



Universiteit
Leiden
The Netherlands

Functional aspects of the adaptive immune system in arthritis

Jansen, D.T.S.L.

Citation

Jansen, D. T. S. L. (2017, March 8). *Functional aspects of the adaptive immune system in arthritis*. Retrieved from <https://hdl.handle.net/1887/47913>

Version: Not Applicable (or Unknown)

License: [Licence agreement concerning inclusion of doctoral thesis in the Institutional Repository of the University of Leiden](#)

Downloaded from: <https://hdl.handle.net/1887/47913>

Note: To cite this publication please use the final published version (if applicable).

Cover Page



Universiteit Leiden



The handle <http://hdl.handle.net/1887/47913> holds various files of this Leiden University dissertation

Author: Jansen, D.T.S.L.

Title: Functional aspects of the adaptive immune system in arthritis

Issue Date: 2017-03-08

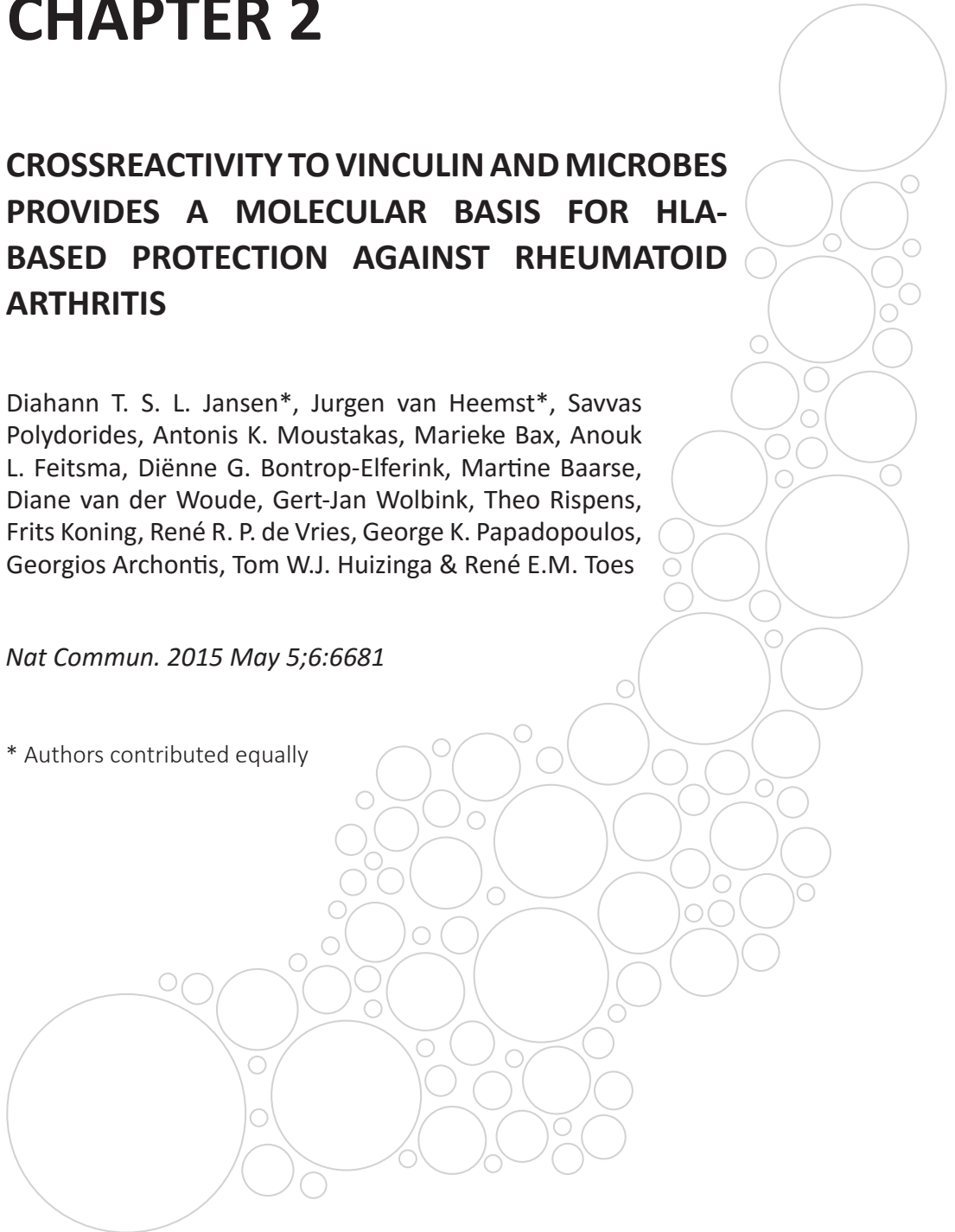
CHAPTER 2

CROSSREACTIVITY TO VINCULIN AND MICROBES PROVIDES A MOLECULAR BASIS FOR HLA-BASED PROTECTION AGAINST RHEUMATOID ARTHRITIS

Diahann T. S. L. Jansen*, Jurgen van Heemst*, Savvas Polydorides, Antonis K. Moustakas, Marieke Bax, Anouk L. Feitsma, Diënné G. Bontrop-Elferink, Martine Baarse, Diane van der Woude, Gert-Jan Wolbink, Theo Rispens, Frits Koning, René R. P. de Vries, George K. Papadopoulos, Georgios Archontis, Tom W.J. Huizinga & René E.M. Toes

Nat Commun. 2015 May 5;6:6681

* Authors contributed equally



ABSTRACT

The HLA locus is the strongest risk factor for anti-citrullinated protein antibody (ACPA)+ rheumatoid arthritis (RA). Despite considerable efforts in the last 35 years, this association is poorly understood. Here we identify (citrullinated) vinculin, present in the joints of ACPA+ RA patients, as an autoantigen targeted by ACPA and CD4+ T cells. These T cells recognize an epitope with the core sequence DERAA, which is also found in many microbes and in protective HLA-DRB1*13 molecules, presented by predisposing HLA-DQ molecules. Moreover, these T cells crossreact with vinculin-derived and microbial-derived DERAA epitopes. Intriguingly, DERAA-directed T cells are not detected in HLA-DRB1*13+ donors, indicating that the DERAA epitope from HLA-DRB1*13 mediates (thymic) tolerance in these donors and explaining the protective effects associated with HLA-DRB1*13. Together our data indicate the involvement of pathogen-induced DERAA-directed T cells in the HLA-RA association and provide a molecular basis for the contribution of protective/predisposing HLA alleles.

INTRODUCTION

Rheumatoid arthritis (RA) is a chronic autoimmune disease affecting synovial joints that can lead to severe disability. Pivotal pathophysiological insight has been obtained by the identification of anti-citrullinated protein antibodies (ACPA)^{1,2}. These autoantibodies target proteins that have undergone a post-translational conversion of arginine to citrulline, catalysed by peptidylarginine deiminases (PAD enzymes)³. ACPA are highly specific for RA, enriched in the joints of patients and can crossreact between citrullinated antigens that are expressed in the inflamed joints⁴⁻⁸. It is clear now that RA represents two main syndromes, ACPA+ and ACPA- disease, each with distinct genetic and environmental risk factors and disease outcome⁹⁻¹². Characteristics of ACPA (for example, isotype usage and epitope spreading) indicate the involvement of CD4+ T cell help in shaping the ACPA response¹³.

The most important genetic risk factor for ACPA+ RA is the HLA class II locus and risk is confined to a region with genes encoding for the beta chain of HLA-DR and the alpha and beta chain of HLA-DQ that are in tight linkage disequilibrium (LD) and inherited in haplotypes^{14,15}. An understanding of the HLA class II association and the relative contribution of the HLA-DR and HLA-DQ locus has been lacking for the last 35 years.

Next to the association of the predisposing alleles to ACPA+ RA, other HLA molecules are associated with protection. These protective HLA alleles, mainly HLA-DRB1*13, carry the five amino-acid sequence DERAA at positions 70-74 of the beta chain and protect also in the presence of predisposing alleles^{16,17}. Intriguingly, protection by these alleles is transferred from mother to child, supporting an active protective role of these alleles, possibly via microchimeric cells influencing thymic selection of CD4+ T cells, and indicating a dominant role of HLA-DRB1*13 in disease protection¹⁸.

HLA-derived peptides are a dominant peptide source presented by HLA class II molecules. We therefore proposed that the protective effect of HLA-DRB1*13 is explained by presentation of an HLA-DRB1*13-derived peptide in the context of other (predisposing) HLA class II molecules¹⁹⁻²¹.

It was previously shown that the degradation of HLA-DRB1*13 can result in the presentation of a peptide with the core sequence DERAA by other HLA class II molecules to CD4+ T cells²². This could allow for the negative selection of such 'DERAA-directed' CD4+ thymocytes. Interestingly, the DERAA sequence is also found in many microbes and in the self-protein vinculin. Vinculin is expressed in the synovium and was recently shown to be citrullinated in the synovial fluid of an RA patient^{4,23}. Likewise, T cells directed to vinculin are found under certain infectious conditions, indicating that T cell tolerance to vinculin is not absolute^{24,25}. Molecular mimicry of self-proteins with pathogenic proteins was proposed as an important mechanism to break T cell tolerance^{26,27}. Therefore, we postulate that on priming of DERAA-directed T cells by microbes expressing DERAA-containing proteins, T cells crossreactive to vinculin would be able to provide help to B cells reactive to citrullinated vinculin. This would ultimately result in the production of ACPA. In HLA-DRB1*13-positive donors, these T cells

are conceivably deleted, leading to protection against ACPA+ disease.

Here we show that citrullinated vinculin is novel autoantigen for ACPA antibodies. In addition, we demonstrate the presence of a T cell population in HLA-DRB1*13-negative donors that specifically recognize a DERAAs-containing vinculin epitope and that crossreact with DERAAs sequences derived from pathogens.

METHODS

Cells and sera

HLA-typed buffy coats from healthy volunteers were obtained from the blood bank (Sanquin, The Netherlands). PBMCs and sera from RA patients were derived from patients participating in the Leiden Early Arthritis Clinic cohort³⁷. All RA patients fulfilled the American College of Rheumatology (formerly the American Rheumatism Association) 1987 revised criteria for the classification of RA. A total of 178 RA patients were used in the current analyses. Patient samples were compared with 80 control samples from healthy individuals also living in the Leiden area. PBMCs were isolated using a standard Ficoll procedure. The protocols were approved by the Leiden University Medical Center ethics committee and informed consent was obtained.

Peptides

Peptides were synthesized according to standard Fmoc (*N*-(9-fluorenyl)methoxycarbonyl) chemistry using a Syroil peptide synthesizer (Multi-SynTech, Witten, Germany). The integrity of the peptides was checked using reverse-phase high-performance liquid chromatography and mass spectrometry. For vinculin ELISPOT experiments, HLA class II-binding studies and T cell activation studies 15-mer vinculin epitope VCL₆₂₂₋₆₃₆ (VCL-DERAA; REEVFDERAAANFENH) was used. To determine the peptide-binding register, the VCL₆₂₂₋₆₃₆ peptide was modified: Truncated on the N- or C terminus or alanine-/arginine-substituted at different positions. For ELISPOT assays against common pathogens, pathogenic peptides were used derived from measles virus (SSRASDERAAHLPTS), influenza A virus (VFEFSDERAAANPIVP), human herpesvirus 7 (LAARADERAAFPDVG), *Bordetella pertussis* (SPNLTDER- AAQAGVT), *Staphylococcus aureus* (QDMNDDERAAALMAM), *Haemophilus influenzae* (RFHGDDERAAKVYEN), *Salmonella enteritidis* (PLMMDDERAAKVYEN) and *Propionibacterium acnes* (EEVFTDERAARLSHV). For the generation of T cell lines, these eight pathogenic peptides were pooled.

Cell lines

The following Epstein-Barr virus-transformed lymphoblastoid B cell lines were used: BSM (DRB1*04:01;DQA1*03:01;DQB1*03:02, IHW9032), BOLETH (DRB1*04:01;DQA1*03:01;DQB1*03:02, IHW9031), BM21 (DQA1*05:01;DQB1*03:01, IHW9043), JSM (DQA1*03:02;DQB1*03:01), APD (DRB1*13:01; DQA1*01:03;DQB1*06:03, IHW9291), BC34 (DRB1*13:02;DQA1*01:02;DQB1*06:04), WT8 (DQA1*01:02;DQB1*06:02, IHW9017), KAS116 (DRB1*01:01; DQA1*01:02; DQB1*05:01, IHW9003) and 721.82 (DR/DQ-negative)³⁸.

Cells were maintained in IMDM (Lonza) supplemented with 10% heat-inactivated fetal calf serum. JSM and BC34 cells are derived from healthy HLA-typed donors that were Epstein-Barr virus transformed.

HLA class II competitive peptide-binding assay

Peptide-binding assays were performed, as described previously³⁹. In short, cell lysates from HLA class II homozygous B-lymphoblastoid cell lines were incubated on SPVL3- (anti-HLA-DQ) or B8.11.2- (anti-HLA-DR) coated (10 µg/ml³) FluoroNunc 96-well plates at 4 °C overnight. Titration ranges of the tested peptides (0 to 300 µM) were mixed with a fixed concentration (0.6 µM) of biotinylated indicator peptide and added to the wells. Bound indicator peptide was detected using europium-streptavidin (Perkin Elmer, Boston, MA) and measured in a time-resolved fluorometer (PerkinElmer, Wallac Victor2). IC50 values were calculated based on the observed binding of the test peptide against the fixed concentration indicator peptide. The IC50 value depicts the concentration of test peptide required for a loss of 50% of the indicator peptide signal.

Enzyme-linked immunosorbent spot assay

PBMCs were incubated in 96-well Multiscreen HA plates (Millipore) precoated with 5 µg/ml anti-IFN γ capture antibody (clone 1-D1K; Mabtech) at 5×10^5 cells per well and stimulated in X-VIVO medium (LONZA) with 5 µg/ml peptide or with a mix of recall anti- gens consisting of 0.75 Lf/ml tetanus toxoid (Netherlands Vaccine Institute), 5 µg/ml tuberculin purified protein derivative (Netherlands Vaccine Institute) and 0.005% *Candida albicans* (HAL allergy). After 24 h incubation at 37 °C, plates were extensively washed and incubated with 0.3 µg/ml biotin-labelled anti-IFN γ detection antibody (clone 7-B6-1; Mabtech) for 2h at RT, with 1,000 x diluted ExtrAvidin-Alkaline phosphatase (Sigma-Aldrich) for 1h at RT and with 5-Bromo-4-chloro-3-indolyl phosphate/Nitro blue tetrazolium (Sigma-Aldrich). Spots were analysed using the BioSys Bioreader 3000pro. Circular spots with a size of 80-450 µm were included.

Flow cytometry

Polyclonal CD4+ T cell lines were generated by stimulating 3×10^6 PBMCs per well in a 24-well plate with 5 µg/ml peptide for 7 days in IMDM supplemented with 5% human serum (Sanquin). After 7 days, 1.5×10^6 autologous PBMCs per well were plated in 24-well plates for 2h. After 2h, nonadherent cells were removed and adherent cells were pulsed with 5 µg/ml peptide and used as feeders for 10^6 T cells. After 1h, 10 µg/ml brefeldin A was added. Cells were incubated overnight and used for intracellular cytokine staining. The cells were incubated with fluorochrome-conjugated antibodies recognizing CD4, (Clone RPA-T4; BD biosciences), CD14 (Clone 61D3; eBioscience) and CD25 (Clone M-A251; BD biosciences), after which they were permeabilized using CytoFix CytoPerm Kit (BD Biosciences). After washing, cells were incubated with PE-labelled anti-IFN γ or matching isotype control. Cells were taken up in 1% paraformaldehyde until flow cytometric acquisition. Flow cytometry was performed on FACS

Calibur (BD biosciences) or LSR II (BD biosciences). Analysis was performed using FACS Diva (BD biosciences) and FlowJo software.

T cell cloning

JPT57 was generated from an HLA-DRB1*04:05/01:01;DQ8/DQ5 donor. PBMCs were cultured for 7 days in the presence of 10 µg/ml of VCL-DERAA peptide and restimulated with VCL-DERAA-pulsed antigen-presenting cells. After 1 week, cells were restimulated with 150 U/ml rIL-2. After two rounds of restimulation, T cell lines were tested for their specificity. The wells responding to VCL-DERAA peptide were cloned in a limiting dilution of 0.3 cells per well resulting in the isolation of clone JPT57.

