

Recognition of nonpeptide antigens by human V γ 9V δ 2 T cells requires contact with cells of human origin

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SUMMARY

It is becoming apparent that $\gamma\delta$ T cells form an important part of the adaptive immune response. However, the ligands recognized by $\gamma\delta$ T cell receptors (TCRs) and the exact biological function of the cells that express this receptor remain unclear. Numerous studies have shown that the dominant human peripheral blood subset of $\gamma\delta$ T cells, which express a V γ 9V δ 2 TCR, can activate in response to low molecular weight nonpeptidic molecules. Some of these components have been purified from bacteria or parasites. We examined the activation of polyclonal $\gamma\delta$ T cell lines, clones with V γ 9V δ 2 and V γ 9V δ 1 TCRs, and $\gamma\delta$ T cells directly *ex vivo* in response to multiple phosphate, alkylamine and aminobisphosphonate (nBP) antigens and purified protein derivative from *Mycobacterium tuberculosis* (PPD). V γ 9V δ 2 T cells were able to respond to multiple small organic molecules of highly variable structure whereas cells expressing a similar V γ 9 chain paired with a V δ 1 chain failed to recognize these antigens. Thus, the TCR δ chain appears to make an important contribution to the recognition of these antigens. The kinetics of responses to alkylphosphate and alkylamine antigens differ from those of responses to the nBP pamidronate. These different classes of antigen are believed to have differed mechanisms of action. Such differences explain why nBPs can be pulsed onto antigen presenting cells (APCs) and still retain their ability to activate $\gamma\delta$ T cells while alkylphosphate and alkylamine antigens cannot. We also demonstrate that a substantial proportion of the cells that produce IFN γ directly *ex vivo* in response to PPD are $\gamma\delta$ T cells and that $\gamma\delta$ T cell activation requires contact with cells of human origin.

Keywords T lymphocytes $\gamma\delta$ T cell receptors alkylphosphate alkylamine aminobisphosphonate

INTRODUCTION

The key to adaptive immunity lies in the somatic gene rearrangement of antigen receptors that produces an almost infinite array of specific configurations from a finite number of genes. There are three different types of antigen receptor: the B cell receptor (antibody), the $\alpha\beta$ T cell receptor (TCR), which generally recognizes small peptide antigens in the context of self major histocompatibility complex (MHC) molecules presented on the cell surface, and the $\gamma\delta$ TCR. These three receptors distinguish the main lymphocyte lineages: B cells, $\alpha\beta$ T cells and $\gamma\delta$ T cells. The last 40 years have seen much progress in our understanding of how B cells and $\alpha\beta$ T cells contribute to immunity. Surprisingly, the ligands recog-

nized by the $\gamma\delta$ TCR and the exact biological role of cells that express this receptor remain unclear [1,2].

The potential importance of $\gamma\delta$ T cells is evident from the large number of possible $\gamma\delta$ T cell receptors (TCRs) [3]. Despite a relatively restricted number of V genes compared with $\alpha\beta$ T cells, huge diversity can be generated through the D and J gene segments and it has been estimated that approximately 10^{20} different $\gamma\delta$ TCRs can be created by these gene rearrangements [3]. This is substantially more than the number of possible antibodies or $\alpha\beta$ TCRs that the human body can generate although it is not clear to what extent $\gamma\delta$ T cells exploit this potential. Hints at an important role for $\gamma\delta$ T cells in immunity come from the observed large expansions of these cells that occur in response to a wide range of pathogens. In humans, bacterial infections including *Mycobacterium tuberculosis* [4], *Mycobacterium leprae* [5], *Salmonella typhi* [6], *Brucella melitensis* [7], *Francisella tularensis* [8], *Listeria monocytogenes* [9,10] and *Ehrlichia chaffeensis* [11], viral infections including HIV [12–14], Epstein-Barr Virus [15] and cytome-

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galovirus [16,17], yeast infection with *Candida albicans* [18] and parasitic infections including *Plasmodium falciparum* [19], *Plasmodium vivax* [20] and *Leishmania donovani* [21] can all stimulate a dramatic expansion of $\gamma\delta$ T cells *in vivo*. While murine studies have shown that $\gamma\delta$ T cells are redundant and nonessential for immunity to many infectious agents (reviewed in [1]) they are indispensable and nonredundant for immunity to other pathogens. Depletion of the $\gamma\delta$ T cell subset in mice with TCR-specific antibodies can result in an increase in bacterial titres by 2–3 orders of magnitude after infection with *L. monocytogenes* [22,23]. A recent study showed that while intranasal inoculation of $\gamma\delta^{-/-}$ mice with *Nocardia asteroides* led to 100% fatality within two weeks there were no deaths in $\gamma\delta^{+/+}$ control mice [24]. Other studies in $\gamma\delta^{-/-}$ mice reveal a role for $\gamma\delta$ T cells in protection against *Mycobacterium bovis* bacillus Calmette-Guerin (BCG) [25], *M. tuberculosis* [26] and *L. monocytogenes* [27]. In addition, recent experiments in primates highlight the adaptive nature of $\gamma\delta$ T cell responses. Macaques infected with BCG showed up to 25 fold increases in circulating V γ 9V δ 2 T cells and exhibited larger expansions during re-infection [28], demonstrating that $\gamma\delta$ T cells, in common with their $\alpha\beta$ TCR-expressing cousins, possess the capacity for immunological memory. The clonotypic expansions of V γ 9V δ 2 T cells during BCG infection and re-infection infer antigen-specificity in primary and memory $\gamma\delta$ T cell-mediated immune responses [28]. Vaccination of humans with BCG induces a marked increase in the capacity of $\gamma\delta$ T cells to expand *in vitro* following stimulation with live BCG or *M. tuberculosis* lysate [29]. These data suggest that human $\gamma\delta$ T cells can mount a memory response [29]. The immune potency of human peripheral blood $\gamma\delta$ T cells *in vivo* has also been demonstrated in the SCID mouse model. Inoculation of SCID mice with human PBMC rescues mice from lethal infection with *E. coli* [30]. Depletion of V δ 2-expressing T cells from PBMC resulted in a higher bacterial load and lower survival rate [30].

Despite the large number of organisms known to activate $\gamma\delta$ T cells, relatively few antigens have been identified. Recent advances have described low molecular weight, nonpeptidic compounds that appear to be able to activate the dominant subset of $\gamma\delta$ T cells in human peripheral blood, those with a V γ 9V δ 2 TCR. These compounds include alkylphosphates, alkylamines and aminobisphosphonates (nBPs) [31–33]. Recognition of all these groups of molecules appears to be subject to strict chemical constraints as minor modifications in antigen structure abolish activity [32,34,35]. Many of the $\gamma\delta$ T cell phosphoantigens and the antigenic alkylamines are produced by bacteria, suggesting that the recognition of such compounds might be immunologically relevant.

In this study, we examine the recognition requirements of human $\gamma\delta$ T cells in detail. We show that V γ 9 T cells expressing a V δ 2 chain recognize multiple small nonpeptidic ligands. In contrast, cells isolated from human peripheral blood that express a V γ 9 chain paired with a V δ 1 chain fail to recognize any of these ligands. A single V γ 9V δ 2 T cell clone activated in response to alkylphosphate, alkylamine, and nBP antigens despite the obvious differences in their chemistry. Recognition of all these compounds was dependent on cell-cell contact. nBP ligands could be pulsed onto antigen presenting cells (APCs), while alkylphosphate and alkylamine antigens needed to be continuously present in the media to activate $\gamma\delta$ T cells. Furthermore, all these molecules were only able to activate $\gamma\delta$ T cells when the presenting cell was of human origin.

