RESEARCH ARTICLE

Human V γ 9V δ 2-T cells efficiently kill influenza virus-infected lung alveolar epithelial cells

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$\gamma\delta$ -T cells play an indispensable role in host defense against different viruses, including influenza A virus. However, whether these cells have cytotoxic activity against influenza virus-infected lung alveolar epithelial cells and subsequently contribute to virus clearance remains unknown. Using influenza virus-infected A549 cells, human lung alveolar epithelial cells, we investigated the cytotoxic activity of aminobisphosphonate pamidronate (PAM)-expanded human V γ 9V δ 2-T cells and their underlying mechanisms. We found that PAM could selectively activate and expand human V γ 9V δ 2-T cells. PAM-expanded human V γ 9V δ 2-T cells efficiently killed influenza virus-infected lung alveolar epithelial cells and inhibited virus replication. The cytotoxic activity of PAM-expanded V γ 9V δ 2-T cells was dependent on cell-to-cell contact and required NKG2D activation. Perforin–granzyme B, tumor-necrosis factor-related apoptosis-inducing ligand (TRAIL) and Fas–Fas ligand (FasL) pathways were involved in their cytotoxicity. Our study suggests that targeting $\gamma\delta$ -T cells by PAM can potentially offer an alternative option for the treatment of influenza virus.

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INTRODUCTION

Although they represent only a small population of immune cells, $\gamma\delta$ -T cells exhibit features characteristic of both innate and adaptive immunity and play an indispensable role in host defense, immune surveillance and homeostasis.^{1–4} In humans, $\gamma\delta$ -T cells constitute approximately 1%–5% of circulating T cells, and most of them bear the V γ 9V δ 2 T-cell receptor (TCR).^{5,6} Human V γ 9V δ 2-T cells can be specifically activated in an HLA-unrestricted manner by small non-peptidic phosphoantigens, which are metabolites of the isoprenoid biosynthesis pathways.⁷ It is known that isopentenyl pyrophosphate (IPP), an intermediate produced through the mevalonate pathway, can selectively activate and expand human V γ 9V δ 2-T cells.^{8–10} Pharmacological compounds, such as the aminobisphosphonate pamidronate (PAM), which is commonly used for the treatment of osteoporosis, can activate and expand

human V γ 9V δ 2-T cells by inducing the intracellular accumulation of IPP.^{8,11}

$\gamma\delta$ -T cells are broadly reactive against different viruses,^{12–18} such as herpes viruses (herpes simplex virus, cytomegalovirus, human herpes virus-6), vaccinia virus, influenza virus, coxsackie B virus and human or simian immunodeficiency virus (HIV/SIV), indicating a role for these lymphocytes in antiviral immune responses. Cell-mediated cytotoxicity is the major mechanism to eliminate virus-infected cells and thus to eliminate potential sources of new virus. Previously, using influenza virus-infected human monocyte-derived macrophages, we showed that IPP-expanded human V γ 9V δ 2-T cells had cytotoxic activity against virus-infected monocyte-derived macrophages.¹⁰ However, macrophages are not the major target cells for influenza virus infection, despite the influenza virus' ability to replicate in human macrophages.¹⁹ Human lung alveolar

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epithelial cells, as the major target cells for influenza virus infection, are a more reliable target to investigate the antiviral role of $\gamma\delta$ -T cells. Recently, we demonstrated that IPP-expanded human V γ 9V δ 2-T cells had a non-cytolytic effect on human influenza virus by releasing IFN- γ in virus-infected lung alveolar epithelial cells.²⁰ However, whether these cells have cytolytic effects in influenza virus-infected lung alveolar epithelial cells, and subsequently contribute to virus clearance, is still unknown.

In this study, using influenza virus-infected human lung alveolar epithelial cells, we investigated the cytotoxic activity of PAM-expanded human V γ 9V δ 2-T cells and their underlying mechanisms. We found that PAM-expanded human V γ 9V δ 2-T cells efficiently killed influenza virus-infected lung alveolar epithelial cells and inhibited virus replication. The cytotoxic activity of PAM-expanded V γ 9V δ 2-T cells was dependent on cell-to-cell contact and required NKG2D activation. The pathways of perforin–granzyme B, tumor-necrosis factor-related apoptosis-inducing ligand (TRAIL) and Fas–Fas ligand (FasL) mediated their cytotoxicity.