T cell clone D2C18 was generated from an HLA-DRB1*04:01;DQ2/DQ8-positive donor. CD4+ T cells and CD14+ monocytes were isolated from PBMCs using antibodies bound to magnetic beads from, respectively, Dynal and Miltenyi. CD4+ T cells were labelled with 1 µM CFSE (Invitrogen) and incubated in a 2:1 ratio with CD14+ monocytes with 30 µg/ml PathMix. After 6 days, CD3_{pos} CD4_{pos} CD25_{pos} CD14_{neg} DAPI_{neg} CFSE_{low} cells were sorted by FACS aria (BD). Isolated CD4+ T cells were rested in medium containing 20 IU/ml rIL-2 (Peprotech). After 3 days, the cells were cloned in a limiting dilution of 0.3 cells per well resulting in the isolation of CD4+ T cell clone D2C18.

T cell activation

To determine IFN γ production by T cell clones in response to peptide stimulation, 50,000 T cells were incubated in a 1:1 ratio with B-LCL lines pulsed with 10 µg/ml of peptide for 3-6h. After 3 days, the supernatant was collected and an IFN γ ELISA (eBioscience) was used to determine the concentration of IFN γ . For measurement of T cell proliferation, 50,000 T cells were incubated in a 1:1 ratio with irradiated (3000RAD) autologous PBMCs pulsed with 10 µg/ml of peptide for 3-6h in IMDM supplemented with 5% human serum. After 3 days, cells were cultured for 16-20h with [³H]thymidine (0.5 µCi per well). ³H incorporation was measured by liquid scintillation counting (1450 MicroBeta TriLux; PerkinElmer). Blocking experiments were performed by preincubating antigen-presenting cells for 1h with 20 µg/ml anti-DQ (SPVL3), anti-DR (B8.11.2) and anti-DP (B7.21) blocking antibodies.

Detection of anti-citrullinated vinculin antibodies

Citrullinated vinculin was generated by incubation of 50 µg vinculin protein (Sanbio) in a volume of 200 µl containing 0.1 M Tris-HCl pH 7.6, 0.15 M CaCl₂, and 10 U PAD4 (Sigma) for 4h at 37 °C. Unmodified vinculin protein was generated by incubation of vinculin with PAD4 without CaCl₂. Citrullinated and unmodified vinculin were loaded onto 10% SDS-polyacrylamide gels and transferred onto blotting membranes. Blots were blocked, washed and incubated in 1:500 diluted serum overnight at 4 °C. The sera were either ACPA+ or ACPA- as determined by ELISA. Blots were incubated with horseradish peroxidase-labelled rabbit anti-human IgG (Dako) and visualized with chemiluminescence (ECL, Amersham). To analyse

reactivity to vinculin by a monoclonal ACPA (anti-cFIB1.1, citrullinated fibrinogen), vinculin- or citrullinated vinculin-coated Nunc plates were incubated with 0.2 µg/ml anti-cFIB1.1 for 2h at room temperature²⁸. Bound antibody was detected using horseradish peroxidase-labelled rabbit anti-human IgG (Dako) and visualized with ABTS. Peptide ELISA was performed as described previously using biotinylated peptides coated on streptavidin-precoated plates⁴⁰.

B cell activation by JPT57 cells

B cells were isolated from PBMCs of HLA-DQ8-positive healthy donors or ACPA-positive RA patients by magnetic anti-CD19 beads (Invitrogen). B cells were cultured in IMDM supplemented with 10% fetal calf serum, penicillin, streptomycin and glutamax. From healthy subjects, 30,000 B cells were co-cultured with different number of JPT57 cells in the presence of 5 µg/ml anti-IgM (Jackson Immunoresearch Laboratories) and with 10 µg/ml VCL-DERAA or 1 µg/ml PHA in round-bottom 96-well plates. After 7 days, IgG production by B cells was determined using a total IgG ELISA (Bethyl laboratories). From RA patients, 20,000 B cells were pulsed with 10 µg/ml VCL-DERAA peptide and co-cultured with 20,000 JPT57 cells in the presence of 5 µg/ml anti-IgM in round-bottom 96-well plates. After 7 days, ACPA production was determined by ELISA measuring reactivity against the CCP2-peptide in individual wells (EuroDiagnostica).

Energy minimization

Molecular simulations of HLA-DQ8 (A1*0301/B1*302) and HLA-DQ7.3 (A1*0302/B1*0301) complexed with various peptides, experimentally shown to bind to these molecules, were carried out as previously described using the Discover Suite (programmes InsightII and Discover) of Accelrys (San Diego, CA, release of 2005) on a Silicon Graphics Fuel instrument, using a minimization approach previously described⁴¹, that is, 1,000 steps of the steepest gradient method, followed by 1,000 steps of the conjugate gradient method. Records of the energy of every step showed a continuous decrease in energy without any local minima and an energy asymptote for the last 300-400 steps of the conjugate gradient method. The base molecule was the crystal structure of HLA-DQ8 (A1*0301/B1*0302) with bound the insulin B11-23 peptide⁴². The region HLA- DQβ105-112 for which full coordinates were not available in the original data, was constructed by molecular replacement of the respective region from HLA-DR1⁴³, after superposition of the β-plated sheet regions of the two molecules in the α1β1 domains. The binding registers of the vinculin and the bound microbial peptides were decided from the binding of truncated peptides as well as Arg-substituted peptides in presumed anchor positions; energy minimization of successive registers confirmed the registers predicted from the binding data. The rotamers for the peptide residues were chosen from a library of rotamers provided by the software database, to have no molecular clashes with the residues of HLA-DQ8. Minimizations were carried out either at pH 5.4 (endosomal pH) or 7.4 (extracellular). There are no similarly charged residues (for example, Glu-Glu) with their charged groups so close to each other as to require that one of the residues be

uncharged. Occasionally runs were performed on a Silicon Graphics Octane instrument with previous releases of the same software with very similar results. Figures are drawn using the WebLabViewer v.3.5 and DSViewerPro software of Accelrys, the latter currently freely available on the web. In the figures a side view of the bound peptide with the eye level placed at the level of the peptide backbone and parallel to the HLA-DQ β -sheet floor is visualized with an atomic colour code as in the respective figure legend. In the Supplementary Figures, the $\alpha 1\beta 1$ domains of the modelled HLA-DQ molecules in complex with given bound peptides are depicted in van der Waals surface representation, with colour and depiction conventions identical to those for the other figures. Several visible residues from the HLA-DQ molecule in contact with the antigenic peptide and potential contact with a cognate TCR in canonical orientation are shown in stick form with a transparent surface (atomic colour code: oxygen, red; nitrogen, blue; hydrogen, white; carbon, orange; sulfur, yellow). The antigenic peptide in the groove is shown in space filling form with the same atomic colour code as in Figure 6D-G. Coordinates of all complexes shown in the various figures have been deposited in the Figshare repository under accession codes 1294716.

MD simulations

The simulated system consisted of the entire DQ8 molecule and the 13-residue peptide with the vinculin sequence Glu-Glu-Val-Phe-Asp-Glu-Arg-Ala-Ala-Asn-Phe-Glu-Asn. Titratable residues were assigned their most common ionization state at physiological pH, with the exception of DQ8 residue α Glu31, which was protonated. In the crystallographic structure of the DQ8:insulin complex, residues α Glu31 and β Glu86 are in direct contact⁴² and their geometry and interactions with nearby residues and a crystallographic water suggest that at least one of them is protonated. A similar conclusion was reached for a pair of Glu and Asp residues in the pocket P6 of DR and I-E molecules⁴⁴. We used the empirical model Propka⁴⁵ and a constant-pH Monte Carlo approach implemented in the program PROTEUS⁴⁶ to compute the pK of titratable groups in the crystallographic structure of HLA-DQ8, both in the absence and the presence of the peptides insulin and VCL-DERAA. Both methods agreed that the residues α Glu31 and β Glu86, near pocket P1, are strongly correlated; the predicted pKa values suggested that one of them should be protonated, most probably α Glu31.

The initial coordinates of the protein heavy atoms were taken from the crystallographic structure of the DQ8:insulin complex (PDB accession code 1JK8)⁴². The peptide main chain heavy atoms were placed at the corresponding coordinates of the insulin main chain. Hydrogens were positioned by the HBUILD algorithm of the CHARMM programme⁴⁷. The peptide side chain initial conformations were optimized with the program PROTEUS⁴⁵. An additional control simulation studied the DQ8:insulin complex; for this system, the initial coordinates of the protein and peptide were taken from the corresponding crystallographic structure (PDB accession code 1JK8)⁴².

The initial set-up of the simulation system was performed with the CHARMM- GUI interface⁴⁸. A total of 70 crystallographic waters of the DQ8:insulin complex were retained. For the vinculin

complex, crystallographic waters were minimized for 100 steps before the simulation, with the protein and peptide atoms kept fixed. The ligand complexes were immersed in a periodically replicated water box with the shape of a 117-Å truncated octahedron. Overlapping water molecules were omitted and 17 potassium anions were added (14 ions in the insulin complex), to neutralize the total charge. The final complex had 120,324 atoms (6,166 protein- ligand atoms); the insulin complex had 120,504 atoms (6,172 protein-ligand atoms).

The simulations employed the molecular mechanics program CHARMM c37b2 (ref. 47). Protein atomic parameters were taken from the CHARMM36 all-atom force field with a CMAP backbone phi/psi energy correction^{49,50}. Water parameters corresponded to the modified TIP3P water model^{51,52}. Electrostatic interactions were calculated without truncation by the particle-mesh Ewald method⁵³, with a parameter $\kappa = 0.34$ for the charge screening, and sixth-order splines for the mesh interpolations. The Lennard-Jones interactions between atom pairs were switched to zero at a cutoff distance of 12Å. The temperature was kept at $T = 300$ K by a Nose-Hoover thermostat⁵⁴ with a mass of 2,000 kcal mol⁻¹ps⁻² for the thermostat. The pressure was maintained at $P = 1$ atm with a Langevin piston^{55,56}. The piston mass was set to 1/20 of the total mass of the system and the collision frequency was set to 20 ps⁻¹. The classical equations of motion were solved by the leap-frog integrator. Bond lengths to hydrogen atoms and the internal water geometry were constrained to standard values via the SHAKE algorithm⁵⁷, implemented into CHARMM.

The structure was initially optimized by 100 energy minimization steps with the steepest-descent and adopted-basis Newton-Raphson algorithms. This was followed by an equilibration run, consisting of three 200-ps segments, in which the harmonic force constants were gradually lowered from 10 to 0 kcalmol⁻¹Å⁻². The production simulation had a duration of 3 ns (vinculin complex) and 4 ns (vinculin complex with ionized α Glu31 and insulin complex). A total of 300 and 400 snapshots were analysed for the vinculin- and insulin-DQ8 complex, respectively, extracted every 10 ps. All simulations were conducted with version c37b2 of the CHARMM programme⁴⁷. Hydrogen bond occupancies and averaged intermolecular interaction energies were computed by in-house scripts. Molecular visualization was performed using the program VMD⁵⁸.

The interaction energies of selected peptide-protein residue pairs were computed by the following equation^{59,60}.

$$\Delta G_{RR'}^{\text{inte}} = \underbrace{\sum_{i \in R} \sum_{j \in R'} (E_{ij}^{\text{Coul}} + E_{ij}^{\text{GB}})}_{\Delta G_{RR'}^{\text{polar}}} + \underbrace{\sum_{i \in R} \sum_{j \in R'} E_{ij}^{\text{vW}} + \sigma \sum_{i \in R, R'} S_i}_{\Delta D_{RR'}^{\text{nonpolar}}}$$

The first and second group of terms on the right-hand side of equation (1) describe, respectively, polar and non-polar interactions between R and R'; 'Coul' denotes Coulombic interaction, 'GB' denotes generalized born interaction, ΔS_i is the change in the solvent accessible surface area of atom on binding and σ is a surface tension coefficient. The residue-

pair interaction energies of equation (1) include solvent-mediated effects via the above GB and surface area terms. In the calculations of Supplementary Figure 15, R and R' are distinct ligand and protein residues.

We employed the GBSW Generalized Born model^{61,62}, as implemented in the CHARMM programme. The coefficient σ was set to 0.005 kcal mol⁻¹Å⁻², for consistency with the GBSW parameterization. To compute the GB contributions, we removed all waters and ions from the simulation system and set the charge to zero for protein and peptide atoms other than those belonging to residues R and R', respectively. The last term contains the difference in solvent accessible surface areas of groups R and R' in the complex and unbound states. Coordinates of all complexes shown in the various figures have been deposited in the Figshare repository under accession code 1294716.

RESULTS

Citrullinated vinculin is a novel target for ACPA

We speculate that the protective effects of HLA-DRB1*13 on the development of ACPA+ RA are related to a T cell response reactive to a sequence that is commonly present in the HLA-DRB1*13 molecule, microorganisms and self-proteins that are targeted by ACPA. Indeed, a 5 amino acid long HLA-DRB1*13-derived sequence (DERAA) is present in many microorganisms and a few self-proteins. Of these self-proteins, the citrullinated protein vinculin attracted our attention. Therefore, we first analysed whether citrullinated vinculin is recognized by ACPA. To this end, we citrullinated vinculin *in vitro* with PAD enzymes and tested both native and citrullinated vinculin for recognition with serum of an ACPA+ RA patient. We observed citrulline-specific recognition of vinculin by serum of an ACPA+ RA patient, but not with an ACPA- patient (Figure 1A). We further confirmed that citrullinated vinculin is a target of ACPA using an ACPA monoclonal antibody (Figure 1B)²⁸.

Recently, several reports showed the presence of citrullinated vinculin in the synovial fluid of ACPA+ RA patients. In addition, three sites of *in vivo* citrullination on this protein were identified: Arg285, Arg622 and Arg823^{4,23}. When we studied antibody responses to these citrullination sites, we could show recognition by 34% of tested sera from ACPA+ RA patients versus 5% of sera from ACPA- patients (Figure 1C). Sera of ACPA+ RA patients are highly (cross-) reactive towards multiple citrullinated antigens. Indeed, when we quantified responses to citrullinated alpha-enolase, fibrinogen, vimentin and myelin basic protein in patient sera that react to citrullinated vinculin peptides, we could readily demonstrate additional reactivities indicating that ACPA are not exclusively directed against citrullinated vinculin as expected (Figure 1D).

Together these data show that citrullinated vinculin is a self-protein recognized by RA autoantibodies in a citrulline-dependent fashion.