MATERIALS AND METHODS

$\gamma\delta$ T cell culture

$\gamma\delta$ T cells were isolated from human peripheral blood using a magnetic bead-based separation kit (TCR γ/δ Microbead Kit and MS columns, Miltenyi Biotech Ltd., Bisley, UK). Once isolated, cells were resuspended in T cell medium (RPMI supplemented with penicillin, streptomycin and glutamine, 10% FCS, 10% T-STIM (BD Biosciences, Cowley, UK), 200 U/ml Proleukin (Chiron)) containing 2×10^6 γ -irradiated human peripheral blood mononuclear cells (PBMC)/ml from three unrelated donors and 2 μ g/ml phytohaemagglutinin (PHA). Cells were maintained with T cell medium, and restimulated with mixed irradiated PBMC and PHA every three weeks. For cloning, the same mix was used to grow cells by limiting dilution in 96-well U-bottomed plates. The IMGT system of TCR nomenclature is used throughout this work [36]. Cells that grew were confirmed to be $\gamma\delta$ T cells with V γ 9 antibody. 59/60 of V γ 9-expressing clones also stained with V δ 2 antibody. The remaining clone expressed a V δ 1 receptor. Two of the best growing V γ 9V δ 2 clones and the V γ 9V δ 1 clone were taken forward for further examination. Sequencing showed that both V γ 9V δ 2 T cells expressed an identical TCR.

Sequencing $\gamma\delta$ TCRs

TCR usage in $\gamma\delta$ T cell clones was characterized with a molecular analysis of gene expression. RNA was isolated from 10^6 cells in TriReagent (Sigma-Aldrich Company Ltd., Gillingham, UK). cDNA was prepared using a poly T oligonucleotide and reverse transcriptase enzyme. Polymerase chain reaction (PCR) was carried out for the gamma chain and for the delta chain using a full panel of variable region primers and a reverse primer to anneal to the constant region. The results of the PCR were consistent with the flow cytometric analysis and demonstrated both a V γ 9V δ 1 and a V γ 9V δ 2 clone.

Antibodies

The following antibodies were used for flow cytometric analysis: FITC-conjugated mouse anti-human V δ 2 mAb clone B6-1 (BD Biosciences Pharmingen, Cowley, UK), FITC-conjugated mouse anti-human V γ 9 mAb clone 7A5 (Endogen), FITC-conjugated mouse anti-human V δ 1 mAb clone TS8-2 (Endogen), PE-conjugated mouse anti-human V γ 9 mAb clone B3-1 (BD Biosciences Pharmingen, Cowley, UK), FITC-conjugated mouse anti-human pan- $\gamma\delta$ mAb clone 5A6.E9 (Endogen), PerCP-conjugated mouse anti-human CD3 mAb clone SK7 (BD Biosciences), FITC-conjugated mouse anti-human CD69 mAb clone L78 (BD Biosciences), allophycocyanin (APC)-conjugated mouse anti-human IL2 clone MQ1-17H12 (Caltag-MedSystems Ltd., Siverstone, UK), APC-conjugated mouse anti-human TNF α clone Mab11 (BD Biosciences Pharmingen, Cowley, UK) and APC-conjugated mouse anti-human IFN γ clone B27 (BD Biosciences Pharmingen, Cowley, UK).

$\gamma\delta$ T cell antigens

Alkylamines (Sigma) were diluted to a 1 M stock in RPMI media and brought to pH 7.0 with concentrated HCl. Alkylamine stocks were maintained in small aliquots at -80°C . 1 mM isopentenyl pyrophosphate (Sigma), 1 mM pamidronate (Calbiochem) and 100 μ M risedronate (Procter & Gamble) stocks in RPMI were stored in the same way. Tuberculin purified protein derivative (PPD) was purchased from the Statens Seruminstitut

(Denmark). A cross-linking anti-CD3 monoclonal antibody (clone UCHT-1; Amersham) was used as a positive control.

IFN γ ELISpot

$\gamma\delta$ T cells were washed in RPMI and incubated overnight in R10 at 37°C. 96-well nitrocellulose plates (Millipore (UK) Ltd., Watford, UK) were incubated overnight at 4°C with 15 μ g/ml anti-human-IFN γ primary antibody (clone 1-D1K; Mabtech, AB, Nacka Strand, Sweden). The plates were then washed twice with RPMI and blocked with R10 for 3 h at 37°C. R10 was decanted by inversion and assays applied to each well before incubation at 37°C as detailed below. Assays were terminated by washing once in water, followed by 5 washes in PBS. Secondary antibody (anti-human-IFN γ -Biotin antibody clone 7-B6-1; Mabtech) was applied at 1 μ g/ml and the plate incubated for 100 min at room temperature (RT). The plate was washed 6 times with PBS before application of streptavidin-ALP (1 : 1000 in PBS; Mabtech) for 40 min at RT. After 5 further washes in PBS, spots were revealed by incubation for 15 min at RT with developing buffer (Bio-Rad AP conjugate substrate kit) and counted mechanically using an ELISpot Reader System ELR02 (Autoimmun Diagnostika; Strassberg) after drying. The data displayed has not undergone any manual manipulation. A variety of cell lines were used as antigen presenting cells in IFN γ ELISpot assays. These included the MHC class I-deficient cell lines 721.221 and C1R, the TAP-deficient cell line 721.174, Jurkat T cells, HeLa cells, the β 2m-deficient cell line Daudi and several EBV-transformed B cell lines and primary tumour lines.

Intracellular cytokine staining

10⁶ fresh PBMC were incubated in FACS tubes with brefeldin A (10 μ g/ml in R10) for 5 h after incubating with relevant antigens for 1 h. The cells were then washed, permeabilized in 20% FACSLyse (BD Biosciences), washed twice in ice cold PBS/0.1% BSA and stained on ice with pretitrated APC-conjugated monoclonal antibodies specific for IFN γ , TNF α and IL-2, together with PE-conjugated anti-V γ 9, for 20 min. Cells were then washed, resuspended in PBS and analysed immediately on a FACSCalibur flow cytometer with CellQuest software (BD Biosciences, Cowley, UK).

MIP1 β ELISA

5 \times 10⁴ clonal V γ 9V δ 1 or V γ 9V δ 2 T cells were incubated with 5 \times 10⁴ Spinner HeLa cells for 4 h at 37°C in a 96-well U-bottom plate, in the presence or absence of 1.5 μ g/ml anti-CD3 antibody (clone UCHT-1; Amersham plc., Amersham, UK), 10 mM secbutylamine, 10 μ M IPP, 1 μ M risedronate (Procter & Gamble). MIP-1 β concentrations in the cell culture supernatant were determined with a Quantikine MIP1 β ELISA kit (R & D Systems, Europe Ltd., Abingdon, UK) as previously described [37].

