᶜα₀, upper panel) and kinetic reverse-rate (kᵣ₀, lower panel) measured using these 11 batches of OTI T cells were plotted against the CD8αα/CD8αβ ratio and compared with the average affinity or average reverse rate (dash lines). Data were presented as mean ± SEM.

FIGURE 8. Reduction of MHC-CD8 affinity by cholesterol oxidase treatment. A and B, 2D binding affinities of H-2Kb (A) or H-2Db (B) MHC for CD8 on three T cells with (open bars) or without (solid bars) cholesterol oxidase treatment. Data were presented as mean ± SEM. ND = not detectable. C, The CD8αα and CD8αβ expression levels of F5 T cells treated with 1 U/ml cholesterol oxidase (dash histogram) were compared with untreated T cells (open histogram, which overlaps with the dash histogram). Isotype control for nonspecific binding is shown as shaded histogram.
Table III.  Binding affinities of MHC-CD8 with (+) or without (−) cholesterol oxidase treatment

<table>
<thead>
<tr>
<th>T Cell</th>
<th>H-2Kb (−)</th>
<th>H-2Kb (+)</th>
<th>H-2Db (−)</th>
<th>H-2Db (+)</th>
</tr>
</thead>
<tbody>
<tr>
<td>OTI (10^−6 μm^−1)</td>
<td>5.75 ± 0.31</td>
<td>0.47 ± 0.15</td>
<td>0.51 ± 0.06</td>
<td>ND^a</td>
</tr>
<tr>
<td>P14 (10^−6 μm^−1)</td>
<td>3.67 ± 0.49</td>
<td>0.65 ± 0.24</td>
<td>0.18 ± 0.02</td>
<td>ND^a</td>
</tr>
<tr>
<td>F5 (10^−6 μm^−1)</td>
<td>2.80 ± 0.12</td>
<td>0.48 ± 0.25</td>
<td>ND^a</td>
<td>ND^a</td>
</tr>
</tbody>
</table>

^a Not detectable.

(Fig. 7C, upper panel) or reverse-rates (Fig. 7C, lower panel) measured using eleven batches of OTI T cells vs the CD8αα/CD8αβ ratio. It is evident that the two kinetic parameters A and k_d measured from different batches of T cells scatter evenly around a horizontal line equal to their respective average values (dotted line in each panel of Fig. 7C), suggesting a lack of dependence on the CD8αα/CD8αβ coreceptor potencies between CD8α and CD8αβ in activating T cells does not manifest as difference in their binding kinetics for MHC.

Disruption of membrane rafts differentially reduces affinity for MHC of CD8 from different T cells and abolishes the affinity differences

The micropipette frequency assay measured interactions between molecules anchored on the apposing membranes of a RBC and a T cell. These interactions can be impacted by the cell surface environment, as previously shown for membrane microtopology and stiffness (33, 34). Another aspect of the membrane environment may be membrane rafts into which CD8 can partition (4). Membrane rafts, and/or the CD8 partitioning therein, could vary among the three transgenic T cells used in our study, thus altering the apparent MHC-CD8 affinity. To test this hypothesis, we measured 2D binding affinities after disrupting membrane rafts by incubating T cells with 1 U/ml cholesterol oxidase (MP Biomedicals) for 30 min (42, 43). Disruption of membrane rafts substantially reduced the MHC binding affinities for CD8 on all three T cells for both H-2Kb (Fig. 8A, compare the solid and open bars in each group, and Table III) and H-2Db (Fig. 8B, adhesion became undetectable after cholesterol oxidase treatment despite that we used highest possible MHC site density, and Table III) alleles. This was not due to a decrease in the CD8 expression by the cholesterol oxidase treatment, as flow cytometry measurement using OTI or F5 T cells with and without such treatment showed identical CD8 expression levels (Fig. 8C). Also, treating RBCs with cholesterol oxidase alone did not affect MHC-CD8 binding (data not shown). Remarkably, cholesterol oxidase treatment reduced the H-2Kb MHC-CD8 binding affinity by different amounts in different T cells: 12-, 5.6-, and 5.8-fold in OTI, P14, and F5 cells, respectively. Thus, the H-2Kb MHC binding affinities for CD8 on different T cells were similar after rafts disruption (Fig. 8A, compare the three open bars and Table III). These results suggested that partitioning of CD8 in membrane rafts differentially enhanced its MHC binding affinity in the three T cells tested. The enhancement was abolished after the membrane rafts were disrupted, resulting in comparable CD8-MHC binding affinities regardless of the T cell specificity.