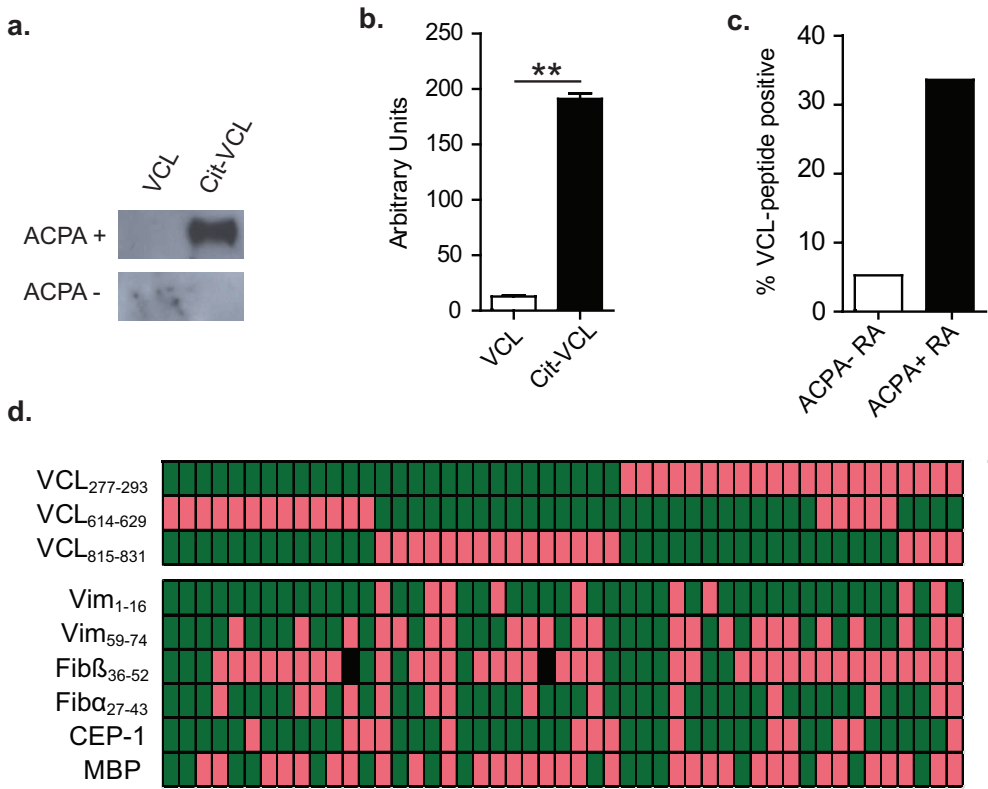


Figure 1. Citrullinated vinculin is a novel autoantigen targeted by ACPA. (A) Western blot (Supplementary Figure 22) of native and citrullinated vinculin protein, stained with the serum of an ACPA+ or an ACPA- RA patient. (B) Enzyme-linked immunosorbent assay (ELISA) of vinculin- or citrullinated vinculin protein-coated plates stained with ACPA monoclonal antibody anti-citFib1.1. Two-sided statistical analysis was performed using a Student's t-test with ** indicating $P < 0.001$. Error bar is S.E.M. (C) ELISA of citrullinated vinculin peptides VCL-CIT285_{277-293'}, VCL-CIT622_{614-651'}, VCL-CIT823_{815-831'} in serum of anti-CCP2⁺ ($n = 140$) versus anti-CCP2⁻ ($n = 38$) RA patients. Plot depicts the percentage of patient sera positive for one or more citrullinated vinculin peptides. (D) Heat map depicting all patients positive for one or more citrullinated vinculin epitopes ($n = 49$) and their serum reactivity towards different citrullinated epitopes derived from vimentin, fibrinogen and alpha-enolase. Red, positive; green, negative; black, undetermined. Error bars are S.E.M.

Vinculin is recognized by T cells from HLA-DRB1*13 negative donors

We next wished to determine whether the DERRA sequence from vinculin is recognized by human CD4⁺ T cells. To this end, peripheral blood mononuclear cells (PBMCs) were stimulated with a 15-mer vinculin-peptide (VCL_{622-636'} REEVFDERRAANFENH) (VCL-DERRA) for 24h. We observed a clear reactivity towards this epitope as determined in an interferon- γ (IFN γ)-ELISPOT-assay (Supplementary Figure 1). HLA-DRB1*13, especially HLA-DRB1*13:01, is strongly associated with protection against ACPA+ RA¹⁷. We hypothesize that HLA-DRB1*13 protects against ACPA+ disease by affecting the generation of DERRA-directed T cells. Therefore, we stratified donors for HLA-DRB1*13:01 status. We observed a striking difference

in IFN γ -producing cells depending on HLA-DRB1*13:01-status. The lack of reactivity in HLA-DRB1*13:01-positive donors was not due to a hampered ability of such donors to respond to T cell antigens as we observed strong response to microbial antigens in both HLA-DRB1*13:01 carriers and non-carriers when stimulated with recall antigens (Figure 2A). We further confirmed this finding in PBMCs stimulated for 4 days (Supplementary Figure 2). The differential ability of HLA-DRB1*13:01-positive donors to respond to VCL-DERAA is most likely not due to a general deficiency to present the VCL-DERAA epitope as these donors were heterozygous and thus expressed other HLA molecules that could potentially present VCL-DERAA. Nonetheless, to further confirm that HLA-DRB1*13:01 affects the ability to generate VCL-DERAA-directed T cell responses, we repeated the experiments in a set-up stratified for HLA using PBMCs from HLA-DRB1*04, HLA-DRB1*13:01 and HLA-DRB1*04/*13:01 heterozygous donors. A significant reduction in IFN γ -producing cells was observed in both HLA-DRB1*13:01 carriers and DRB1*04/*13:01 heterozygous donors as compared with HLA-DRB1*04 carriers. Again, no difference was observed for recall antigens (Figure 2B). These data indicate that the lack of detecting VCL-DERAA-directed T cells is not explained by the inability of HLA-DRB1*13 molecules to present VCL-DERAA, but rather the result of a dominant effect associated with the presence of HLA-DRB1*13.

Thus, vinculin is an autoantigen recognized by circulating VCL-DERAA-directed CD4+ T cells. In HLA-DRB1*13:01 carriers these T cells are absent. This effect, like the protective effects of HLA-DRB1*13:01 on the development of ACPA+ RA, was present in a dominant fashion, consistent with the notion that HLA-DRB1*13 affects the generation of DERAA-directed T cells, possibly during thymic selection.

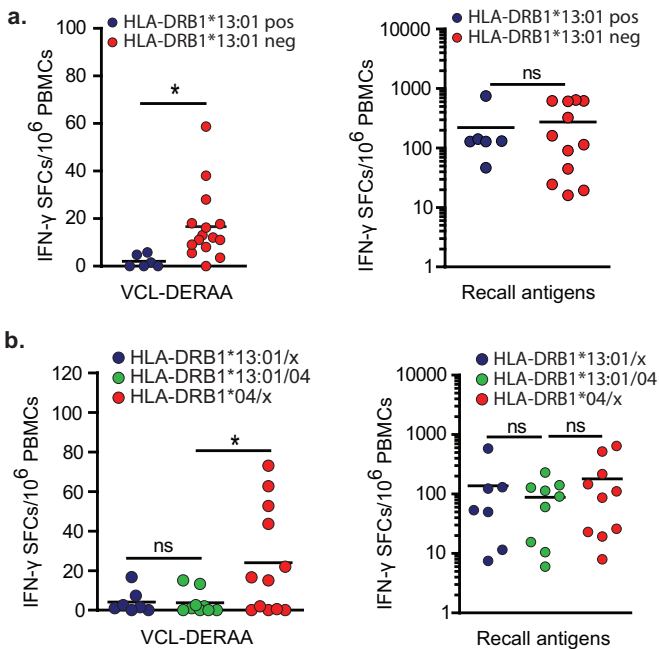


Figure 2. Citrullinated vinculin is a novel autoantigen targeted by CD4+ T cells in HLA-DRB1*13-negative donors. (A) IFN γ ELISPOT of PBMCs from healthy individuals stimulated for 24h with VCL-DERAA peptide REEVFDERAANFENH ($n = 21$) or with a mix of recall antigens (tetanus, tuberculin, candida, $n = 18$). Donors were stratified based on HLA-DRB1*13:01 status. (B) IFN γ ELISPOT of PBMCs of HLA-DRB1*13:01 carriers ($n = 7$), HLA-DRB1*04 carriers ($n = 12$) and HLA-DRB1*13:01/DRB1*04 donors ($n = 9$), stimulated for 24 h with VCL-DERAA peptide or with a mix of recall antigens. Each dot represents a unique donor. Two-sided statistical analyses of ELISPOT data were performed using a Mann-Whitney U-test with * indicating $p < 0.05$. Neg, negative; NS, not significant; Pos, positive

Presentation of Vinculin-DERAA by RA-predisposing HLA-DQ molecules

HLA-DRB1 alleles predisposing to ACPA+ RA in the Caucasian population are collectively called HLA shared epitope (SE) alleles^{14,15,29}. The most common HLA-SE alleles are HLA-DRB1*04:01 and HLA-DRB1*01:01. Using HLA-DR-binding assays, we examined the ability of VCL-DERAA to bind to these HLA molecules, but we could not detect any binding of VCL-DERAA to these alleles (Table 1, Supplementary Figure 3A, 3B). The HLA class II region is known for its strong LD. Genes that encode for the HLA-DR-beta chains are inherited in haplotypes with genes that encode for the alpha and beta chain of HLA-DQ. The predisposing HLA-SE alleles are in tight LD with HLA-DQ5 (DQA1*01;DQB1*05:01), HLA-DQ7.3 (DQA1*03:02;DQB1*03:01) and HLA-DQ8 (DQA1*03:01;DQB1*03:02) (Figure 3A, 3B). In Figure 3B, the distribution of HLA-DQ molecules in SE+ healthy donors is depicted, showing that these donors are HLA-DQ5-, HLA-DQ7.3- or HLA-DQ8-positive. We therefore next studied the binding of VCL-DERAA to these HLA-DQ alleles and observed binding of the VCL-DERAA peptide to HLA-DQ5, DQ7.3 and DQ8 (Table 1, Supplementary Figure 3C-E). These data indicate that VCL-DERAA can be presented by HLA-DQ molecules in LD with predisposing HLA-SE alleles.

To obtain an indication whether the VCL-DERAA can be presented by more HLA molecules, we also studied the ability of additional HLA-DR and HLA-DQ molecules encoded by HLA haplotypes that protect or have no influence on the development of ACPA+ RA. We could not detect binding of VCL-DERAA to these HLA class II alleles (Table 1, Supplementary Figure 3F-K). Likewise, using an IFN γ ELISPOT, VCL-DERAA-directed T cells were absent in donors negative for predisposing HLA-DQ molecules (Supplementary Figure 4).

These data indicate that HLA-DQ molecules that predispose to ACPA+ disease present the VCL-DERAA peptide, whereas the analysed HLA-DQ and HLA-DR molecules not associated with disease do not present VCL-DERAA.

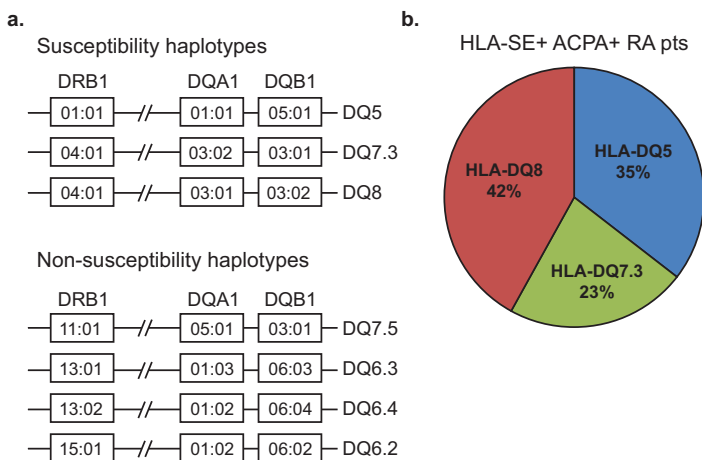


Figure 3. Presentation to vinculin-DERAA-directed T cells is restricted to RA-predisposing HLA-DQ molecules. (A) Schematic representation of the LD between the HLA-DRB1, HLA-DQA1 and HLA-DQB1 gene in studied haplotypes associated with susceptibility to ACPA+ RA. (B) Distribution of HLA-DQ molecules in ACPA+ RA patients positive for HLA-DR SE alleles ($n = 31$).

Table 1 | Presentation to vinculin-DERAA-directed T cells is restricted to RA-predisposing HLA-DQ molecules.

Association	Subtype	HLA allele	IC50 (μM)
Susceptibility haplotypes	HLA-DR	DRB1*01:01	> 300
		DRB1*04:01	> 300
	HLA-DQ	DQ5	39
		DQ7.3	11
Non-susceptibility haplotypes	HLA-DR	DRB1*13:01	> 300
		DRB1*13:02	> 300
	HLA-DQ	DQ6.2	> 300
		DQ6.3	> 300
		DQ6.4	> 300
		DQ7.5	> 300

Affinity of VCL-DERAA for associated HLA class II molecules. Affinity was determined by a competitive binding assay using high-affinity biotinylated peptides. Results are depicted as IC50 value, the concentration of test peptide (μM) where 50% of biotinylated peptide is bound. Experiments were performed at least three times and the average IC50 value between the different experiments is shown.

Table 1. Presentation of vinculin-DERAA-directed T cells is restricted to RA-predisposing HLA-DQ molecules.

Affinity of VCL-DERAA for associated HLA class II molecules. Affinity was determined by a competitive binding assay using high-affinity biotinylated peptides. Results are depicted as IC50 value, the concentration of test peptide (μM) where 50% of biotinylated peptide is bound. Experiments were performed at least three times and the average IC50 value between the different experiments is shown.

HLA-DQ-restricted recognition of VCL-DERAA by T cells

To further confirm the presence of VCL-DERAA-directed T cells and their HLA-restriction, we isolated CD4+ T cell clone JPT57, which was specific for VCL-DERAA. As shown in Supplementary Figure 5, this clone proliferated readily and produced large amounts of IFN γ when stimulated with the VCL-DERAA peptide.

When cultured with B cells from ACPA+ RA patients pulsed with the VCL-DERAA epitope, the clone not only upregulated CD40L, but also enhanced the production of ACPA in culture, indicating that such T cells have a phenotype compatible with the ability to provide 'help' to ACPA-producing B cells (Supplementary Figure 6).

To further confirm that HLA-DQ molecules present the VCL-DERAA peptide, we stimulated the clone with VCL-DERAA-pulsed HLA-typed antigen-presenting cells preincubated with HLA class II blocking antibodies (Supplementary Figure 7). In concordance with the binding studies, anti-HLA-DQ antibodies abrogated T cell recognition. Interestingly, we observed that JPT57 can recognize VCL-DERAA, presented by both HLA-DQ7.3 and HLA-DQ8 suggesting that this epitope is presented in a similar binding register by these HLA molecules (Supplementary Figure 8).

Together HLA class II presentation and T cell recognition of VCL-DERAA was restricted to HLA-DQ molecules that are associated with RA susceptibility.

Identification of microbe-specific T cells targeting DERAA epitopes

We next investigated the presence of the DERAA sequence in microorganisms. A Blast search showed that DERAA is found in 66% of bacteria and 4% of viruses (Figure 4A). This large number represents a major challenge to identify potential 'crossreactive microorganisms'. To select for relevant microorganisms, we restricted the search to those microorganisms that can cause disease or symptoms in humans and are present in the western world. This approach