$\gamma\delta$ T cell depletion of PBMC

10⁸ human PBMC were incubated with 10 μ g/ml anti-human pan- $\gamma\delta$ antibody for 30 min on ice. Cells were washed once in 20X labelling volume with PBS/0.1% BSA and resuspended at 4 \times 10⁷ cells/ml in PBS/0.1% BSA. 2 \times 10⁷ anti-mouse IgG1 Dynabeads (DynaL Biotech Ltd., Bromborough, UK) washed in PBS/0.1% BSA were added and the cells incubated for 1 h at 4°C with gentle rotation. The tube was then placed in the magnetic particle concentrator (DynaL) and left to separate for 30 min. The supernatant was transferred to a fresh tube. The success of depletion

was determined by FACS analysis with the anti-CD3 and pan- $\gamma\delta$ antibodies detailed above.

Stimulation of $\gamma\delta$ T lymphocytes for subsequent immunoblotting

Cells were washed twice in RPMI and incubated overnight in R10. The following day, FCS was washed off with two changes of RPMI and 10⁶ $\gamma\delta$ T cells were resuspended in 10 μ l of RPMI. After 10 min at 37°C in 5% CO₂, T cells were stimulated by incubation with 10 μ g/ml anti-CD3 antibody (clone UCHT-1; Amersham) for 3 min. The reaction was stopped by washing once with 1 ml ice-cold PBS, and resuspending the pellet in cold lysis buffer (140 mM NaCl, 20 mM Tris pH 8.0, 10 mM NaF, 2 mM EDTA, 20% glycerol, 1% IGEPAL, 1 mM Na₃VO₄, 10 μ g/ml aprotinin, 10 μ g/ml leupeptin) at 5 \times 10⁷ cells/ml. Blots were performed as described previously [38].

RESULTS

Activation of human peripheral blood $\gamma\delta$ T cells directly ex vivo

Recent studies have described low molecular weight, nonpeptidic compounds that appear to be able to activate the V γ 9V δ 2 dominant subset of $\gamma\delta$ T cells in human peripheral blood. These compounds include alkylphosphates [31,39], alkylamines [32] and nBPs [33]. We examined IFN γ production in response to activating alkylphosphate and alkylamine ligands and compared this to PPD by ELISpot (Fig. 1). Significant spot formation was observed with the previously reported [32] $\gamma\delta$ antigens secbutylamine and isobutylamine. Between 15 and 220 spots were observed per 5 \times 10⁴ peripheral blood mononuclear cells (PBMC) from nine separate healthy donors (Fig. 2a). Curiously, ethylamine, isopropylamine, ethanolamine, and isoamylamine did not appear to activate (Fig. 2b) despite having been previously described as antigenic at the concentration used [32]. However, alkylamines only activate $\gamma\delta$ T cells at concentrations close to those that are cytotoxic [32]. It is possible that the assays or cells we used were less sensitive than in previous studies so that cells died at alkylamine concentrations below the activation threshold. Magnetic depletion of $\gamma\delta$ T cells from PBMC (judged to be >90% successful by FACS analysis) removed the vast majority of alkylamine-induced spots (Fig. 1a,b). Magnetic depletion similarly removed the majority of spots induced by the phosphate antigen isopentenyl pyrophosphate (IPP) (Fig. 1a,b). Titration showed that IPP was a significantly more potent antigen than the alkylamines (Fig. 2b). High concentrations of isobutylamine were toxic [32] (Figs 1a and 2b) and also prevented peptide-induced spot formation by $\alpha\beta$ TCR-expressing cytotoxic T lymphocytes (data not shown). Magnetic-depletion of $\gamma\delta$ T cells was observed to remove approximately a third of IFN γ spot forming cells in response to PPD during overnight ELISpot assay (Fig. 1). Similar results were obtained using PBMC from two other individuals (data not shown). V γ 9 T cells were shown to respond to IPP, risedronate and PPD in direct *ex vivo* intracellular cytokine staining (ICS) assays (Fig. 2c). Examination of T cell activation by CD69 up-regulation (Fig. 2d) confirmed the presence of PPD-reactive $\gamma\delta$ T cells in the PBMC of more than half of the individuals studied. The responses measured by 6 h ICS were significantly lower than those measured by overnight ELISpot or CD69 up-regulation. Far fewer peripheral blood $\gamma\delta$ T cells up-regulated surface expression of CD69 (<5%) after only 6 h incubation with PPD (data not shown); thus, the difference between ICS and other assays likely reflects the duration of antigen exposure. Overnight ICS assay was toxic to cells.

Characterization of $\gamma\delta$ T cell clones

In order to characterize peripheral blood $\gamma\delta$ T cells further, we grew $\gamma\delta$ T cell clones from human PBMC and screened for those bearing the dominant V γ 9 receptor by flow cytometry with monoclonal antibody clone 7A5 (Endogen). Curiously, 1/60 of the clones that grew expressed a V γ 9 receptor paired with a V δ 1 chain. Two robust V γ 9V δ 2-expressing and the V γ 9V δ 1-expressing clone were taken forward for further analysis. The TCR sequences

of these clones (Fig. 3) confirmed these designations but showed that both V γ 9V δ 2-expressing cells expressed an identical TCR. The CDR3 regions of the V γ 9 chain differed in the V δ 1 and V δ 2 clone (Fig. 3). Both V γ 9 chains used a joining segment encoded by the TCRGJP (J γ P) gene (IMGT nomenclature [36]). The V γ 9V δ 1 and V γ 9V δ 2 clones expressed high levels of cell surface TCR (Fig. 4a) and could be activated through this TCR by CD3-crosslinking (Fig. 4b). The V γ 9V δ 2 clone made substantial