Discussion

The present work represents a first step toward dissecting the 2D molecular interactions between T cells and APCs, which involve a number of molecular players, including TCR, coreceptors, adhesion molecules, and costimulatory molecules. Some molecules provide physical linkages to bridge the two cell membranes and form junctional structures, while others deliver signals upon Ag recognition. Some interactions take place between binding partners respectively residing on apposing membranes of the T cell and the APC, while others occur between molecules both residing on the same membrane. Kinetic rates and binding affinity are important determinants for these molecular interactions, for such parameters are believed to correlate with T cell activation. Most published work used SPR for kinetic measurement, which uses purified molecules that have been isolated from their native T cell surface environment (3, 4, 10, 11, 23–25, 44, 45). Some studies use MHC tetramer or MHC-Ig dimeric molecules to stain T cells (6, 7, 16, 25, 46–50), but still measure 3D kinetic parameters. A few experiments measured 2D binding by a simple cell adhesion assay (21, 37, 51). However, this assay lacks sufficient temporal resolution to measure the kinetics of MHC-CD8 interactions. A different method has been used to measure 2D affinity, which visualizes bond formation between a receptor-expressing cell and a glass-supported lipid bilayer reconstituted with fluorescently labeled, lipid-anchored, freely mobile ligands (52–54). At physiological receptor and ligand densities, thousands of bonds are typically formed in a time scale of ~10 min, which is a large enough number of bonds and sufficiently long time to smoothen the self-assembled contact area. By comparison, the micropipette method detects binding in seconds on a rough T cell surface full of microvilli. These differences probably underlie the orders of magnitude of differences in the 2D affinities measured by the supported bilayer method and the micropipette method (31). Moreover, it required extension for this method to be used for measuring 2D kinetics (TP Tolentino, J Wu, VI Zarnitsyna, F Ying, ML Dustin, and C Zhu, unpublished data). Our paper reports the first 2D kinetic measurements using T cell CD8 to interact with pMHC presented on the membrane of RBCs. The measured effective binding affinities of TCR-independent MHC-CD8 interactions are much lower than those of selectin-ligand interactions (32, 55) and the high affinity integrin-ligand interactions (56) but are similar to the IgG binding affinities of the low affinity FcγRs CD16b (57) and CD32a (58).

There are two known receptors on the surface of CTLs for MHC: TCR and CD8. Multiple lines of evidence suggest that CD8 facilitates or enhances TCR to recognize Ag and initiate T cell activation, and it was shown that after abolishing the CD8 binding by making a mutation on the α3 domain of MHC, not only the tetramer staining efficiency for T cell was reduced, but the T cell activation was dramatically diminished (48). However, the detail mechanisms of cooperation between CD8 and TCR are not yet clear (1). One hypothesis proposes that binding of one molecule (CD8 or TCR) holds MHC to an optimal configuration, thereby decelerating the dissociation of both molecules (CD8 or TCR) to MHC and the APC, while others occur between molecules both residing on the same membrane. Kinetic rates and binding affinity are important determinants for these molecular interactions, for such parameters are believed to correlate with T cell activation. More...
TCR and CD8 at the level of MHC binding. Although SPR experiments did not find significant difference in the binding of solution MHC to immobilized TCR in the presence or absence of solution CD4/8 (3, 44), these negative data cannot rule out the possibility of cooperation when both TCR and CD8 reside on the same T cell membrane. The data in Fig. 3 show that inhibiting the TCR-MHC interaction by a blocking mAb does not result in detectable change in the kinetics of MHC binding to CD8 at a resting state. In other words, TCR-MHC and MHC-CD8 interactions appear to be independent at the level of adhesion measured by the present assay and the adhesion levels mediated by these two interactions are additive rather than synergistic, which provides a partial answer to the question of cooperation.