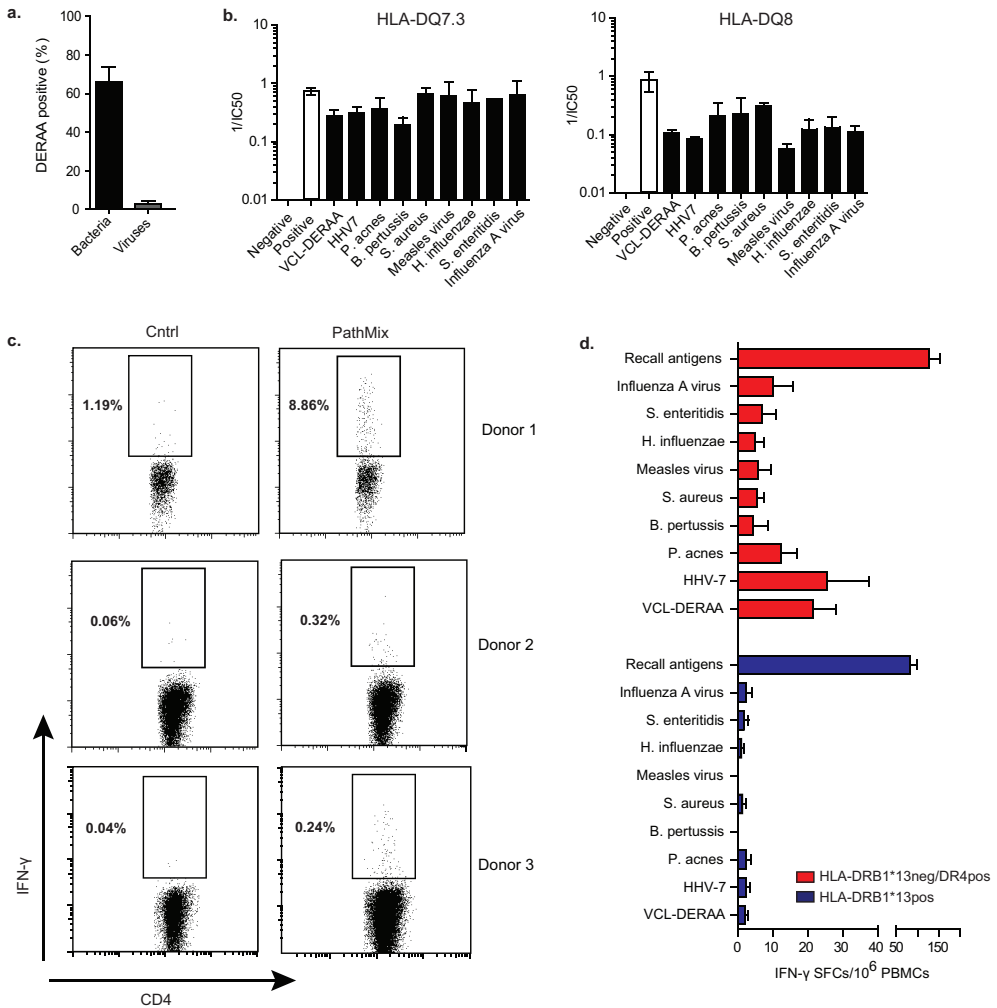


Figure 4. Identification of pathogen-specific T cells targeting DERRA epitopes in HLA-DRB1*13 negative donors. (A) Frequency of bacteria or viruses positive for a DERRA-containing protein as identified using the NCBI BLAST. List of all bacteria or viruses in blast database was generated and for three random sets of 50 bacteria/viruses a blast search for DERRA was performed. Error bars indicate the S.D. between the three analysed sets. (B) Competitive binding of pathogen-derived DERRA epitopes to HLA-DQ7.3 or HLA-DQ8. Binding experiments were performed at least three times and plots show pooled experiments, the error bars show the variation between the different experiments. (C) Flow cytometry staining of PathMix-directed T cell lines restimulated with PathMix pulsed or unpulsed antigen-presenting cells. Plots are gated on CD14neg/CD4pos/CD25pos T cells. (D) ELISPOT of PBMCs of healthy individuals of HLA-DRB1*13:01 donors ($n = 5$) and HLA-DQ7.3/DQ8 donors ($n = 8$) stimulated for 24h with DERRA-containing peptides and recall antigens. Error bars indicate the variation between different donors. Error bars are S.E.M.

left us with 219 candidate sequences. We then synthesized eight DERRA-containing peptides from common recall microbes including several bacteria (*P. acnes*, *S. enteritidis*, *B. pertussis*, *S. aureus*) and viruses (measles virus, influenza A and human herpesvirus 7). Interestingly, all of these peptides were presented by HLA-DQ7.3 and DQ8 molecules showing that these

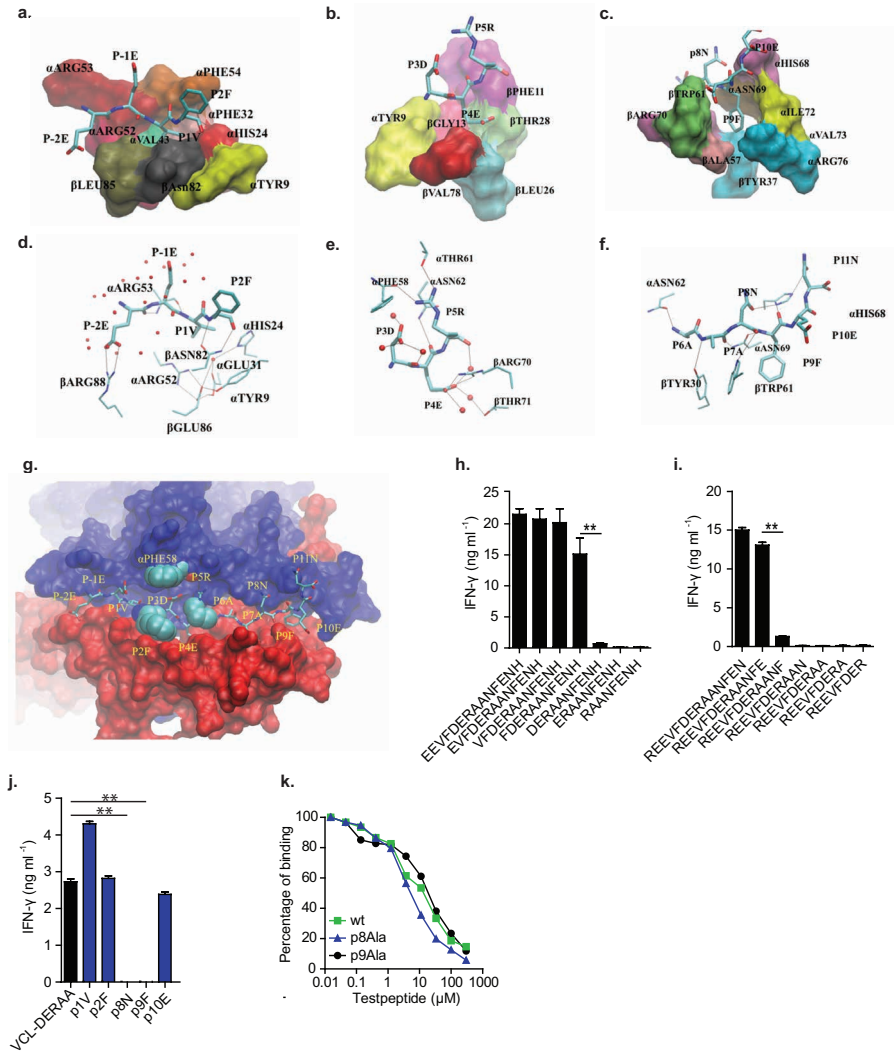
HLA molecules can efficiently accommodate both the VCL-DERAA epitope as well as microbe-derived DERAA epitopes (Figure 4B).

To determine whether these microbial-derived epitopes are recognized by human T cells, we generated T cell lines by stimulating PBMCs from three healthy HLA-DQ8-positive donors for 7 days with a pool of the eight pathogen-derived peptides (PathMix). Next we determined the presence of PathMix-specific CD4+ IFN γ -producing T cells. As shown in Figure 4C, such T cells were readily detectable. Using limiting dilutions, we isolated T cell clone D2C18, further confirming the presence of ‘microbial-DERAA-directed’ T cells (Supplementary Figure 9). Next we also analysed the presence of ‘microbial-DERAA- directed’ T cells by ELISPOT directly *ex vivo*, allowing the analyses of more donors at higher throughput. Interestingly, PBMCs from HLA-DRB1*13:01 carriers displayed a significantly reduced reaction against these peptides, indicating that the presence of HLA-DRB1*13 also affects the formation of T cell responses against DERAA epitopes from microbes (Figure 4D).

Together these data indicate that HLA-DQ alleles associated with ACPA+ RA efficiently present both VCL-DERAA and microbe-derived DERAA epitopes and that microbe-specific T cells directed to DERAA epitopes are readily detected. The presence of HLA-DRB1*13:01 affects the formation of these T cell responses.

Predicting crossreactive microbes by modelling vinculin-DERAA presentation

The data presented above demonstrate the presence of microbial- and VCL-DERAA-directed T cells, but do not show if a single T cell receptor (TCR) can react with both epitopes. To facilitate the search for possible ‘crossreactive’ DERAA epitopes out of all microbe-derived DERAA sequences, we first determined the binding register of VCL-DERAA using HLA class II-binding assays with amino (N)- and carboxyl (C)- terminally truncated and amino-acid-substituted VCL-DERAA peptides as detailed in the Supplementary Note and in Supplementary Figures 10-14. Together these experiments support **VFDERAANF** (anchors in bold) as the core binding register for both HLA-DQ7.3 and HLA-DQ8. All-atom molecular dynamics (MD) simulations showed that residues Val625 (P1), Glu628 (P4) and Phe633 (P9) make numerous intermolecular polar and non-polar interactions in the respective pockets. In Figure 5A-F, we highlight key intermolecular interactions between the VCL-DERAA epitope and the HLA-DQ8 molecule. A detailed discussion of the MD results and a quantitative assessment of the intermolecular interactions can be found in the supplementary data (Supplementary Figures 15 and 16, Supplementary Table 1 and Supplementary Note). The MD simulations also indicated that the long protruding side chains of Glu623 (P-2) Glu624 (P-1), Phe626 (P2), Asp627 (P3), Arg629 (P5), Asn632 (P8), Glu634 (P10) and Asn635 (P11) are exposed and could potentially interact with crossreactive TCRs (Figure 5G). Subsequently, we also further confirmed the obtained model using a second type of molecular modelling: energy minimization, which further confirmed possible TCR-contact residues (Supplementary Figures 17 and 18).



To functionally confirm whether a TCR would indeed interact with (some of the) potential TCR-contact residues within the VCL-DERAA peptide, we next determined how the epitope is recognized by the VCL-DERAA-directed T cell clone JPT57. Phe626 (P2) interacts with the JPT57-TCR as its removal results in a large decrease in recognition without affecting HLA-DQ8 binding. (Figure 5H, Supplementary Figure 12). C-terminal truncations resulted in a large decrease in T cell recognition after the removal of Glu634 (P10), showing that this residue is also important for JPT57-TCR interaction (Figure 5I). Thus, the data obtained using the VCL-DERAA-directed T cell clone as a functional read-out, are in line with the HLA-binding and molecular modelling studies indicating the sequence **VFDERAANFE** (anchors in bold) as the minimal epitope required for activation of JPT57. Next we performed alanine substitutions within the minimal epitope to remove critical TCR-interacting residues. Substituting Asn632 (P8) and Phe633 (P9) dramatically impacted T cell recognition, without affecting the binding affinity of VCL-DERAA (Figure 5J, 5K).

Together these data indicate **VFDERAANF** (anchors in bold) as the most likely core binding register for HLA-DQ7.3 and HLA-DQ8 and Asn632 and Phe633 as important residues for JPT57-TCR recognition.

TCR crossreactivity between vinculin and bacterial antigens

Identifying microbes that crossreact with vinculin was challenging due to the large number of potential candidates. Therefore, we used the data presented above to determine if a single TCR can crossreact both to DERAA sequences from microbes and the self-protein vinculin.

As we identified the Asn632 and Phe633 at P8 and P9, respectively, as important for JPT57-TCR interactions, we use the molecular modelling to predict which viral and bacterial epitopes are likely to bind to HLA-DQ8 and harbor P8Asn or P9Phe and are thus likely to crossreact with JPT57. From the 15 peptides identified (Supplementary Table 2), three were able to activate JPT57 in a T cell stimulation assay at concentrations similar to those used for the vinculin peptide. These epitopes were derived from *Campylobacter coli*, *Lactobacillus curvatus* and *Lactobacillus sakei* and crossreacted with JPT57 in an HLA-DQ-dependent manner (Figure 6A). Peptide-binding studies revealed that these three peptides bind with an intermediate binding affinity to HLA-DQ7 and HLA-DQ8 (Figure 6B-6C). Molecular models of these bacterial peptides in HLA-DQ8 illustrate the similarities with VCL-DERAA (Figure 6D-G, Supplementary Figures 19-21).

Thus, these data show that the RA-susceptibility alleles HLA-DQ7 and HLA-DQ8 can present both VCL-DERAA and related microbe-derived epitopes to T cells and that such T cells can be crossreactive to vinculin and bacterial epitopes, thereby providing an explanation for the presence of activated self-reactive CD4+ T cells directed to vinculin in peripheral blood.

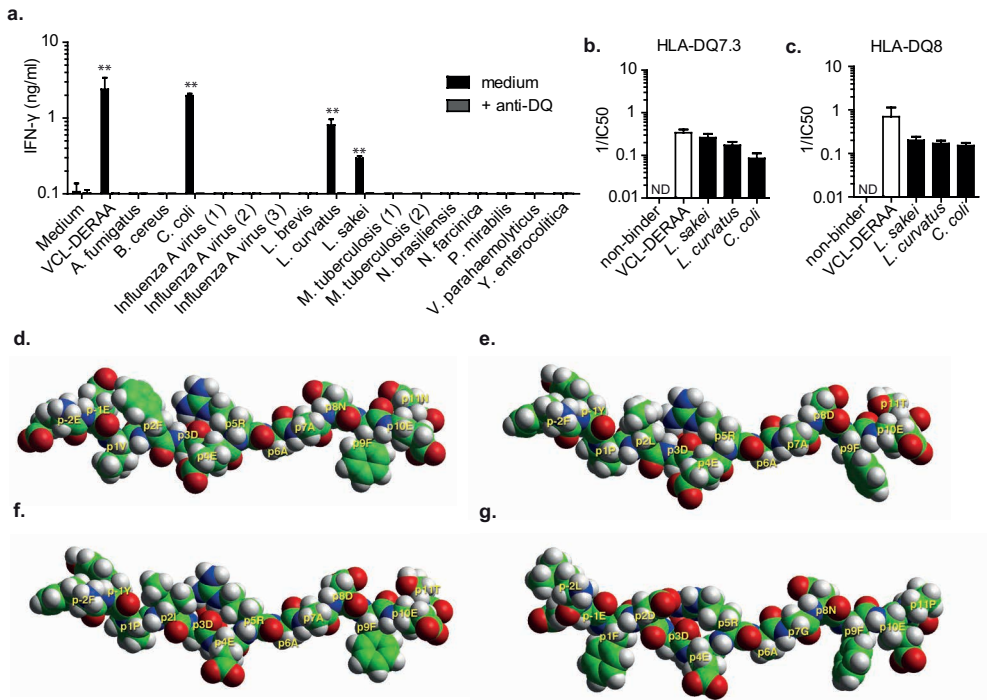


Figure 6. TCR crossreactivity of vinculin and bacterial DERAAs epitopes. (A) IFN γ ELISA on supernatant of T cell clone JPT57 stimulated with HLA-DQ8-positive B-LCLs and alanine-substituted VCL-DERAA peptides. This experiment was performed three times, the plot shows a representative experiment. Two-sided statistical analysis was performed using a Student's t-test with * indicating $p < 0.05$ and ** indicating $p < 0.001$. (B-C) Competitive binding of HLA-DQ7.3 (B) and HLA-DQ8 (C) to a non-binding negative control peptide, an unbiotinylated positive control peptide and epitopes from *Lactobacillus curvatus*, *Lactobacillus sakei* and *Campylobacter coli*. Binding experiments were performed three times. The error bars show the variation between the different experiments. (D-G) Side views of HLA-DQ8 complexed with the vinculin-DERAA peptide (D) and peptides derived from *Lactobacillus curvatus* (E), *Lactobacillus sakei* (F) and *Campylobacter coli* (G) in space-filling mode at pH 7.4 (extracellular), obtained by energy minimization as described in the methods section. Each peptide is shown in atomic colour code: oxygen, red; nitrogen, blue; hydrogen, white; carbon, green; sulfur, yellow. Error bars are S.E.M.

DISCUSSION

The strong connection between the HLA locus and RA has been known for more than 35 years. The complex HLA class II associations and the diverse ACPA responses in RA patients suggest the presence of multiple aetiological pathways. To unravel these pathways, identifying the relevant autoantigens is crucial. We here present evidence favouring the involvement of vinculin in the emergence of ACPA+ disease. This cytoskeletal protein was recently found to be citrullinated *in vivo* in the synovial fluid. We now show that it is recognized by ACPA as well, thereby adding it to a still selective list of targets. Moreover, we identified an epitope from vinculin recognized by CD4+ T cells restricted to HLA-DQ molecules predisposing to ACPA+ RA. The core amino-acid sequence (DERAA) is also present in many pathogens and in HLA-DRB1*13, a molecule encoded by an HLA locus associated with protection against ACPA+ RA.

We have also shown that a single TCR can recognize both a vinculin-DERAA epitope as well as DERAA epitopes from microbes, indicating the crossreactive nature of 'DERAA'-directed T cell responses. More importantly, such T cell responses appear absent from donors carrying HLA-DRB1*13 as DERAA-directed T cell responses to either pathogen- or vinculin-derived DERAA epitopes were lacking in these subjects. Even donors that harboured HLA-DRB1*13 next to predisposing HLA alleles were unable to respond to vinculin-DERAA.