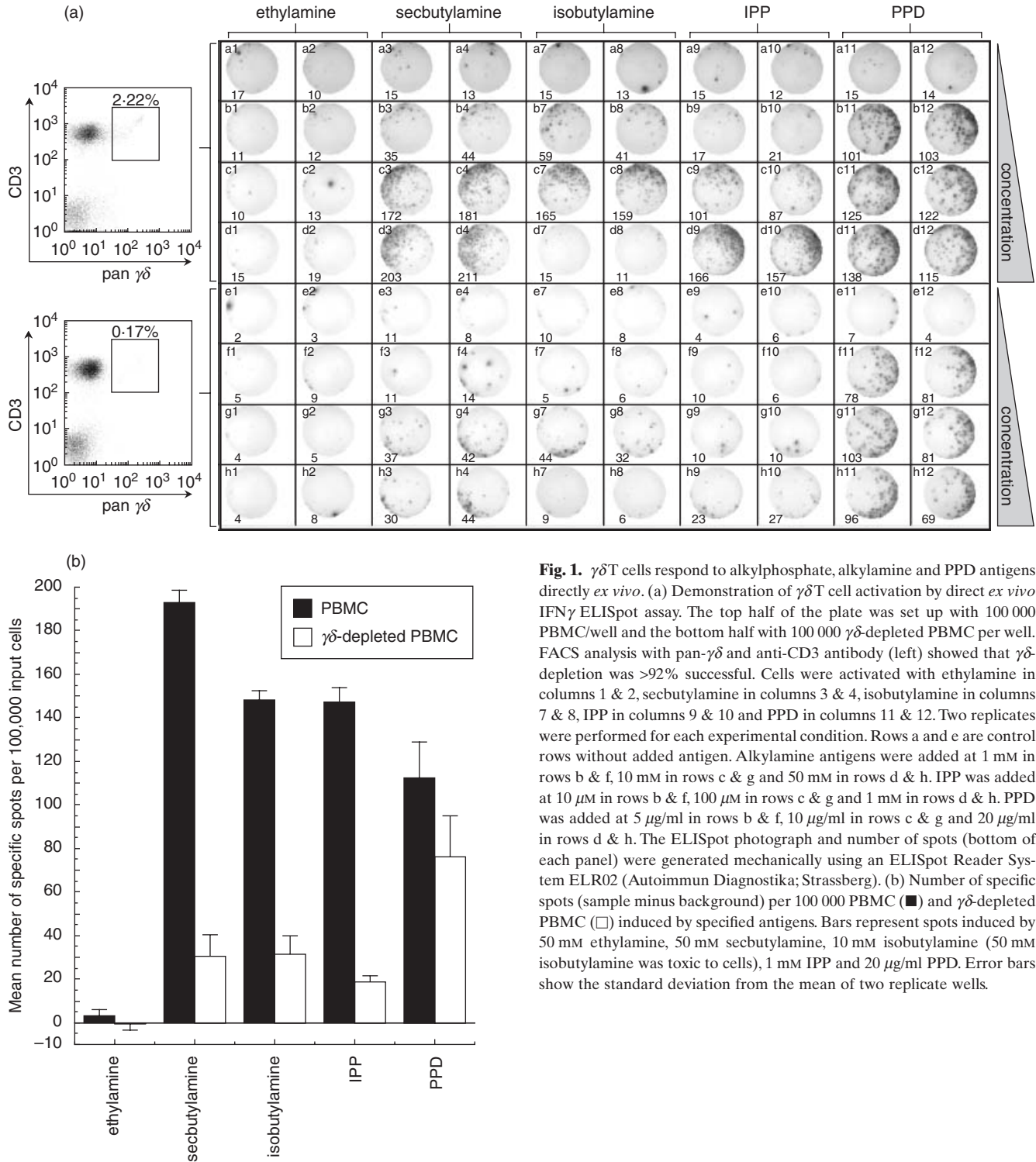


Fig. 1. $\gamma\delta$ T cells respond to alkylphosphate, alkylamine and PPD antigens directly *ex vivo*. (a) Demonstration of $\gamma\delta$ T cell activation by direct *ex vivo* IFN γ ELISpot assay. The top half of the plate was set up with 100 000 PBMC/well and the bottom half with 100 000 $\gamma\delta$ -depleted PBMC per well. FACS analysis with pan- $\gamma\delta$ and anti-CD3 antibody (left) showed that $\gamma\delta$ -depletion was >92% successful. Cells were activated with ethylamine in columns 1 & 2, secbutylamine in columns 3 & 4, isobutylamine in columns 7 & 8, IPP in columns 9 & 10 and PPD in columns 11 & 12. Two replicates were performed for each experimental condition. Rows a and e are control rows without added antigen. Alkylamine antigens were added at 1 mM in rows b & f, 10 mM in rows c & g and 50 mM in rows d & h. IPP was added at 10 μ M in rows b & f, 100 μ M in rows c & g and 1 mM in rows d & h. PPD was added at 5 μ g/ml in rows b & f, 10 μ g/ml in rows c & g and 20 μ g/ml in rows d & h. The ELISpot photograph and number of spots (bottom of each panel) were generated mechanically using an ELISpot Reader System ELR02 (Autoimmun Diagnostika; Strassberg). (b) Number of specific spots (sample minus background) per 100 000 PBMC (■) and $\gamma\delta$ -depleted PBMC (□) induced by specified antigens. Bars represent spots induced by 50 mM ethylamine, 50 mM secbutylamine, 10 mM isobutylamine (50 mM isobutylamine was toxic to cells), 1 mM IPP and 20 μ g/ml PPD. Error bars show the standard deviation from the mean of two replicate wells.