MATERIALS AND METHODS

Cells

Human peripheral blood was obtained from healthy donors in accordance with the Hong Kong University Institutional Review Board. Peripheral blood mononuclear cells (PBMCs) were isolated by Ficoll-Hypaque (Pharmacia, Piscataway, NJ, USA) gradient centrifugation, as previously described.²¹ PBMCs were cultured in 10% fetal bovine serum RPMI-1640 medium with 9 μ g/ml of PAM. Recombinant human IL-2 (Invitrogen, Carlsbad, CA, USA) was added to reach a final concentration of 500 IU/ml every 3 days from day 3. After being cultured for 20 days, V γ 9V δ 2-T cells were purified by negative selection with a TCR $\gamma\delta$ -T-cell isolation kit, according to the manufacturer's instruction (Miltenyi Biotec, Bergisch Gladbach, Germany). The purity of $\gamma\delta$ -T cells, as determined by flow cytometry, was consistently >97%. Immortalized human alveolar type II epithelial cells (A549) were cultured in RPMI 1640 medium supplemented with 10% bovine calf serum.

Virus preparation, titration and infection

Influenza H1N1 virus (A/PR/8/34) was used. The virus was cultured in Madin–Darby canine kidney cells as previously described.^{22,23} The virus titer was determined by daily observation of the cytopathic effect in cells infected with serial dilutions of virus stock, and the median tissue culture infective dose (TCID₅₀) was calculated according to the Reed–Muench formula. A549 cells were infected by influenza virus at a multiplicity of infection (MOI) of 2. After 1 h of viral adsorption, the cells were washed by phosphate-buffered saline to remove unadsorbed virus.

Flow cytometry

Cells were stained for surface markers with the following antibodies: anti-CD3, anti-TCR γ 9, anti-TCR δ 2, anti-NKG2D, anti-CD69, anti-TRAIL, anti-TCR- $\gamma\delta$, anti-MICA/B, anti-

Fas, anti-FasL, anti-TRAIL and their ligands DR4 and DR5. For intracellular staining, cells were fixed, permeabilized and then stained with anti-perforin and anti-granzyme B (GrB, GB11) antibodies or their relevant isotype controls. All samples were acquired by flow cytometry and analyzed by FlowJo software (Tree Star, Inc., Ashland, OR, USA).

Cytotoxicity assay

Purified V γ 9V δ 2-T cells (effector) were cocultured with H1N1 virus-infected or mock-infected A549 cells (target) at different effector-to-target (E/T) ratios for 5–6 h. Afterward, non-adherent cells were harvested directly. Adherent cells were detached with 0.25% trypsin-ethylenediaminetetraacetic acid. All of the adherent and non-adherent cells were then stained with anti-human CD3 to identify V γ 9V δ 2-T cells and ethidium homodimer-2 (EthD-2) to identify dead cells. The cytotoxicity of V γ 9V δ 2-T cells against A549 cells was assessed by flow cytometry as the percentage of EthD-2⁺ cells in the CD3⁺ population, as previously described.¹⁰

Transwell coculture

To evaluate the cell–cell contact requirement for V γ 9V δ 2-T-cell cytotoxicity, a transwell culture system (24-well, pore size 0.4 μ m; Millipore, Bedford, MA, USA) was used. A549 cells (Target, T) in the bottom well were infected with influenza virus at an MOI of 2, and V γ 9V δ 2-T cells (Effector, E) were added directly into the bottom wells or into transwell inserts at an E/T ratio of 10:1. After culturing for 5 h, the A549 cells in the bottom wells were harvested and analyzed for cell death, as described above. The supernatants in the transwell coculture of V γ 9V δ 2-T cells and A549s were collected at the indicated time for determining the virus titer, as described above.

Blocking assay

Purified V γ 9V δ 2-T cells (E) were cocultured with influenza virus-infected A549 cells (T) at an E/T ratio of 10:1 for 5 h. The neutralization antibodies anti-NKG2D (10 μ g/ml; BD Biosciences, San Jose, CA, USA), anti-FasL (10 μ g/ml; R&D, Minneapolis, MN, USA), anti-TRAIL (10 μ g/ml; R&D) and their relevant isotype control mouse IgG1 (mIgG1) were added to the coculture for blocking NKG2D, FasL and TRAIL mediated pathways, respectively. For blocking perforin and granzyme B, the perforin inhibitor concanamycin A (CMA) (1 μ g/ml; Sigma, St Louis, MO, USA) and granzyme B inactivator Bcl-2 (1 μ g/ml; R&D) were used, as in previous reports.²⁴ The cytotoxicities were analyzed by flow cytometry as described above and calculated as % inhibition relative to those of the control.

Statistical analysis

Data are expressed as the mean $\pm$ standard errors of the mean. Statistical significance was determined by the Student's *t*-test or nonparametric tests using Graphpad Prism software (version 5). A *P* value of <0.05 was considered to be significant.