Together these data indicate a novel pathway that explains several of the protective and predisposing HLA-effects associated with ACPA+ RA (Figure 7). In short, recognition of citrullinated vinculin by B cells will lead to the presentation of the 'VCL-DERAA' epitope in the context of HLA class II molecules. The HLA-DQ molecules genetically linked to the predisposing HLA-SE-molecules are particularly good in presenting DERAA-containing peptides. The DERAA-directed T cells primed against various pathogens harbouring DERAA-containing proteins cross-react with the VCL-DERAA peptide and provide help to the B cells, ultimately leading to a strong ACPA response. Subjects born with HLA-DRB1*13, will present the HLA-DRB1*13-derived DERAA-peptide in the thymus, leading to tolerization of the DERAA-directed T cell response and hence the inability to provide help to ACPA-producing B cells via this pathway and thereby the emergence of ACPA+ RA (Figure 7).

The variation of HLA-DR and HLA-DQ molecules in the human population is enormous. We have shown that predisposing HLA-DQ molecules are particularly good at presenting the VCL-DERAA epitope. Interestingly, the absence of VCL-DERAA affinity for a wide variety of other tested HLA-DR or HLA-DQ molecules could indicate a selective presentation by these HLA risk molecules. Next to a role of predisposing haplotypes, we also focussed on the protective effect of HLA-DRB1*13 alleles. However, the DERAA sequence can also be found in other HLA-DRB1 alleles (*04:02, *11:02, *11:03), which are rare in Caucasian populations. These alleles have previously been implicated in protection from ACPA+ RA, but their allele frequency hampers functional studies³⁰⁻³². Interestingly, it has been previously reported that the processing of these alleles results in the generation of a similar HLA-DERAA epitope suggesting that these alleles could all protect via the pathway that we have described²².

The place in time at which HLA-DR13 mediates protection from the development of ACPA and/or ACPA+ disease, or its relation to epitope spreading of the ACPA response is currently not known and would be relevant to determine in future studies. Recent evidence showed that the ACPA response matures before disease onset and that the HLA system could be involved in this^{13,33}. It is intriguing to speculate that infections such as by DERAA-containing microbes are involved in this expansion. Molecular mimicry of self-proteins with pathogenic proteins was proposed as a mechanism to break T cell tolerance, allowing the development of autoimmune disease^{26,27}. Interestingly, the DERAA sequence is also present in proteins from many (common) microbes allowing priming of DERAA-directed T cells. In mouse models, it was shown that low-avidity T cells to tissue-restricted antigens can persist without signs of anergy and unresponsiveness. Infection lowers the threshold for T cell activation

resulting in the induction of autoimmunity and memory formation³⁴. Infection could also induce autoimmunity via molecular mimicry of microbial proteins with self-proteins. We now identified crossreactive epitopes from the gut-residing bacteria *L. sakei*, *L. curvatus* and *C. coli*. It was shown that acute gastrointestinal infections can induce loss of T cell tolerance to (commensal) gut microbes, resulting in the activation of microbiota-specific T cells, their differentiation to inflammatory effector cells and formation of memory T cells³⁵. A recent study on the fecal microbiota of RA patients compared with controls demonstrated a significant increase in *Lactobacillus* species, together providing a rationale for a role of such bacterial species in the formation of DERAAs-directed T cell responses³⁶. Together our study provides a mechanistic clue on the HLA-RA connection, including both predisposing and protective HLA effect, and warrants further studies addressing the possibility to target DERAAs-directed T cells in the prevention of ACPA+ RA.

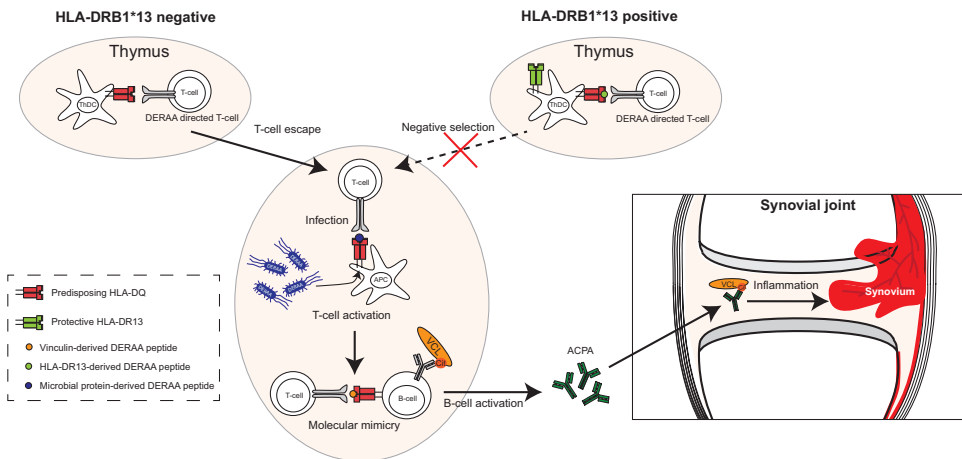


Figure 7. Schematic representation of the proposed role of DERAAs-directed CD4+ T cells in ACPA+ RA. DERAAs-directed T cells are restricted to RA-predisposing HLA-DQ5, -DQ7.3 and -DQ8 molecules. In carriers of these HLA-DQ molecules, DERAAs-directed T cells can become activated on contact with microbes. These activated T cells can subsequently crossreact with a DERAAs epitope derived from vinculin resulting in the activation of citrulline-directed B cells and the production of ACPA directed to citrullinated vinculin or vinculin-linked proteins. Citrullinated vinculin is present in the synovial compartment and is a target of ACPA. Binding of ACPA to its target can induce antibody-mediated effector mechanisms thereby contributing to synovial inflammation. In HLA-DRB1*13-positive individuals, a HLA-DRB1*13-derived DERAAs epitope is presented by predisposing HLA-DQ molecules to CD4+ T cells resulting in their negative selection, thereby protecting against the development of ACPA+ RA.

ACKNOWLEDGEMENTS

We thank B. Malissen (Centre d'Immunologie de Marseille-Luminy, Université de la Méditerranée, Marseille, France) for providing us with the B8.11.2 hybridoma. The Silicon Graphics Fuel instrument and the accompanying software were obtained via a grant from the Epirus Regional Development Programme to the Epirus Institute of Technology, through the third Community Support Framework of the European Union (80% European Union funds, 20% Hellenic state funds). This work was further supported by an NWO-ZonMW VICI grant from the Netherlands Organization for Scientific Research, by the IMI JU funded project BeTheCure, contract no 115142-2 and by a University of Cyprus grant to G.A. and S.P.

REFERENCES

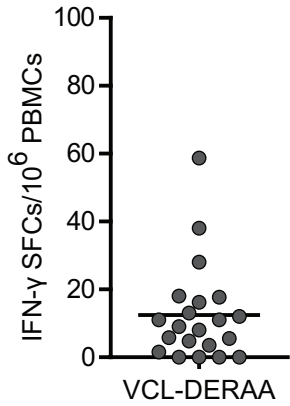
1. Schellekens, G. A., de Jong, B. A., van den Hoogen, F. H., van de Putte, L. B. & van Venrooij, W. J. Citrulline is an essential constituent of antigenic determinants recognized by rheumatoid arthritis-specific autoantibodies. *J. Clin. Invest.* 101, 273-281 (1998).
2. Girbal-Neuhausser, E. et al. The epitopes targeted by the rheumatoid arthritis-associated antifilaggrin autoantibodies are posttranslationally generated on various sites of (pro)filaggrin by deimination of arginine residues. *J. Immunol.* 162, 585-594 (1999).
3. Anzilotti, C., Pratesi, F., Tommasi, C. & Migliorini, P. Peptidylarginine deiminase 4 and citrullination in health and disease. *Autoimmun. Rev.* 9, 158-160 (2010).
4. van Beers, J. J. et al. The rheumatoid arthritis synovial fluid citrullinome reveals novel citrullinated epitopes in apolipoprotein E, myeloid nuclear differentiation antigen, and beta-actin. *Arthritis Rheum.* 65, 69-80 (2013).
5. Vossenaar, E. R. et al. The presence of citrullinated proteins is not specific for rheumatoid synovial tissue. *Arthritis Rheum.* 50, 3485-3494 (2004).
6. Ioan-Facsinay, A. et al. Anti-cyclic citrullinated peptide antibodies are a collection of anti-citrullinated protein antibodies and contain overlapping and non-overlapping reactivities. *Ann. Rheum. Dis.* 70, 188-193 (2011).
7. Amara, K. et al. Monoclonal IgG antibodies generated from joint-derived B cells of RA patients have a strong bias toward citrullinated autoantigen recognition. *J. Exp. Med.* 210, 445-455 (2013).
8. Snir, O. et al. Antibodies to several citrullinated antigens are enriched in the joints of rheumatoid arthritis patients. *Arthritis Rheum.* 62, 44-52 (2010).
9. van Gaalen, F. A. et al. Autoantibodies to cyclic citrullinated peptides predict progression to rheumatoid arthritis in patients with undifferentiated arthritis: a prospective cohort study. *Arthritis Rheum.* 50, 709-715 (2004).
10. Rantapaa-Dahlqvist, S. et al. Antibodies against cyclic citrullinated peptide and IgA rheumatoid factor predict the development of rheumatoid arthritis. *Arthritis Rheum.* 48, 2741-2749 (2003).
11. Huizinga, T. W. et al. Refining the complex rheumatoid arthritis phenotype based on specificity of the HLA-DRB1 shared epitope for antibodies to citrullinated proteins. *Arthritis Rheum.* 52, 3433-3438 (2005).
12. Verpoort, K. N. et al. Association of HLA-DR3 with anti-cyclic citrullinated peptide antibody-negative rheumatoid arthritis. *Arthritis Rheum.* 52, 3058-3062 (2005).
13. Willemze, A., Trouw, L. A., Toes, R. E. & Huizinga, T. W. The influence of ACPA status and characteristics on the course of RA. *Nat. Rev. Rheumatol.* 8, 144-152 (2012).
14. Stastny, P. Mixed lymphocyte cultures in rheumatoid arthritis. *J. Clin. Invest.* 57, 1148-1157 (1976).
15. Raychaudhuri, S. et al. Five amino acids in three HLA proteins explain most of the association between MHC and seropositive rheumatoid arthritis. *Nat. Genet.* 44, 291-296 (2012).
16. Shadick, N. A. et al. Opposing effects of the D70 mutation and the shared epitope in HLA-DR4 on disease activity and certain disease phenotypes in rheumatoid arthritis. *Ann. Rheum. Dis.* 66, 1497-1502 (2007).
17. van der Woude, D. et al. Protection against anti-citrullinated protein antibody-positive rheumatoid arthritis is predominantly associated with HLA-DRB1*1301: a meta-analysis of HLA-DRB1 associations with anti-citrullinated protein antibody-positive and anti-citrullinated protein antibody-negative rheumatoid arthritis in four European populations. *Arthritis Rheum.* 62, 1236-1245 (2010).
18. Feitsma, A. L. et al. Protective effect of noninherited maternal HLA-DR antigens on

- rheumatoid arthritis development. *Proc. Natl Acad. Sci. USA* 104, 19966-19970 (2007).
19. Chicz, R. M. et al. Predominant naturally processed peptides bound to HLA- DR1 are derived from MHC-related molecules and are heterogeneous in size. *Nature* 358, 764-768 (1992).
 20. Collado, J. A. et al. Composition of the HLA-DR-associated human thymus peptidome. *Eur. J. Immunol.* 43, 2273-2282 (2013).
 21. Adamopoulou, E. et al. Exploring the MHC-peptide matrix of central tolerance in the human thymus. *Nat. Commun.* 4, 2039 (2013).
 22. Snijders, A. et al. An HLA-DRBI-derived peptide associated with protection against rheumatoid arthritis is naturally processed by human APCs. *J. Immunol.* 166, 4987-4993 (2001).
 23. Romero, V. et al. Immune-mediated pore-forming pathways induce cellular hypercitrullination and generate citrullinated autoantigens in rheumatoid arthritis. *Sci. Transl. Med.* 5, 209ra150 (2013).
 24. di Marzo Veronese, F. et al. Autoreactive cytotoxic T lymphocytes in human immunodeficiency virus type 1-infected subjects. *J. Exp. Med.* 183, 2509-2516 (1996).
 25. Propato, A. et al. Apoptotic cells overexpress vinculin and induce vinculin- specific cytotoxic T cell cross-priming. *Nat. Med.* 7, 807-813 (2001).
 26. Oldstone, M. B., Nerenberg, M., Southern, P., Price, J. & Lewicki, H. Virus infection triggers insulin-dependent diabetes mellitus in a transgenic model: role of anti-self (virus) immune response. *Cell* 65, 319-331 (1991).
 27. Wucherpfennig, K. W. & Strominger, J. L. Molecular mimicry in T cell- mediated autoimmunity: viral peptides activate human T cell clones specific for myelin basic protein. *Cell* 80, 695-705 (1995).
 28. van de Stadt, L. A. et al. Monoclonal anti-citrullinated protein antibodies selected on citrullinated fibrinogen have distinct targets with different cross- reactivity patterns. *Rheumatology* 52, 631-635 (2013).
 29. Gregersen, P. K., Silver, J. & Winchester, R. J. The shared epitope hypothesis. An approach to understanding the molecular genetics of susceptibility to rheumatoid arthritis. *Arthritis Rheum.* 30, 1205-1213 (1987).
 30. de Vries, N., Tijssen, H., van Riel, P. L. & van de Putte, L. B. Reshaping the shared epitope hypothesis: HLA-associated risk for rheumatoid arthritis is encoded by amino acid substitutions at positions 67-74 of the HLA-DRB1 molecule. *Arthritis Rheum.* 46, 921-928 (2002).
 31. van der Helm-van Mil, A. H. et al. An independent role of protective HLA class II alleles in rheumatoid arthritis severity and susceptibility. *Arthritis Rheum.* 52, 2637-2644 (2005).
 32. Shadick, N. A. et al. Opposing effects of the D70 mutation and the shared epitope in HLA-DR4 on disease activity and certain disease phenotypes in rheumatoid arthritis. *Ann. Rheum. Dis.* 66, 1497-1502 (2007).
 33. Hensvold, H. A. et al. Environmental and genetic factors in the development of anticitrullinated protein antibodies (ACPAs) and ACPA-positive rheumatoid arthritis: an epidemiological investigation in twins. *Ann. Rheum. Dis.* 74, 375-380 (2013).
 34. Enouz, S., Carrie, L., Merkler, D., Bevan, M. J. & Zehn, D. Autoreactive T cells bypass negative selection and respond to self-antigen stimulation during infection. *J. Exp. Med.* 209, 1769-1779 (2012).
 35. Hand, T. W. et al. Acute gastrointestinal infection induces long-lived microbiota-specific T cell responses. *Science* 337, 1553-1556 (2012).
 36. Liu, X., Zou, Q., Zeng, B., Fang, Y. & Wei, H. Analysis of fecal lactobacillus community structure in patients with early rheumatoid arthritis. *Curr. Microbiol.* 67, 170-176 (2013).
 37. de Rooy, D. P., van der Linden, M. P., Knevel, R., Huizinga, T. W. & van der Helm-van Mil, A. H. Predicting arthritis outcomes—what can be learned from the Leiden Early Arthritis Clinic? *Rheumatology* 50, 93-100 (2011).