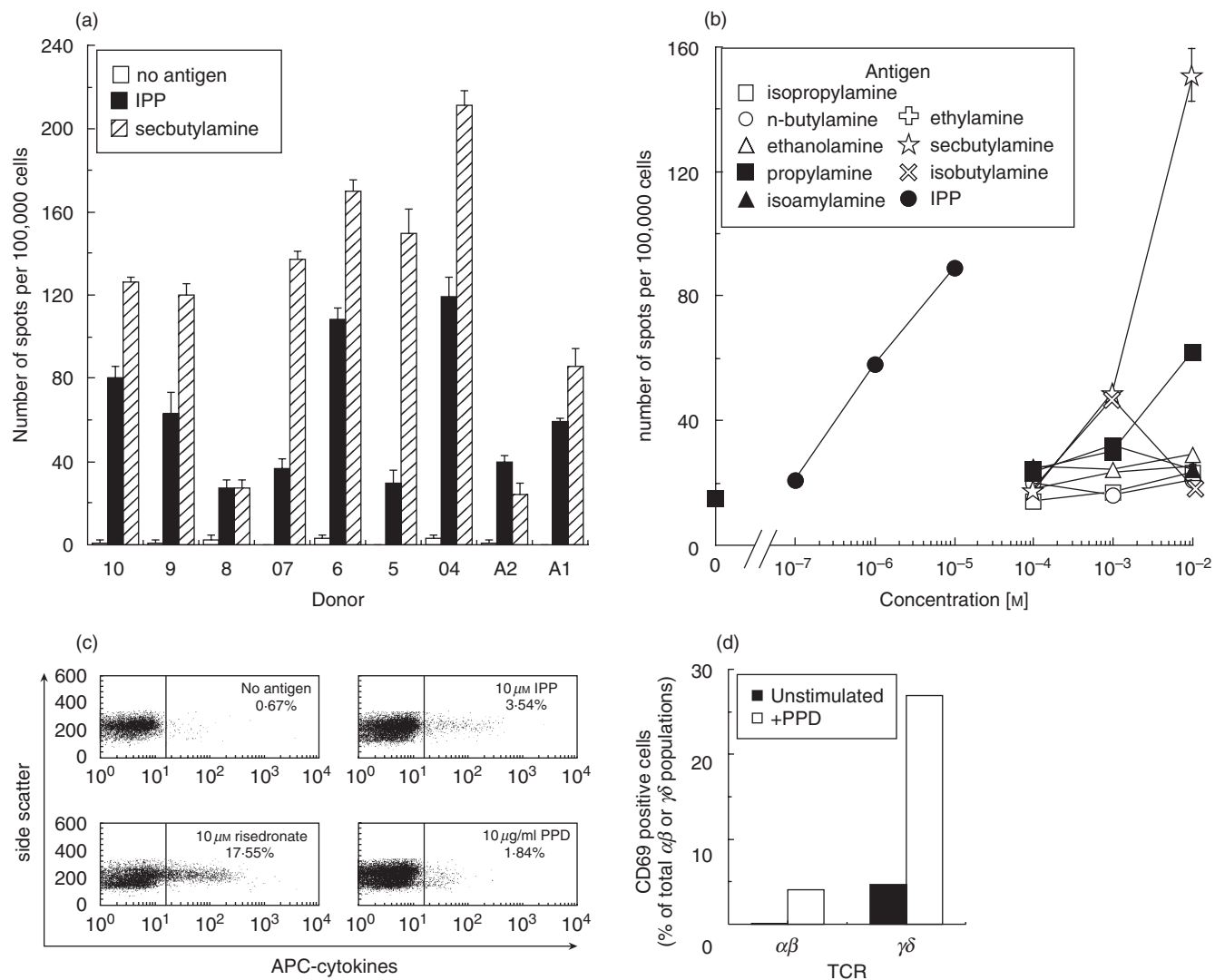


Fig. 2. Further characterization of direct *ex vivo* responses to $\gamma\delta$ T cell antigens. (a) IFN γ ELISpot responses to 100 μ M IPP and 10 mM secbutylamine in PBMC from 9 separate individuals. All individuals also responded to n-butylamine, propylamine and isobutylamine (data not shown) although the response to these antigens was generally lower than that with secbutylamine. Bars show the standard deviation from the mean of two replicate assays. (b) The effect of antigen concentration on the activation of $\gamma\delta$ T cells directly *ex vivo*. The number of spots per 100 000 input PBMC is shown. PBMC from other individuals gave similar results (data not shown). Isobutylamine is toxic at high concentration [32]. Ethylamine, isopropylamine, ethanolamine, and isoamylamine, previously reported to be antigenic [32], did not induce spots. Error bars show standard deviation from the mean of two replicate assays. The errors were generally smaller than the plot symbols. (c) 10^6 fresh PBMC from a BCG vaccinated healthy donor were exposed to indicated antigens for 6 h and then stained with anti-V γ 9 antibody and for intracellular cytokines (IFN γ , TNF α and IL2) as described in the Materials and Methods. Plots show side scatter vs. staining with APC-cytokine antibodies for the V γ 9 positive population and indicate the fraction of cytokine positive cells. (d) PPD induced CD69 up-regulation in almost 30% of human peripheral blood $\gamma\delta$ T cells. PBMCs were washed in RPMI and resuspended in RPMI + 10% FCS at 10^6 cells/ml and incubated \pm 10 μ g/ml PPD at 37°C for 12 h. Cells were stained with FITC-conjugated anti-CD69 antibody, PE-conjugated anti-V γ 9 antibody and PerCP-conjugated anti-CD3 antibody prior to FACS analysis.

amounts of MIP1 β in response to anti-CD3 cross-linking, IPP, secbutylamine and risedronate (Fig. 4c). In contrast, the V γ 9V δ 1 clone only responded to anti-CD3 cross-linking and was unable to recognize alkylphosphate, alkylamine or nBP antigens (Fig. 4c). IFN γ ELISpot analysis confirmed that the V γ 9V δ 2 clone activated in response to IPP, secbutylamine and isobutylamine (Fig. 4d). Activation of the V γ 9V δ 2 clone by low molecular weight nonpeptidic compounds required the presence of both antigen-presenting cells and antigen (data not shown). Curiously,

PPD was unable to activate the V γ 9V δ 2 T cell clone. This result contrasts with those of another study which indicated that all V γ 9V δ 2 T cells respond to *M. tuberculosis* extract including those that are CD4-CD8-, CD8+ or CD4+ [40]. However, these differences might be due to the fact that *M. tuberculosis* extracts are known to contain pyrophosphate antigens that are lost in the production of PPD. Neither alkylphosphate or alkylamine antigens were able to activate the V γ 9V δ 1-expressing clone in IFN γ ELISpot assay (data not shown).



Fig. 3. The γ and δ chain sequences of $\gamma\delta$ T cell clones. The TCR sequences for the V γ 9V δ 1 (top box) and V γ 9V δ 2 (bottom box) T cell clones (IMGT nomenclature [36] is used throughout this work). The γ and δ chain sequences for the V γ 9V δ 1 clone have the accession numbers AJ583014 and AJ583015, respectively, at the EMBL Nucleotide Sequence Database. The accession numbers AJ583012 and AJ583013 have been assigned to the γ and δ chains for the V γ 9V δ 2 clone. The CDR1, 2 and 3 sequences are indicated by shaded boxes. The differences in the V γ 9 CDR3 region between the V δ 1 and V δ 2 clones are underlined. Both V γ 9 chains use the J γ P joining region which contains two germline encoded lysine residues (shown in bold text within the CDR3 region) that have been shown to be essential for recognition of alkylphosphate and alkylamine ligands [46].

V γ 9V δ 2 T cells respond to alkylphosphate/alkylamine antigens and nBP antigens with different kinetics

Recent studies have shown that nBP antigens can be pulsed onto tumour cells [41,42]. We used IFN γ ELISpot to examine whether alkylphosphate, alkylamine and aminobisphosphonate antigens could be pulsed onto antigen presenting cells (Fig. 5). Alkylphosphate and alkylamine antigens were unable to induce lasting changes to antigen presenting cells and activation was only observed when these antigens were present throughout the assay. In contrast, the nBP antigen, pamidronate, was able to effect changes in antigen presenting cells so that they remained antigenic even after washing (Fig. 5).

$\gamma\delta$ T cell activation by phosphate and alkylamine antigens requires contact with cells of human origin

Analysis of T cell activation by IFN γ ELISpot has the benefit of utilizing very few cells. Such a low number of cells spread out in a 96-well flat bottomed plate are unlikely to contact each other. Indeed no spots were seen if antigen was added in the absence of presenting cells (Fig. 6). Antigen recognition could be restored by

the addition of cell lines of human origin. The appearance of spots when V γ 9V δ 2 cells were plated at a higher density indicated that $\gamma\delta$ T cells were able to present antigen to themselves. The mouse myeloma P3/NS1/1-Ag4-1 (ECACC 85011427) and the rat myeloma Y3.AG.1.2.3 (ECACC 85110502) cell lines were unable to 'present' phosphate or alkylamine antigens to V γ 9V δ 2 T cells. We tested nine different human cell lines, including those deficient in β 2-microglobulin and MHC class I, and were unable to find a nucleated human cell line that did not present these antigens, or nBP antigens, to V γ 9V δ 2 T cells (data not shown). Activation always required both antigen presenting cells (APC) and antigen. Human erythrocytes were unable to function as APC in our assays (data not shown).

DISCUSSION

Recent studies have documented that T cells expressing a V γ 9V δ 2 TCR, usually the dominant subset of $\gamma\delta$ T cells in human peripheral blood, can activate in response to alkylphosphate, alkylamine or nBP antigens [31–33,39]. We have extended these observations