Cocrystal structures of both human and mouse pMHC:CD8 complexes reveal that CD8 mainly binds to the invariant α3 domain of the H chain and to the β2-microglobulin subunit of MHC (12, 18). By comparison, cocrystal structures of TCR:pMHC complexes reveal that TCR interacts with the α1 and α2 domains of the H chain of MHC where the peptide cleft is located (2, 59). Thus, TCR and CD8 bind to spatially separated sites on MHC. Crystallographic studies also reveal little structural differences in MHC complexed with different peptides (2, 59). It is therefore not surprising to find that the kinetic parameters of the TCR-independent, CD8-mediated MHC binding are peptide independent (Figs. 3, 4D, and 5B, Tables I and II). Indeed, this conclusion has been supported by previous studies using SPR measurement (24) and the cell adhesion assay (37). Thus, unlike the TCR-MHC interaction, MHC-CD8 interaction is independent on the peptide potency in triggering T cell activation.

By comparison, different MHC alleles, such as H-2Kb and H-2Dβ, may have sufficient structural differences to impact their CD8 binding kinetics (25, 45). In support of this hypothesis, we found that CD8 bound H-2Kb with a much higher affinity than H-2Dβ for the three types of T cells tested (Fig. 6). This is consistent with previous SPR measurements, which found that binding of CD8 to H-2Kb MHC was at least 2-fold better than H-2Dβ (25). Similar results were obtained for HLA-CD8 interactions in a human system, where CD8αα bound differentially to different alleles of HLA with subtle conformational differences in the α3 domain (45). The agreement of the present results with previous observations obtained using MHC tetramers, CD8 transfectants, and SPR supports the validity of the micropipette assay as applied to the T cells.

Cocrystal structures of human CD8αα:HLA-A2 (12) and mouse CD8αα: H-2Kb (18) complexes show that CD8αα primarily binds the α3 domain of MHC asymmetrically in an Ab-like fashion. The first α subunit interacts with MHC α3 and β2-microglobulin domains, which comprises of >70% of the contact surface. The second α subunit engages less and interacts only with the MHC α3 domain (12, 18). Although no crystal structure of MHC:CD8αβ complex has been published yet, such structure has been modeled by replacing either CD8α subunit of CD8αα with a CD8β subunit (17). Based on the model, Chang et al. proposed that, although CD8αβ could bind MHC with two distinct orientations, the primary binding site was still the α3 domain of MHC and the basic interaction mode was the same as that of CD8αα (17). By comparison, Devine et al. suggested that only the CD8α subunit could make a major contact with MHC class I (60). Nevertheless, both CD8αα and CD8αβ interact with MHC and promote TCR-MHC recognition (2), yet CD8αβ is two orders of magnitude more efficient as a coreceptor than CD8αα (20). However, it is controversial as to whether the different coreceptor efficiencies correspond to different MHC-CD8 binding. Sun et al. found that CD8αα and CD8αβ mediated similar levels of cell-cell adhesion (21). Similarly, SPR measurement found that CD8αα and CD8αβ had similar 3D affinities for H-2Kb MHC (22, 23, 25). By comparison, MHC tetramer staining experiment found that CD8αβ mediated greater interactions with MHC than CD8αα, although this discrepancy might come from either the CD8 expressing transgenic mice or the H-2Dβ MHC used (16). Our data suggest that the different coreceptor functions of CD8αα and CD8αβ are not due to their different 2D binding for MHC as they bind the same MHC allele with comparable 2D affinities and off-rates (Fig. 7).

Surprisingly, we found that CD8 expressed on different T cells bound the same MHC allele with different affinities (Fig. 6), and cholesterol oxidase treatment differentially reduced these affinities in different T cells. Remarkably, the 2D MHC-CD8 binding affinities became comparable for cholesterol oxidase treated T cells. These data are consistent with the hypothesis that partitioning of CD8 into membrane rafts differentially enhances its 2D binding affinity for MHC depending on the rafts in a particular type of T cells. Further studies are required to fully test this hypothesis and understand its implications to the regulation of MHC-CD8 interaction.

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Disclosures

The authors have no financial conflict of interest.

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