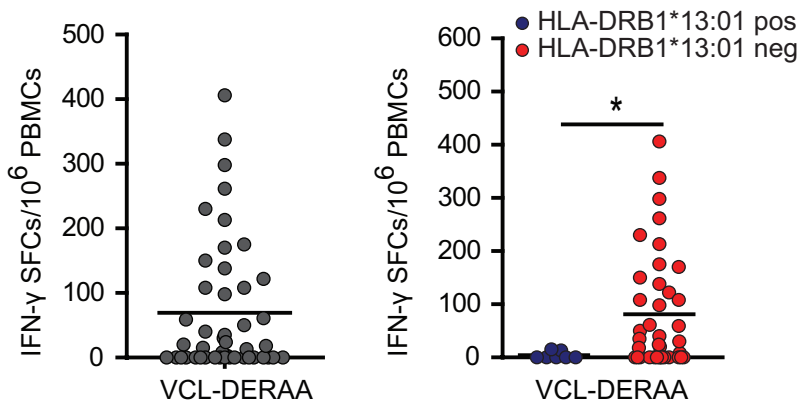
38. Erlich, H., Lee, J. S., Petersen, J. W., Bugawan, T. & DeMars, R. Molecular analysis of HLA class I and class II antigen loss mutants reveals a homozygous deletion of the DR, DR, and part of the DP region: implications for class II gene order. *Hum. Immunol.* 16, 205-219 (1986).
39. Stepniak, D. et al. Large-scale characterization of natural ligands explains the unique gluten-binding properties of HLA-DQ2. *J. Immunol.* 180, 3268-3278 (2008).
40. Shi, J. et al. Autoantibodies recognizing carbamylated proteins are present in sera of patients with rheumatoid arthritis and predict joint damage. *Proc. Natl Acad. Sci. USA* 108, 17372-17377 (2011).
41. Moustakas, A. K. et al. Structure of celiac disease-associated HLA-DQ8 and non-associated HLA-DQ9 alleles in complex with two disease-specific epitopes. *Int. Immunol.* 12, 1157-1166 (2000).
42. Lee, K. H., Wucherpfennig, K. W. & Wiley, D. C. Structure of a human insulin peptide-HLA-DQ8 complex and susceptibility to type 1 diabetes. *Nat. Immunol.* 2, 501-507 (2001).
43. Murthy, V. L. & Stern, L. J. The class II MHC protein HLA-DR1 in complex with an endogenous peptide: implications for the structural basis of the specificity of peptide binding. *Structure* 5, 1385-1396 (1997).
44. Fremont, D. H., Hendrickson, W. A., Marrack, P. & Kappler, J. Structures of an MHC class II molecule with covalently bound single peptides. *Science* 272, 1001-1004 (1996).
45. Rostkowski, M., Olsson, M. H., Sondergaard, C. R. & Jensen, J. H. Graphical analysis of pH-dependent properties of proteins predicted using PROPKA. *BMC Struct. Biol.* 11, 6 (2011).
46. Simonson, T. et al. Computational protein design: the Proteus software and selected applications. *J. Comput. Chem.* 34, 2472-2484 (2013).
47. Brooks, B. R. et al. CHARMM: the biomolecular simulation program. *J. Comput. Chem.* 30, 1545-1614 (2009).
48. Jo, S., Kim, T., Iyer, V. G. & Im, W. CHARMM-GUI: a web-based graphical user interface for CHARMM. *J. Comput. Chem.* 29, 1859-1865 (2008).
49. MacKerell, A. D. et al. All-atom empirical potential for molecular modeling and dynamics studies of proteins. *J. Phys. Chem. B* 102, 3586-3616 (1998).
50. Mackerell, Jr. A. D., Feig, M. & Brooks, III C. L. Extending the treatment of backbone energetics in protein force fields: limitations of gas-phase quantum mechanics in reproducing protein conformational distributions in molecular dynamics simulations. *J. Comput. Chem.* 25, 1400-1415 (2004).
51. Jorgensen, W. L., Chandrasekhar, J., Madura, J. D., Impey, R. W. & Klein, M. L. Comparison of simple potential functions for simulating liquid water. *J. Chem. Phys.* 79, 926-935 (1983).
52. Neria, E., Fischer, S. & Karplus, M. Simulation of activation free energies in molecular systems. *J. Chem. Phys.* 105, 1902-1921 (1996).
53. Darden, T., York, D. & Pedersen, L. Particle Mesh Ewald: an N.Log(N) method for Ewald sums in large systems. *J. Chem. Phys.* 98, 10089-10092 (1993).
54. Nose, S. A. Unified formulation of the constant temperature molecular-dynamics methods. *J. Chem. Phys.* 81, 511-519 (1984).
55. Hoover, W. G. Canonical dynamics - equilibrium phase-space distributions. *Phys. Rev. A* 31, 1695-1697 (1985).
56. Feller, S. E., Zhang, Y. H., Pastor, R. W. & Brooks, B. R. Constant-pressure molecular-dynamics simulation - the Langevin Piston method. *J. Chem. Phys.* 103, 4613-4621 (1995).
57. Ryckaert, J. P., Ciccotti, G. & Berendsen, H. J. C. Numerical-integration of cartesian equations of motion of a system with constraints - molecular-dynamics of N-alkanes. *J. Comput. Phys.* 23, 327-341 (1977).

58. Humphrey, W., Dalke, A. & Schulten, K. VMD: visual molecular dynamics. *J. Mol. Graph.* 14, 33-38 (1996).
59. Tamamis, P., Morikis, D., Floudas, C. A. & Archontis, G. Species specificity of the complement inhibitor compstatin investigated by all-atom molecular dynamics simulations. *Proteins* 78, 2655-2667 (2010).
60. Tamamis, P. et al. Insights into the mechanism of C5aR inhibition by PMX53 via implicit solvent molecular dynamics simulations and docking. *BMC Biophys.* 7, 5 (2014).
61. Im, W., Lee, M. S. & Brooks, III C. L. Generalized born model with a simple smoothing function. *J. Comput. Chem.* 24, 1691-1702 (2003).
62. Chen, J., Im, W. & Brooks, III C. L. Balancing solvation and intramolecular interactions: toward a consistent generalized Born force field. *J. Am. Chem. Soc.* 128, 3728-3736 (2006).

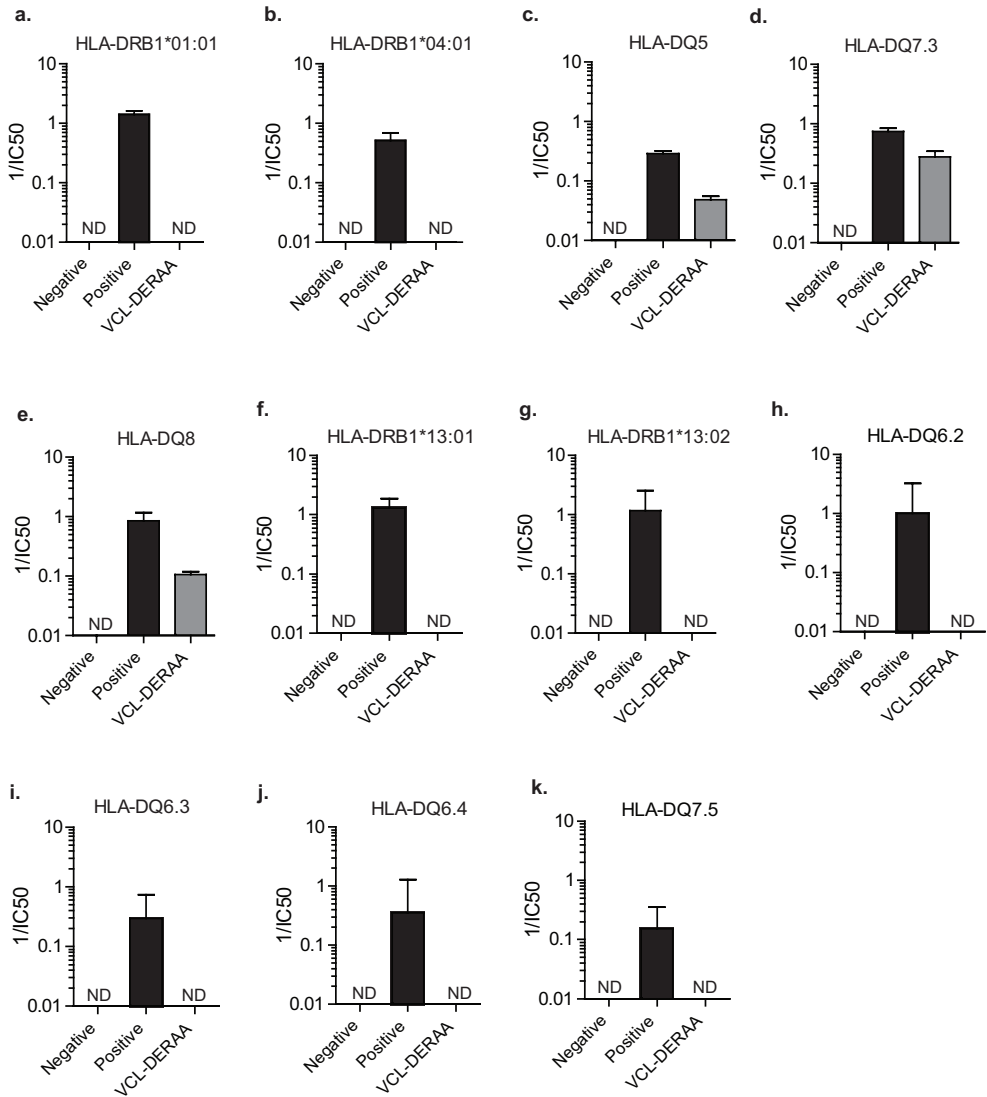
SUPPLEMENTARY FIGURES



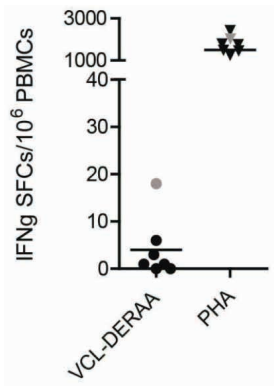
Supplementary Figure 1. Vinculin is an autoantigen recognized by circulating CD4+ T cells. IFN γ ELISPOT of PBMC from healthy individuals ($n=21$) stimulated for 24h with VCL-DERAA peptide REEVFDERAANFENH. Each dot represents a unique donor.



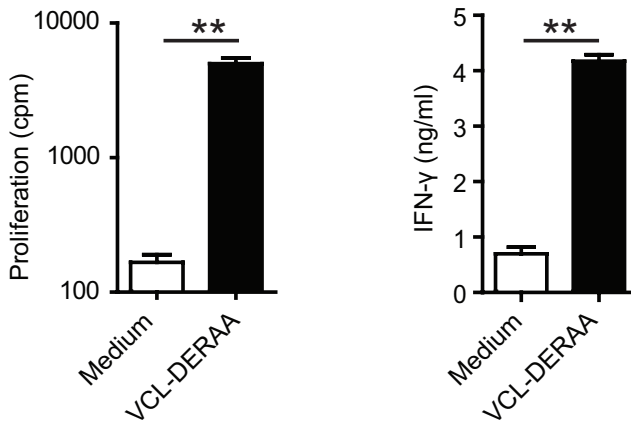
Supplementary Figure 2. Vinculin is an autoantigen recognized by circulating CD4+ T cells of HLA-DRB1*13-negative donors. IFN γ ELISPOT of PBMC from healthy individuals stimulated for 4d with VCL-DERAA peptide REEVFDERAANFENH (Left panel). Stratification of donors in two distinct groups based on the presence or absence of HLA-DRB1*13:01 (Right panel). Each dot represents a unique donor ($n=46$). Two-sided statistical analyses of ELISPOT data were performed using a Mann-Whitney U test with * indicating $p > 0.05$.



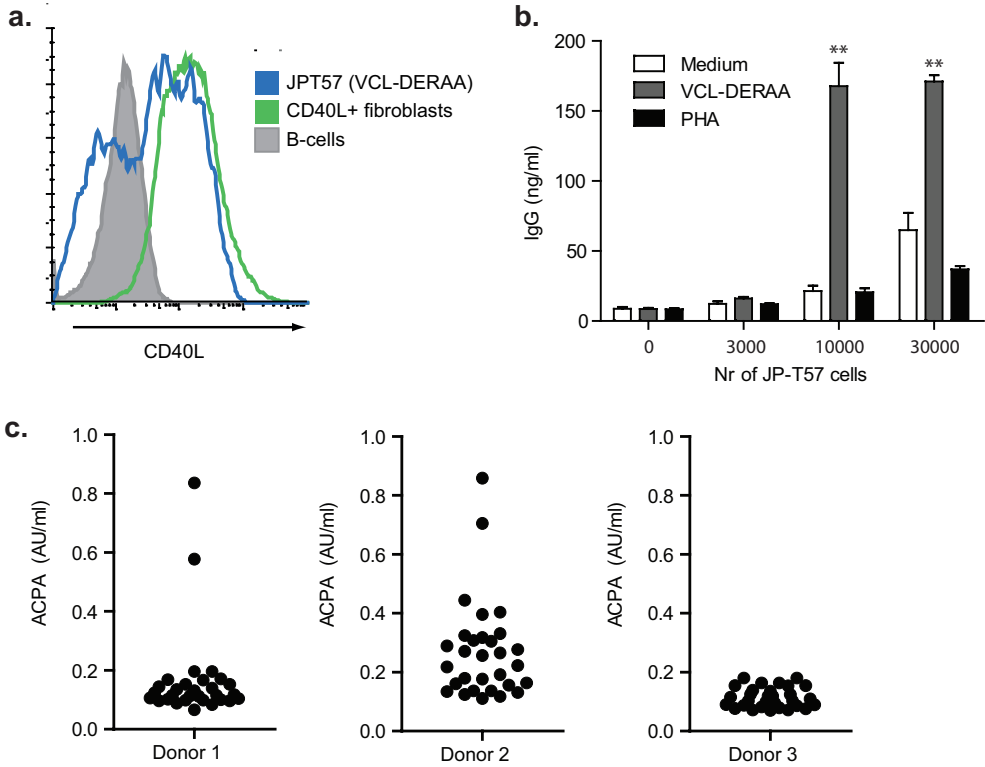
Supplementary Figure 3. Presentation to vinculin-DERAA-directed T cells is restricted to RA-predisposing HLA-DQ molecules. Competitive binding of a non-binding negative control peptide, an unbiotinylated positive control peptide and the VCL-DERAA peptide to HLA-DRB1*01:01 (A), HLA-DRB1*04:01 (B), HLA-DQ5 (C), HLA-DQ7.3 (D), HLA-DQ8 (E), HLA-DRB1*13:01 (F), HLA-DRB1*13:02 (G), HLA-DQ6.2 (H), HLA-DQ6.3 (I), HLA-DQ6.4 (J) or HLA-DQ7.5 (K). IC50 is the concentration of test-peptide (pM) where 50% of biotinylated peptide is bound, ND = non-detectable (IC50 > 300 pM). All experiments were performed at least three times and bars show pooled experiments, the error bars show the variation between the different experiments in S.E.M.



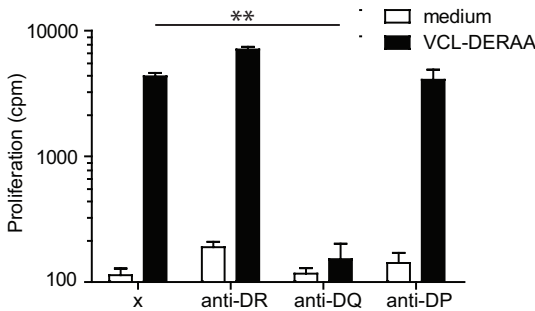
Supplementary Figure 4. Vinculin is not recognized by circulating CD4+ T cells of donors negative for predisposing HLA-DQ molecules. IFN γ ELISPOT of PBMC from healthy individuals negative for HLA-DQ5, HLA-DQ7.3 or HLA-DQ8 ($n=6$) stimulated for 24h with VCL-DERAA peptide REEVDERAANFENH or PHA. PBMC from an HLA-DQ8+ individual was used as a positive control (Gray dot).



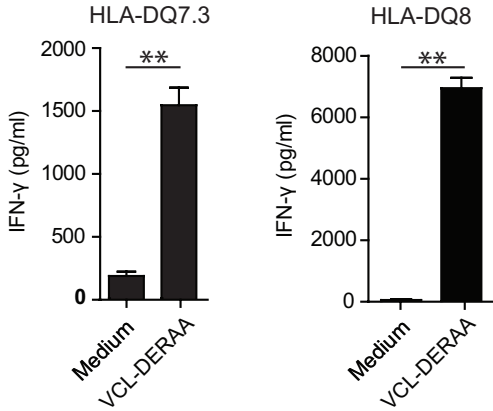
Supplementary Figure 5. T cell clone JPT57 specifically recognized the VCL-DERAA epitope. ³H-thymidine incorporation and IFN γ production upon stimulation of T cell clone JPT57 with unpulsed and VCL-DERAA pulsed autologous APCs. Data are representative of at least three independent experiments. Two-sided statistical analysis was performed using a student's t-test with ** indicating $p < 0.001$. Error bar is S.E.M.