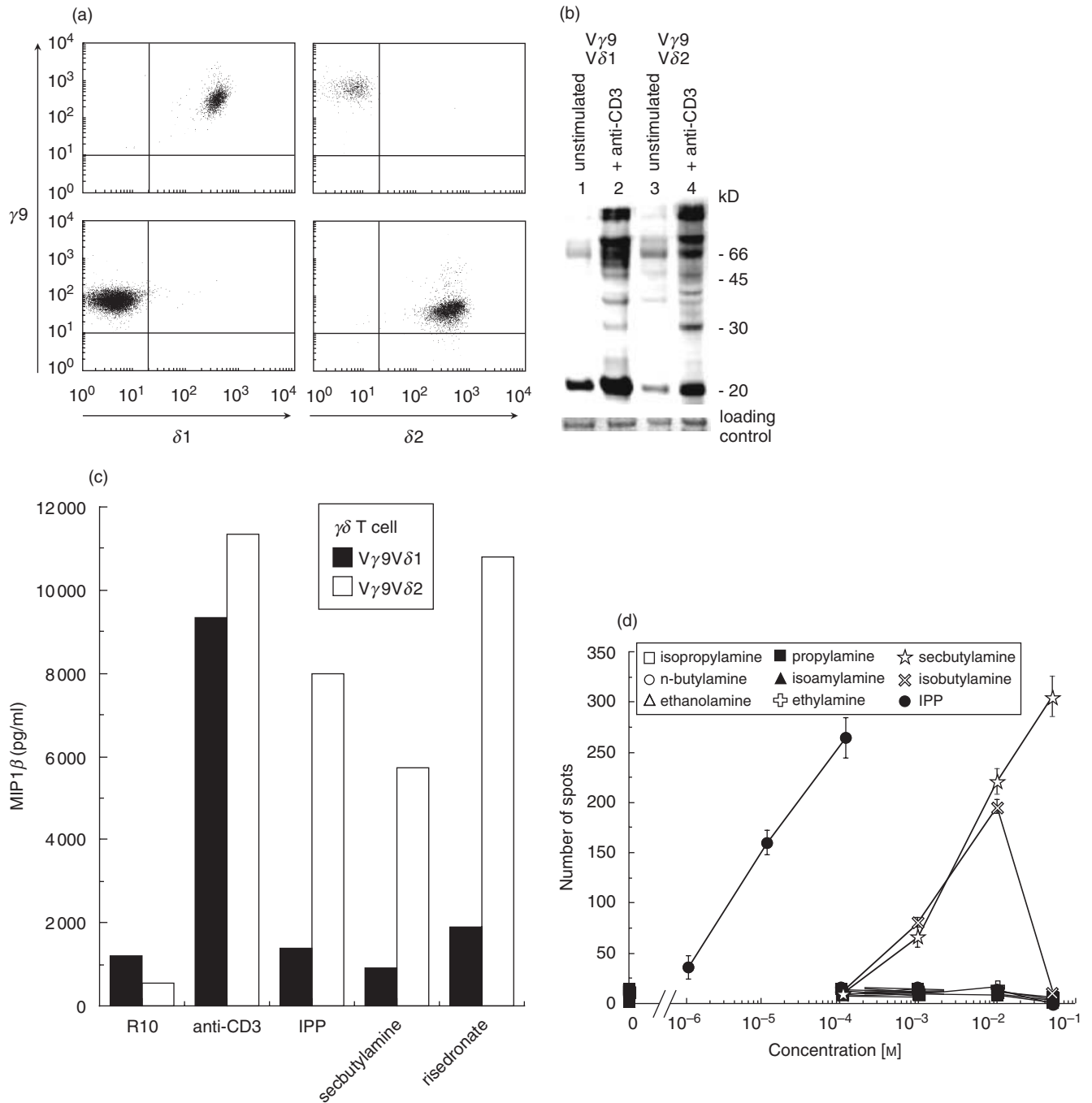


Fig. 4. V $\gamma 9$ V $\delta 1$ and V $\gamma 9$ V $\delta 2$ T cells respond to different antigens. (a) Both the V $\gamma 9$ V $\delta 1$ (upper panels) and V $\gamma 9$ V $\delta 2$ (lower panels) T cell clones were shown to express high levels of TCR by FACS. Clones were stained with PE-conjugated anti-human V $\gamma 9$ mAb clone B3-1 (all panels), FITC-conjugated anti-human V $\delta 1$ mAb clone TS8-2 (left panels) and FITC-conjugated anti-human V $\delta 2$ mAb clone B6-1 (right panels). (b) Both clones can be activated through their TCR by anti-CD3 antibody. Clones were stimulated with anti-CD3 antibody (10 μ g/ml) for 3 min, washed once with PBS, and lysed, before separation by SDS-PAGE alongside unstimulated controls. Proteins bearing phosphorylated tyrosine residues were revealed by immunoblotting with anti-phosphotyrosine antibody clone 4G10 as described [38]. The lysates of 10^6 cells per lane were loaded as follows: V $\gamma 9$ V $\delta 1$ T lymphocytes before (lane 1) and after (lane 2) stimulation; V $\gamma 9$ V $\delta 2$ T lymphocytes before (lane 3) and after (lane 4) stimulation. (c) V $\gamma 9$ V $\delta 2$, but not V $\gamma 9$ V $\delta 1$ T cells, are able to recognize alkylphosphate, alkylamine and nBP antigens. MIP1 β release from 5×10^4 clonal T cells as described in the Materials and Methods section is shown in response to 1.5 μ g/ml anti-CD3 antibody clone UCHT-1, 10 μ M IPP, 10 mM secbutylamine and 1 μ M risedronate. (d) IFN γ ELISpot assay with a titration of potential antigens. 1000 V $\gamma 9$ V $\delta 2$ T cells were used with 25 000 EBV transformed B cells (Sparky line) per well as antigen presenting cells. The V $\gamma 9$ V $\delta 1$ clone failed to activate in response to any of these antigens (data not shown).

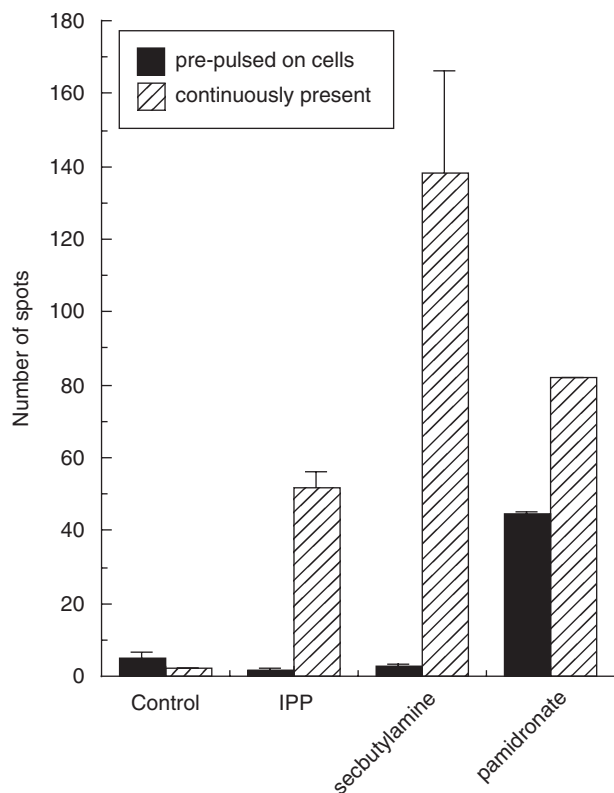


Fig. 5. Alkylamine and alkylphosphate antigens exhibit different pharmacokinetics than nBP antigens. Mean number of spots induced by IPP and secbutylamine when 1000 cells from the $V\gamma 9V\delta 2$ T cell clone Bob were incubated with 25 000 BCL line Sparky in the continuous presence (striped bars) of 100 μM IPP, 10 mM secbutylamine and 1 mM pamidronate during a 12 h $\text{IFN}\gamma$ ELISpot. The filled bars are from a parallel assay where the BCL were incubated for 12 h in R10 containing the above concentrations of antigens and then washed once and resuspended in fresh media without antigen prior to the experiment. The 'control' targets were not pulsed with antigen. Bars show the standard deviation from the mean of two replicate assays. Only pamidronate could be 'pulsed' onto cells. Similar results were obtained with $\gamma\delta$ T cell lines from three separate individuals (data not shown).