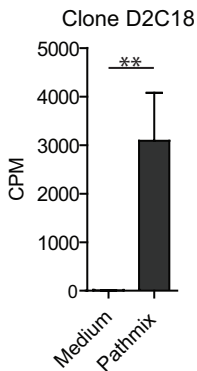
Supplementary Figure 6. Activated JPT57 cells can activate ACPA-producing B cells. (A) Flow cytometry analysis of CD40L expression on JPT57 cells stimulated with VCL-DERAA-pulsed APCs, CD40L-transfected fibroblasts and EBV-transformed B cells. (B) Co-culture of VCL-DERAA-pulsed HLA-DQ8-positive B cells from healthy subjects with increasing numbers of JPT57 cells in the presence of anti-IgM. The experiment was repeated three times and the plot shows a representative experiment. (C) Co-culture of VCL-DERAA-pulsed HLA-DQ8-positive B cells from three ACPA-positive RA patients with JPT57 (1:1) in the presence of anti-IgM. Activation of ACPA-producing B cells was determined by CCP2 ELISA. Each dot represents a different well. Statistical analysis was performed using a student's t-test with ** indicating $p > 0.001$. Error bars are S.E.M.



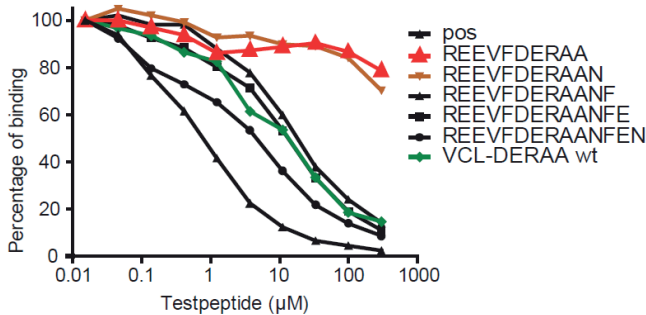
Supplementary Figure 7. Vinculin-DERAA-directed T cell clone JPT57 is HLA-DQ-restricted. ^3H -thymidine incorporation of JPT57 stimulated with feeders, pulsed or unpulsed with VCL-DERAA in the presence of HLA class II blocking antibodies. Data are representative of at least three independent experiments. Two-sided statistical analysis was performed using a student's t-test with ** indicating $p < 0.001$. Error bars are S.E.M.



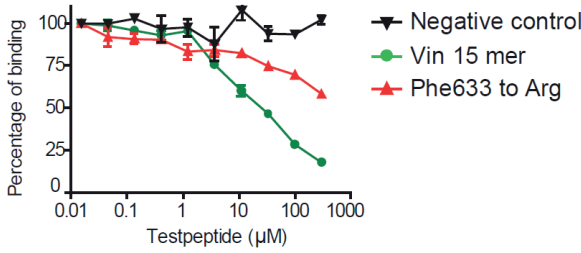
Supplementary Figure 8. Vinculin-DERAA-directed T cell clone JPT57 is restricted to HLA-DQ7.3 and HLA-DQ8. IFN γ production by JPT57 cells stimulated with HLA-DQ7.3 or HLA-DQ8 homozygous EBV-transformed B cells. Data are representative of at least three independent experiments. Two-sided statistical analysis was performed using a student's t-test with ** indicating $p < 0.001$. Error bars are S.E.M.



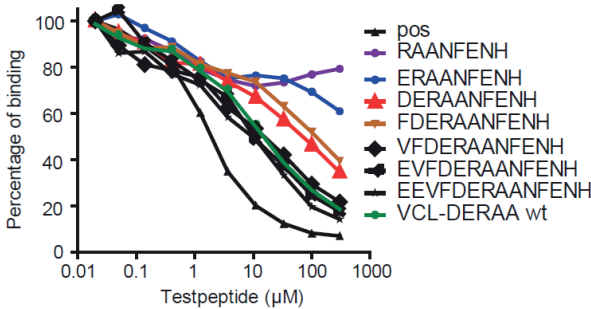
Supplementary Figure 9. Identification of a Pathmix directed T cell clone in an HLA-DRB1*13 negative donor. ³H-thymidine incorporation upon stimulation of T cell clone D2C18 with unpulsed and salmonella-DERAA-pulsed autologous monocytes. The experiment was repeated two times and the plot shows a representative experiment. Two-sided statistical analysis was performed using a student's t-test with ** indicating $p > 0.001$. Error bar is S.E.M.



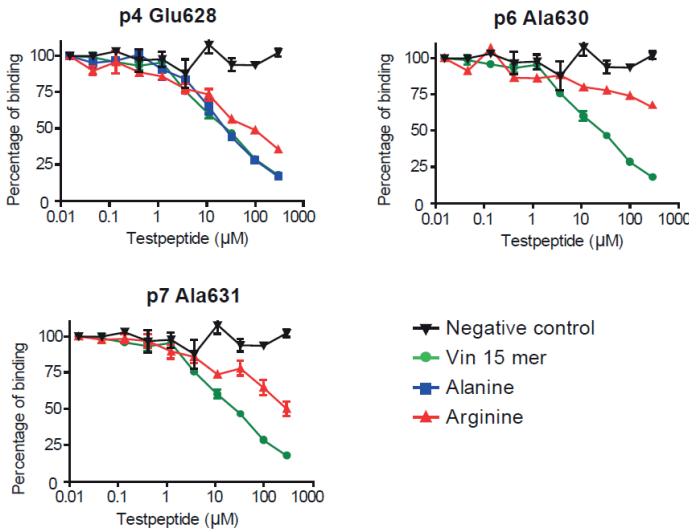
Supplementary Figure 10. Truncation of Phe633 results in loss of binding to HLA-DQ8. Competitive binding of HLA-DQ8 to an unbiotinylated positive control peptide and C-truncated VCL-DERAA peptides. The experiment was repeated three times and plots show a representative experiment.



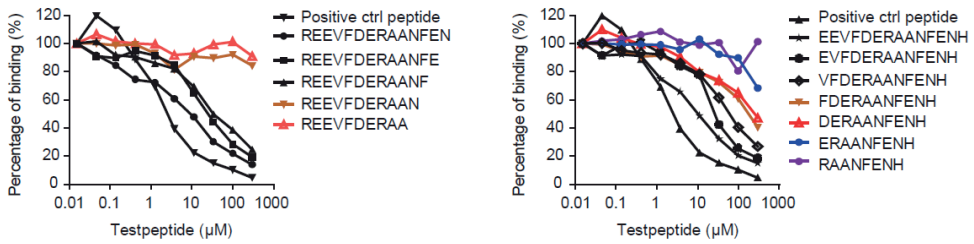
Supplementary Figure 11. Arginine substituting Phe633 negatively impacts binding affinity to HLA-DQ8. Competitive binding of a negative control peptide, the native VCL-DERAA peptide and the VCL-DERAA peptide with Phe633 substituted to arginine. Experiments were performed three times and plots show a representative experiment.



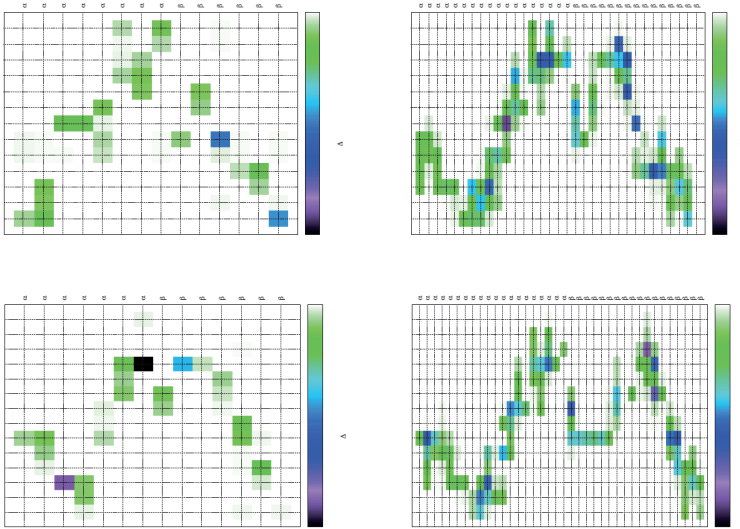
Supplementary Figure 12. N-terminal truncation of Val625 lowers the binding affinity to HLA-DQ8. Competitive binding of HLA-DQ8 to an unbiotinylated positive control peptide and N-truncated VCL-DERAA peptides. The experiment was repeated three times and plots show a representative experiment.



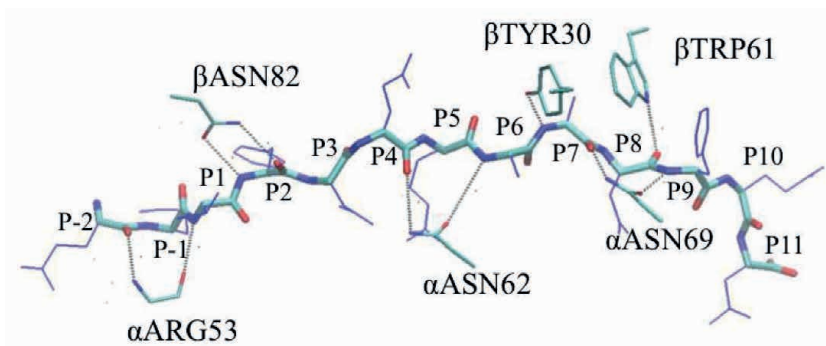
Supplementary Figure 13. Arginine-substitutions of residues implicated to interact with binding pockets negatively impacts binding to HLA-DQ8. Competitive binding of a negative control peptide, the native VCL-DERAA peptide and the VCL-DERAA peptide with Glu628, Ala630 and Ala631 substituted to alanine or arginine. Experiments were performed three times and plots show a representative experiment.



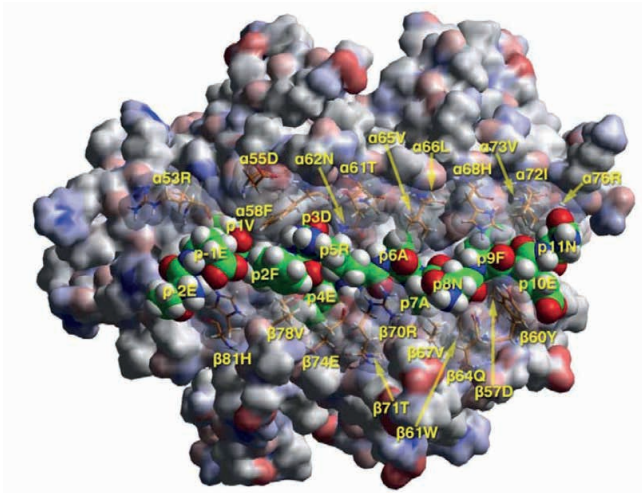
Supplementary Figure 14. HLA-DQ7.3 presents VCL-DERAA in a similar binding register as HLA-DQ8. Competitive binding of HLA-DQ7.3 to an unbiotinylated positive control peptide and C-truncated and N-truncated VCL-DERAA peptides. The experiment was repeated three times and plots show a representative experiment.



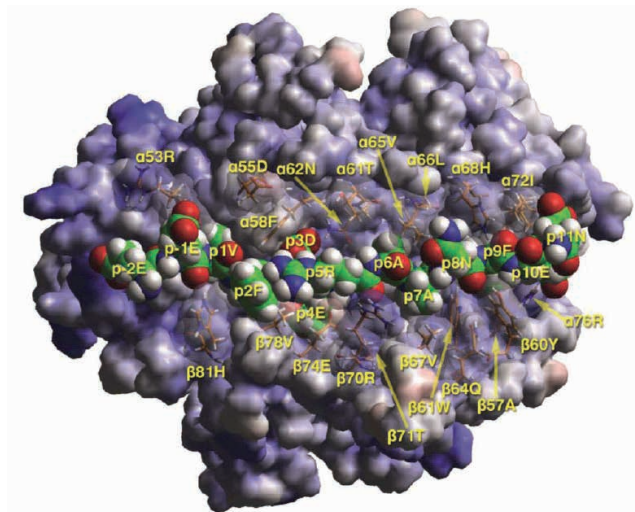
Supplementary Figure 15. Interaction free-energies (in kcal/mol) for selected peptide-ligand residue pairs, averaged over the MD trajectories. The left and right panels correspond, respectively, to polar (GB+Coulomb) and non-polar (vW + SA) interactions, computed with Eq. (1) of the methods. Rows from top to bottom correspond, respectively, to the DQ8:VCL-DERAA and DQ8:insulin complexes.



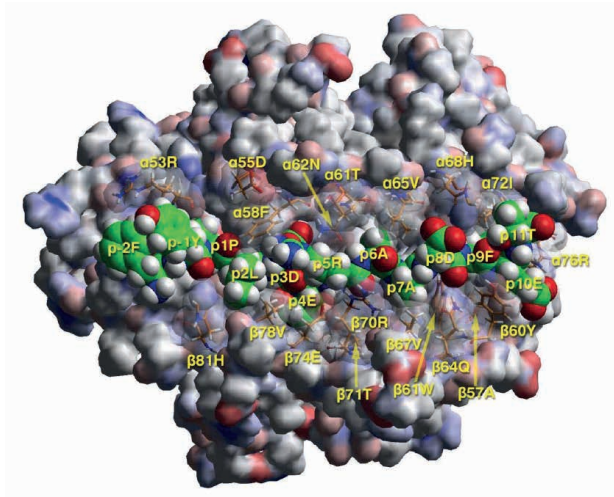
Supplementary Figure 16. Hydrogen bonds between the VCL-DERAA peptide mainchain and DQ8. The peptide mainchain is shown in thick licorice and its sidechains are displayed in blue lines. DQ8 groups are shown in thin licorice. Hydrogen bonds with occupancy smaller than 20% are omitted. A comprehensive list of all hydrogen bonds is included in Supplementary Table 1.



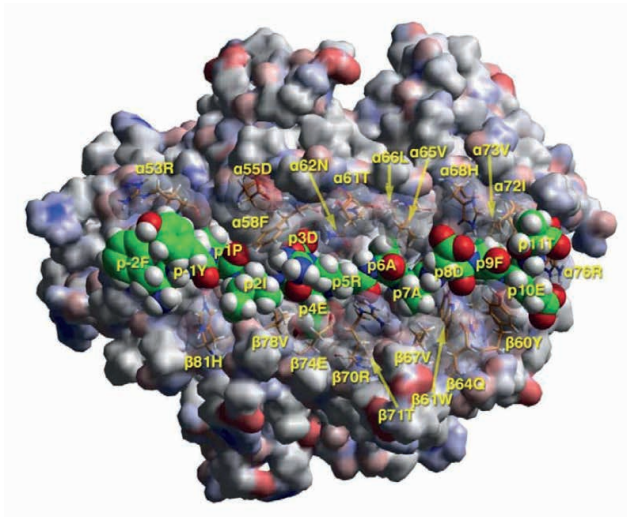
Supplementary Figure 17. TCR view of the complex of HLA-DQ7.3 with the VCL-DERRAA epitope in the groove at pH 7.4 (extracellular). The $\alpha 1\beta 1$ domain of the modelled HLA-DQ molecule is in van der Waals surface representation, colored according to atomic charges (negative = red, positive = blue, neutral = gray, partial charges shades in-between). Several visible residues from the HLA-DQ molecule in contact with the antigenic peptide and potential contact with a cognate TCR in canonical orientation are shown in stick form with a transparent surface (atomic color code: oxygen, red; nitrogen, blue; hydrogen, white; carbon, green; sulfur, yellow). The antigenic peptide in the groove is shown in space-filling form, with identical colour conventions as in Figure 6D-G. Anchors p1V and p9F point into the plane of the paper (screen) and are only partly seen. In vivo, p5R may not interact with p2F (cation- π interaction) because the former might more favorably interact with water molecules from the solvent. In this allele p4E makes an even weaker anchor than in DQ8, because of $\beta 13\text{Gly}\rightarrow\text{Ala}$ and $\beta 26\text{Leu}\rightarrow\text{Tyr}$ substitutions (DQ8 \rightarrow DQ7) that leaves less space available at the base of pocket 4.