by showing that all of these antigens can stimulate a population of $\gamma\delta$ T cells in direct *ex vivo* $\text{IFN}\gamma$ ELISpot assay. The numbers of $V\gamma 9V\delta 2$ T cells measured by FACS analysis of PBMC indicate that only approximately 10% of these cells respond to alkylphosphate, alkylamine and nBP antigens. This may reflect the proportion of cells that recognize these antigens or the proportion that make $\text{IFN}\gamma$ when they respond. However, only approximately 30% of a clonal $V\gamma 9V\delta 2$ T cell population were able to produce spots in $\text{IFN}\gamma$ ELISpot assay (Fig. 4d). This percentage is roughly the same as that observed for more than 20 different $\alpha\beta$ TCR-expressing cytotoxic T lymphocyte (CTL) clones in response to cells presenting their cognate peptide-MHC antigen (unpublished data). We also show that the number of cells that activate in $\text{IFN}\gamma$ ELISpot is dependent on the antigen-presenting cell (Fig. 6). Variation in the ability of different cells to act as presenting cells may reflect differences in the surface expression of ICAM-1 [42] or differences in the surface expression a true $\gamma\delta$ T cell ligand presentation molecule. A recent study defined four subsets of human $V\delta 2$ T cells in peripheral blood based on the expression of CD45RA and CD27 [43]. Only the CD45RA⁺CD27⁻ population

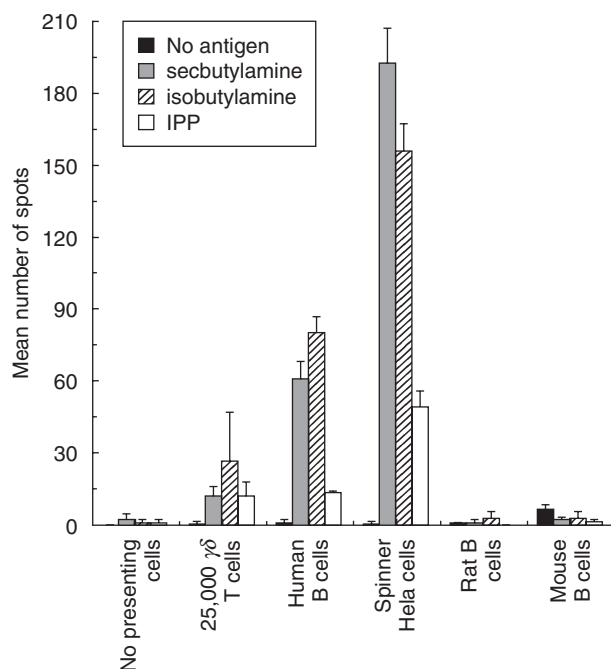


Fig. 6. Recognition of alkylamine and alkylphosphate antigens by $V\gamma 9V\delta 2$ T cells requires contact with cells of human origin. Mean number of spots in $\text{IFN}\gamma$ ELISpot assay with and without antigen presenting cells. 1000 $\gamma\delta$ T cells/well were plated with 25 000 EBV-transformed human B cells, Spinner Hela cells, rat myeloma Y3.AG.1.2.3 or mouse myeloma P3/NS1/1-Ag4.1. Only human cells were able to act as antigen presenting cells in this assay. The use of 25 000 $\gamma\delta$ T cells allowed cell-cell contact and confirms that these cells have some ability to present alkylphosphate and alkylamine antigens to each other. Secbutylamine and isobutylamine were added to assays at 10 mM; IPP was added at 100 μM . Assays were performed over 12 h as described in the Materials and Methods. Standard deviation from the mean of four replicates is shown.

of $V\delta 2$ -expressing cells were able to make large amounts of $\text{IFN}\gamma$ in response to IPP [43]. CD45RA⁺CD27⁻ were found to make up $26.5 \pm 15.3\%$ of $V\delta 2$ T cells in the peripheral blood of 15 healthy adults [43]. It is possible that we are only able to detect this population of $V\delta 2$ cells in our $\text{IFN}\gamma$ ELISpots. We also find that in more than half the individuals we examined, 25–35% of the cells that produce $\text{IFN}\gamma$ in response to PPD express a $\gamma\delta$ TCR (Fig. 1). PPD-induced activation of $\gamma\delta$ T cells was confirmed by FACS analysis (Fig. 2c). These data are in accordance with early studies that show an expansion of $\gamma\delta$ T cells in the lungs of mice exposed to aerosolized PPD [44] as well as those showing expansion of the human $V\gamma 9V\delta 2$ T cell population *in vivo* in *M. tuberculosis* infection [4] and after BCG vaccination [29]. We have not determined the exact nature of the $\gamma\delta$ T cell ligands in PPD and it is possible that the PPD-induced activation of $\gamma\delta$ T cells could have been due to low-level contamination of this preparation with mycobacterial alkylphosphate antigens or to the presence of Toll-like receptor ligands.

To examine their activation requirements further, we cloned $V\gamma 9$ -expressing T cells from peripheral blood. We found that $V\gamma 9V\delta 1$ T cells are unable to recognize the plethora of nonpeptide antigens recognized by cells expressing $V\gamma 9$ in its more common pairing with a $V\delta 2$ chain (Fig. 4c). This result is in agreement with those of a previous study that showed $V\gamma 9V\delta 1$ T cell clones failed to respond to monoethyl phosphate [31]. A previous study

demonstrated that transfection of TCR-negative Jurkat T cells with a V γ 9V δ 2 TCR, but not a V δ 1 TCR (not paired with V γ 9), enabled recognition of alkylphosphate antigen [45]. These elegant experiments further highlight that T cell recognition of such antigens requires the expression of a specific $\gamma\delta$ TCR. Minato and colleagues have recently extended these observations by showing that recognition of alkylphosphate, alkylamines and nBP antigens by V γ 9V δ 2 T cells requires that the V γ 9 segment be joined to a J γ P joining region [46]. Previous experiments have shown that the proportion of V9JPC1 γ chains is significantly higher in the peripheral blood than in thymocytes [47]. The peripheral expansion of V γ 9 cells using J γ P appears to be antigen-driven [48]. The $\gamma\delta$ T cells in cord blood mononuclear cells (CBMC) express a diverse repertoire of $\gamma\delta$ TCRs [48]. Stimulation of CBMC with alkylphosphate antigen, but not with PHA, induces the preferential expansion of T cells bearing V γ 9V δ 2 TCRs [48]. 70% of TCRs expanded with alkylphosphate antigen use a J γ P joining region compared to just 20% of those cells expanded with PHA [48]. The J γ P joining region differs from other possible joining regions in that it contains two germline encoded lysine residues (K₁ & K₂) in addition to a third lysine (K₃) that is conserved in all J γ [36,46]. Mutation of these lysine residues completely abrogated the ability of these TCRs to respond to nonpeptide antigens [46]. Both the V γ 9V δ 1 and V γ 9V δ 2 clones tested here bear V γ 9JP chains (Fig. 3). As only the V δ 2 clone is able to respond to alkylphosphate, alkylamine and nBP antigens (Fig. 4c) it would appear that the V δ 2 chain may also make an essential contribution to the recognition of these ligands. Recent results appear to confirm that specific features of the V δ 2 chain are required for recognition of nonpeptide antigens by V γ 9V δ 2 T cells [48]. It is further noteworthy that the amount of MIP1 β manufactured by V γ 9V δ 2 T cells (Fig. 4c) is comparable to that we have measured after antigen-induced activation of CTL [37]. The release of β -chemokines by $\gamma\delta$ T cells may act to recruit immune cells to sites of infection [49].

We show that a clonal population of V γ 9V δ 2 T cells can recognize alkylphosphate, alkylamine and nBP ligands (Figs 4 and 5). Bukowski *et al.* [32] also demonstrated that a single T cell clone could recognize two types of ligand; alkylamines and IPP. The authors suggest that, given the small sizes of these antigens, a single $\gamma\delta$ TCR might either possess two different binding sites or a single binding site that can accommodate both types of ligand. Our finding that the recognition of both these antigens and nBP antigens requires cell surface presentation makes it highly unlikely that there are separate binding sites for these classes of ligand. Indeed, it is very difficult to envisage just how single V γ 9V δ 2 TCRs are able to respond to such different chemical species and yet still exhibit tight structural constraints within each class of ligand. The most obvious similarity between the recognized ligands is their alkyl chains. While it is known that phosphate antigens require their phosphate and nBPs and alkylamines require their amino groups for their antigenicity [32,33,39] it is possible that this is necessary for anchoring them to a presenting molecule rather than forming a component of the TCR-docking structure. Indeed, the finding that both antigenic alkylphosphate and alkylamine ligands require similar alkyl chains for recognition lends some credence to such a hypothesis and has led to attempts at molecular prediction [50]. However, it is noticeable that the most potent nBP antigen to date, risedronate, which can stimulate $\gamma\delta$ T cell expansion from human PBMC at concentrations as low as 10 nM [35] was not included in this predictive

exercise [50]. We have also shown that risedronate can activate V γ 9V δ 2 T cells in IFN γ ELISpot assay at concentrations of <1 μ M (Fig. 4c and data not shown). Risedronate has an aromatic ring and does not fit with the pattern recognition hypothesis championed by Gossman and Oldfield [50].

It is noteworthy that antigenic nBPs, and not their corresponding nonantigenic nonaminobisphosphonates (BPs), are potent inhibitors of the mevalonate pathway (reviewed in [51]). It must be no coincidence that IPP, a potent activator of V γ 9V δ 2 T cells, is part of this metabolic pathway. Indeed, treatment of cells with the nBPs zoledronate and pamidronate has recently been shown to lead to an accumulation of IPP [52]. It thus seems likely that antigenic nBPs exert their effects via IPP and not directly as previously assumed [41,42,50]. The results of De Libero and colleagues [52] elegantly demonstrate this mode of action for nBPs by two different means. First, they show that nBPs, unlike IPP, require internalization in order to exert their effects [52]. Second, they show that blocking the production of IPP inhibits the ability of nBPs to activate $\gamma\delta$ T cells [52]. A model in which nBPs function via metabolites in the mevalonate pathway conveniently explains some current anomalies in the activation of $\gamma\delta$ T cells by these compounds. Firstly, it indicates how the nBPs risedronate (Fig. 5) and zoledronate [52], the aromatic ring structures of which do not fit with the established pattern recognition motif for $\gamma\delta$ antigens [50], might lead to activation. Secondly, it explains why BPs that do not inhibit the mevalonate pathway [51] fail to act as antigens ([52] and our unpublished data). This mechanism of action also explains why the nBP pamidronate has been described as being 'rather unique' amongst non peptide antigens in that it could be pulsed onto antigen presenting cells and retain its antigenicity after these cells had subsequently been washed ([41,42] and Fig. 5). However, although unlikely, it still remains possible that nBPs activate V γ 9V δ 2 T cells directly. Some of the findings of Gober *et al.* [52] could potentially be due to the toxic effects of statins. Statins and bisphosphonates have inhibitory effects on almost all cells as blocking the mevalonate pathway leads to the loss of Ras and other receptor signals due to the blockade of protein prenylation [53]. nBPs block farnesyl synthase and may act synergistically with inhibitors of 3-hydroxy-3-methylglutaryl coenzyme A (HMG CoA) reductase to kill cells. These effects may be overcome by further addition of intermediates in the mevalonate pathway. The significance of these issues is apparent in the clinical setting. nBPs have become well established as clinically successful antiresorptive agents for the treatment of osteoporosis (reviewed in [51]) and, more recently, multiple myeloma [54,55]. These drugs can induce a transient fever and biological changes, including specific cytokine production, that are suggestive of an acute phase response (e.g. [56–58]). It is possible that the build up of IPP and other metabolites resulting from the actions of these drugs on the mevalonate pathway and the ensuing activation of $\gamma\delta$ T cells *in vivo* might contribute to these side-effects. Further work aimed at testing this hypothesis is required.

Any attempt to understand the recognition requirements of $\gamma\delta$ T cells should also take the absolute requirement for species-specific cell-cell contact into account. The use of IFN γ ELISpot has allowed us to conclusively prove the need for cell-cell contact with a cell of human origin. The results of Morita *et al.* [40] showing that monoethyl phosphate-induced calcium flux in V γ 9V δ 2 T cells required that the cells be centrifuged and resuspended, and those of Lang *et al.* [59] showing that V γ 9V δ 2 T cells responded to

mycobacterial phosphoantigens poorly when kept in suspension first hinted that cell-cell contact might be important for activation. During the preparation of this manuscript another group have shown that recognition of the nBP pamidronate also requires species-specific cell-cell contact [42]. This study showed that 33 tumour cell lines from 12 different species, including other primates, failed to mediate $V\gamma 9V\delta 2$ T cell activation in response to pamidronate [42]. Curiously, pamidronate did not activate $\gamma\delta$ T cells when 'presented' by 5 human tumour cell lines [42]. Recognition of some, but not all, of these cell lines could be restored by transfection with ICAM-1 [42]. However, pamidronate likely functions indirectly via its actions on IPP synthesis rather than as a direct 'ligand' as this study assumed and it remains possible that these cell lines have an abnormal mevalonate pathway. It is important that these experiments be repeated with IPP itself. Human cell lines that fail to present $V\gamma 9V\delta 2$ T cell antigens are likely to be very useful tools in the elucidation of the antigen-presentation mechanism.

That $\gamma\delta$ T cell activation requires cell-cell contact with a cell of human origin (Fig. 6) opens up a number of possibilities. First, it could be indicative of a human-specific $\gamma\delta$ T cell ligand-presentation molecule. The ability of multiple cell lines from different tissues and individuals to act as antigen-presenting cells argues against any kind of large variability in this molecule as seen for the MHC genes. Alkylphosphate and alkylamine antigens are unable to effect stable changes in the antigen presenting cells as their effects are readily removed by dilution and washing [40] (Fig. 5, and our unpublished observation). This argues against major changes in the surface of the presenting cell such as up-regulation of a $\gamma\delta$ TCR ligand or heat shock proteins such as MIC-A and MIC-B [60–62]. Second, the requirement for presentation by a human cell may be indicative of a human specific coreceptor element or the requirement for additional costimulatory signals. Distinguishing between these various models awaits the characterization of the true ligand for peripheral blood $V\gamma 9V\delta 2$ T cells. The recent production of a high purity, soluble $V\gamma 9V\delta 2$ TCR [63] should aid in this process. Only when we understand exactly what the $\gamma\delta$ TCR recognizes will we be able to unlock the forgotten third arm of adaptive immunity that $\gamma\delta$ T cells control.

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