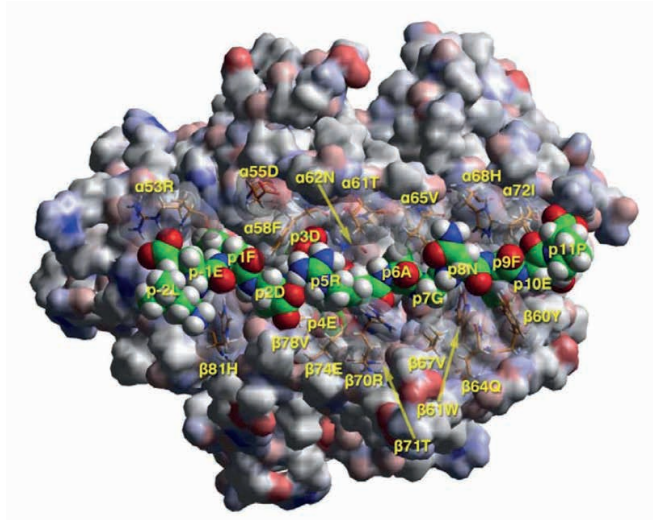
Supplementary Figure 18. TCR view of the complex of HLA-DQ8 with the VCL-DERRAA epitope in the groove at pH 7.4 (extracellular). Depiction and colour conventions as in Suppl. Figure 16. Anchors P1V and P9F point into the plane of the paper (screen) and are only partially seen. In vivo, P5R may not interact with P2F (cation- π interaction) because the former might more favorably interact with water molecules from the solvent.



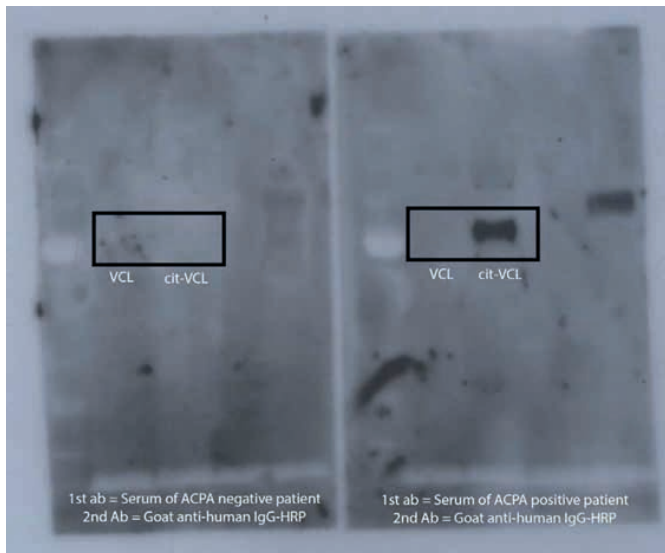
Supplementary Figure 19. TCR view of the complex of HLA-DQ8 with the *L. curvatus* epitope in the groove at pH 7.4 (extracellular). Depiction and colour conventions as in Suppl. Figure 16. Anchors P1P and P9F point into the plane of the paper (screen) and are only partially seen.



Supplementary Figure 20. TCR view of the complex of HLA-DQ8 with the *L. Sakei* epitope in the groove at pH 7.4 (extracellular). Depiction and colour conventions as in Suppl. Figure 16. Anchors P1P and P9F point into the plane of the paper (screen) and are only partially seen.



Supplementary Figure 21. TCR view of the complex of HLA-DQ8 with the *C. coli* epitope in the groove at pH 7.4 (extracellular). Depiction and colour conventions as in Suppl. Figure 16. Anchors P1P and P9F point into the plane of the paper (screen) and are only partially seen.



Supplementary Figure 22. Unedited blot of Figure 1A.

Donor	Acceptor	Occupancy (%)
Water	P-2E (sc)	381
P-2E (Nter)	Water	188
β Arg88 (sc)	P-2E(sc)	165
α Arg53 (mc)	P-2E (mc)	42.7
α Arg53 (sc)	P-2E (sc)	41.3
α Phe51 (mc)	P-2E (sc)	29.8*
α Phe51 (mc)	P-2E (Nter)	18.1*
Water	P-1E (sc)	535
Water	P-1E (mc)	62.9
P-1E (mc)	Water	56.7
β Asn82 (sc)	P-1E(mc)	20.5*
β His81-Side	P-1E(mc)	16.6*
Water	P1V (mc)	79.3
P1V (mc)	α Arg53 (mc)	76.9
P2F (mc)	β Asn82 (sc)	85.6
β Asn82 (sc)	P2F (mc)	68.5
Water	P3D (sc)	301
Water	P3D (mc)	64.2
α AsnN62 (sc)	P3D (sc)	46.9*
α Tyr22 (sc)	P3D (sc)	36.6*
Water	P4E (sc)	269
β Arg70 (sc)	P4E (sc)	160.4
β Thr71 (sc)	E6 (sc)	64.7*
β Thr28 (sc)	E6 (sc)	38.5*
α Asn62 (sc)	P4E(mc)	36

(Continued)

(Continued)

Donor	Acceptor	Occupancy (%)
P5R (sc)	Water	118
Water	P5R (mc)	98.3
P5R(sc)	α Thr61 (sc)	76.9
P5R (sc)	α Phe58 (mc)	74.1
Water	P6A (mc)	118
P6A (mc)	α Asn62 (sc)	88.5
P7A (mc)	β Tyr30 (sc)	65.4
P8N (mc)	Water	70.4
Water	P8N (sc)	70.4
Water	P8N (mc)	47.5
β Trp61 (sc)	P8N(mc)	41.9
P8N (sc)	Water	38.7
P8N (sc)	α Asn69 (sc)	35.0
α His68 (sc)	P8N (sc)	19.2
Water	P9F (mc)	65.6
P9F (mc)	α Asn69 (sc)	38.8
Water	P10E (sc)	55.5
Water	P10E (mc)	62.8
P10E (mc)	Water	48.6
Water	P11N (Cter)	461
Water	P11N (sc)	117.5
P11N (mc)	Water	54.7
P11N (sc)	Water	44.0
P11N(sc)	α His68 (sc)	21.3
α Arg76 (sc)	P11N(sc)	33.2

Supplementary Table 1. Statistics of intermolecular hydrogen bonds in the DQ8:VCL-DERAA complex. Hydrogen bonds were considered present if the D-A distance d_{DA} was smaller than 3.5 Å, and the angle $\theta_{D-H...A}$ was larger than 150°. The (*) denotes water-mediated interactions; “sc” and “mc” denote side-chain and main-chain groups.

	p-2	p-1	p1	p2	p3	p4	p5	p6	p7	p8	p9	p10	p11
VCL-DERAA	E	E	V	F	D	E	R	A	A	N	F	E	N
<i>Aspergillus fumigatus</i>	V	T	A	E	D	E	R	A	A	M	F	F	R
<i>Bacillus cereus</i>	V	A	V	P	D	E	R	A	A	N	A	I	A
<i>Campylobacter coli</i>	L	E	F	D	D	E	R	A	G	N	F	E	P
<i>Influenza A virus</i>	F	E	F	S	D	E	R	A	A	N	P	I	V
<i>Influenza A virus</i>	F	E	L	S	D	E	R	A	A	N	P	I	V
<i>Influenza A virus</i>	F	E	L	S	D	E	R	A	A	N	P	V	V
<i>Lactobacillus brevis</i>	L	V	T	D	D	E	R	A	A	I	F	K	A
<i>Lactobacillus curvatus</i>	F	Y	P	L	D	E	R	A	A	D	F	E	T
<i>Lactobacillus sakei</i>	F	Y	P	I	D	E	R	A	A	D	F	E	T
<i>Mycobacterium tuberculosis</i>	A	L	P	F	D	E	R	A	A	V	F	L	R
<i>Mycobacterium tuberculosis</i>	A	M	P	F	D	E	R	A	A	V	F	L	R
<i>Nocardia brasiliensis</i>	A	L	P	F	D	E	R	A	A	I	F	L	R
<i>Nocardia farcinica</i>	A	L	L	A	D	E	R	A	A	L	F	A	R
<i>Proteus mirabilis</i>	V	I	T	D	D	E	R	A	A	V	F	Y	G
<i>Proteus penneri</i>													
<i>Proteus stuartii</i>													
<i>Vibrio parahaemolyticus</i>	Y	D	E	L	D	E	R	A	A	W	F	Y	E
<i>Yersinia enterocolitica</i>	L	E	D	Y	D	E	R	A	A	N	G	Y	D

Supplementary Table 2. Microbe derived epitopes with molecular mimicry to vinculin-DERAA.

SUPPLEMENTARY NOTE 1**Functional Identification of the primary binding register of VCL-DERAA in HLA-DQ7.3 and DQ8.**

To establish the primary binding register of VCL-DERAA in HLA-DQ8, we performed HLA class II binding assay with N- and C-terminal truncated VCL-DERAA peptides. Upon C-terminal truncations, we observed a dramatic drop in binding affinity upon removal of Phe633 (Supplementary Figure 9). This observation made us speculate that this residue is interacting with the p9 pocket of HLA-DQ8. The p9 pocket of HLA-DQ8 is shaped by several different amino acids including α 68His and α 76Arg. Due to the presence of these positively charged amino acids, the pocket has a preference for negatively charged amino acids. Therefore, we substituted Phe633 for an arginine and observed a drop in binding capacity (Supplementary Figure 10).

Next, we performed N-terminal truncations. We performed a slight drop in binding affinity upon removal of Val625. The removal of Phe626 had no further effect, but removal of Asp627 resulted in a complete loss of binding affinity (Supplementary Figure 11). The N-terminal truncation of Val625 does not result in a complete loss of binding affinity. This is in agreement with previous data on binding of truncated peptides to HLA-DQ molecules and their mouse homologues H2-A, where p1 and p2 positions may be empty, provided the C-terminus of the peptide extends to p11 or beyond^{1, 2}. Indeed, upon truncation of Asp627, the residue proposed to interact with p3, we did observe a complete loss of binding affinity.

Finally, we performed alanine and arginine substitutions of Glu628, Ala630 and Ala631 that are proposed to interact with respectively the p4, p6 and p7 pocket further confirmed the proposed core register (Supplementary Figure 12).

Together this data supports **VFDERAANF** (anchors in bold) as the primary binding register of the VCL-DERAA epitope in HLA-DQ8. Next, we also examined the binding register of this epitope in HLA-DQ7.3 using N- and C-terminal truncated VCL-DERAA epitopes. We observed striking similarities with the previously performed binding assays to HLA-DQ8 (Supplementary Figure 13).

Together these data indicate that VCL-DERAA binds in a similar binding register to both HLA-DQ7.3 and HLA-DQ8.

Molecular Dynamics simulations of HLA-DQ:VCL-DERAA complexes

We studied the structure and interactions of the HLA-DQ8:VCL-DERAA complex in the binding register determined above via atomic-detail MD simulations. Figure 5 shows important hydrogen-bonding patterns and intermolecular nonpolar contacts in pockets P1, P4 and P9. Supplementary Figure 14 displays interaction free energies of selected protein-peptide residue pairs. These values are averaged over the MD trajectories and include solvent effects (see Methods).

The peptide P1V side chain forms nonpolar contacts with several surrounding residues (α Tyr9, α His24, α Phe32, α Val43, α Arg52, α Arg53, α Phe54, β Val78, β His81, β Asn82, β Leu85)

at the P1 pocket (Figure 5 and Supplementary Figure 14). Residues α Arg52, β Glu86, α Glu31, α Tyr9 and α His24 form the same dense network of interactions, as in the crystallographic DQ8:insulin complex³ and additional MD simulations of the DQ8:insulin complex; α His24 replaces a water-mediated interaction with the insulin P1E side chain by a water-mediated interaction with the β Glu86 side chain. In the second anchor pocket, the P4E side chain is surrounded by a large number of water molecules (on average, 6 waters are positioned with their oxygen atom within 4 Å of the side chain), and makes a strong interaction with α Arg70. Pocket P9 has a preference for negatively charged residues, which can form a salt bridge with α Arg76³⁻⁵. In the VCL-DERAA complex this pocket accepts a Phe residue. The P9F side chain is placed between β Trp61 and α Ile72, and makes several additional nonpolar contacts with residues α His68, α Asn69, α Val73, α Arg76, β Tyr37, β Ala57, and β Tyr60. The α Arg76 side chain interacts with the C-terminal residue P11Asn (Supplementary Figure 14 and Supplementary Table 1).

The protein interactions with VCL-DERAA anchor residues P1V, P9F are weaker than the interactions with insulin residues P1E and P9E, in the DQ8:insulin complex (Supplementary Figure 14). In the latter, residues P1E and P9E form salt bridges with α His24 and β Arg76, respectively; similarly, insulin residue P4Y forms stronger nonpolar contacts with the protein, relative to VCL-DERAA residue P4E (Supplementary Figure 14)³. These differences are in line with the weaker, relative to insulin, VCL-DERAA affinity for DQ8.

The interactions described so far involve the side chains of anchor residues. The peptide backbone makes several additional intermolecular hydrogen bonds (Supplementary Table 1 and Supplementary Figure 15) which are consistent with the conserved peptide-protein interactions of the MHCII binding motif⁶⁻⁸. Furthermore, the N- and C-terminal residues form salt bridges with the protein (P-2E- β Arg88, P-2E- β Arg53, P11N- α Arg76).

As discussed above, our IFN- γ ELISA measurements suggest that residues P2F, P8N, P9F and P10E are important TCR contacts, as they affect T cell recognition. The P2F side chain packs against the α -helix of the β 1 domain of HLA-DQ8 and makes direct contacts with several of its residues (Figure 5G and Supplementary Figure 14). Its opposite side is exposed to solvent, and is positioned near the side chain ring of α Phe58. During the simulation, the two rings remain approximately parallel to each other, at an average mutual distance of \sim 8 Å. The P2F and α Phe58 rings and the nonpolar moiety of P5R form a hydrophobic cluster (Figure 5G), which could play a role in TCR recognition. The side chains of the important TCR contact residues P8N and P10E side chains are also solvent-exposed and make few interactions with the protein. Among the other residues of the DERAA sequence, the P3D and P5R side chains face also toward the TCR binding region and form a stable salt-bridge interaction.

SUPPLEMENTARY REFERENCES

1. Scott, C.A., Peterson, P.A., Teyton, L., & Wilson, I.A. Crystal structures of two I-Ad-peptide complexes reveal that high affinity can be achieved without large anchor residues. *Immunity*. 8, 319-329 (1998).
2. He, X.L. et al. Structural snapshot of aberrant antigen presentation linked to autoimmunity: the immunodominant epitope of MBP complexed with I-Au. *Immunity*. 17, 83-94 (2002).
3. Lee, K.H., Wucherpfennig, K.W., & Wiley, D.C. Structure of a human insulin peptide-HLA-DQ8 complex and susceptibility to type 1 diabetes. *Nat. Immunol.* 2, 501-507 (2001).
4. Kwok, W.W., Domeier, M.L., Raymond, F.C., Byers, P., & Nepom, G.T. Allele-specific motifs characterize HLA-DQ interactions with a diabetes-associated peptide derived from glutamic acid decarboxylase. *J. Immunol.* 156, 2171-2177 (1996).
5. Godkin, A. et al. Use of eluted peptide sequence data to identify the binding characteristics of peptides to the insulin-dependent diabetes susceptibility allele HLA-DQ8 (DQ 3.2). *Int. Immunol.* 9, 905-911 (1997).
6. Stern, L.J. et al. Crystal structure of the human class II MHC protein HLA-DR1 complexed with an influenza virus peptide. *Nature* 368, 215-221 (1994).
7. Jardetzky, T.S. et al. Crystallographic analysis of endogenous peptides associated with HLA-DR1 suggests a common, polyproline II-like conformation for bound peptides. *Proc. Natl. Acad. Sci. U. S. A* 93, 734-738 (1996).
8. Painter, C.A. & Stern, L.J. Conformational variation in structures of classical and non- classical MHCII proteins and functional implications. *Immunol. Rev.* 250, 144-157 (2012).

PART II

THE WORKING MECHANISM OF THE BIOLOGICAL ABATACEPT