Figure 1 Phenotypes of fresh and PAM-expanded V γ 9V δ 2-T human cells. The white histograms represent the surface expression of CD69, NKG2D, MIC A/B, Fas, FasL, TRAIL, DR4 (TRAIL receptor 1), DR5 (TRAIL receptor 2), intracellular perforin and granzyme B, and the gray histograms represent isotype controls. Data shown here are representative of four separate experiments. FasL, Fas–Fas ligand; PAM, aminobisphosphonate pamidronate; TRAIL, tumor-necrosis factor-related apoptosis-inducing ligand.

RESULTS

PAM selectively activates and expands human V γ 9V δ 2-T cells

Similar to IPP,^{25,26} PAM and IL-2 can also selectively expand human V γ 9V δ 2-T cells. Freshly isolated PBMCs contained 1%–6% of V γ 9V δ 2-T cells from 16 randomly selected healthy adult donors; after 20 days of culture in the presence of PAM and IL-2, the percentage of V γ 9V δ 2-T cells within PBMC increased to 85%–97% and the absolute numbers of the V γ 9V δ 2-T cells were significantly increased by 93-fold (range: 54–151-fold). In contrast, IL-2 alone did not increase the absolute number of V γ 9V δ 2-T cells.

To determine whether PAM can activate human V γ 9V δ 2-T cells, we examined cell surface markers (CD69, NKG2D, Fas, FasL and TRAIL) and intracellular cytolytic granules (perforin and granzyme B) in fresh and PAM-expanded V γ 9V δ 2-T cells. As shown in Figure 1, fresh V γ 9V δ 2-T cells expressed low levels of FasL, high levels of Fas and medium levels of CD69, NKG2D, TRAIL, perforin and granzyme B. In contrast, PAM-expanded V γ 9V δ 2-T cells had much higher levels of CD69, NKG2D, FasL, TRAIL, perforin and granzyme B expression compared to fresh V γ 9V δ 2-T cells (Figure 1).

Figure 2 PAM-expanded V γ 9V δ 2-T cells efficiently killed influenza virus-infected A549 cells. A549 cells (Target, T) were either mock infected (A549) or infected with influenza H1N1 PR/8 virus at an MOI of 2 (vA549), and then cultured with purified PAM-expanded V γ 9V δ 2 T cells (Effector, E) at various E/T ratios for 5 h. The percentages (mean $\pm$ s.e.m.) of dead A549 cells among target cells (CD3⁺ population), identified as CD3⁺EthD2⁺, for four different experiments are shown. * P <0.05, ** P <0.01. MOI, multiplicity of infection; PAM, aminobisphosphonate pamidronate.

PAM-expanded V γ 9V δ 2-T cells efficiently kill influenza virus-infected A549 cells

To determine the cytotoxic activity of V γ 9V δ 2-T cells against influenza virus-infected A549 cells, purified PAM-expanded V γ 9V δ 2-T cells were cocultured with mock- or influenza virus-infected A549 cells for 5 h. As shown in Figure 2, V γ 9V δ 2-T cells displayed cytotoxic activity against both mock- and virus-infected A549s in a dose-dependent manner. Importantly, the killing of V γ 9V δ 2-T cells against influenza virus-infected A549 cells significantly increased compared to that against mock-treated A549 cells at E/T ratios of 10:1 or 20:1. These results demonstrate that PAM-expanded V γ 9V δ 2-T cells have potent cytotoxic activity against influenza virus-infected lung alveolar epithelial cells.

The killing of virus-infected A549 cells by V γ 9V δ 2-T cells is dependent on cell-to-cell contact

To clarify whether the killing of influenza virus-infected A549 cells by V γ 9V δ 2-T cells requires cell-to-cell contact, we applied a transwell culture system to separate virus-infected A549 cells from V γ 9V δ 2-T cells at an E/T ratio of 20:1. As shown in Figure 3, V γ 9V δ 2-T cells lost their cytolytic activity against influenza virus-infected A549 cells when the physical contact between V γ 9V δ 2-T cells and virus-infected A549 cells was abrogated by a semipermeable membrane. These results indicate that the killing of influenza virus-infected lung alveolar epithelial cells by V γ 9V δ 2-T cells is dependent on cell-to-cell contact.

Influenza virus replication in A549 cells is inhibited by V γ 9V δ 2-T cells

Influenza virus efficiently replicated within A549 cells. The virus titer in the culture supernatants gradually increased from 6 h post-infection and reached a peak level at 72 h post-infection (Figure 4). To determine whether there was a decline of the viral titer in A549 cells after coculture with V γ 9V δ 2-T cells, virus-infected A549 cells were cultured alone or with expanded V γ 9V δ 2-T cells, either in direct contact with or separate from V γ 9V δ 2-T cells. When separated from V γ 9V δ 2-T cells, culturing was performed by a transwell culture system at an E/T ratio of 20:1. As shown in Figure 4, the virus titers in supernatant

Figure 3 Physical contact was required for cytotoxicity of $\gamma 9V\delta 2$ -T cells against influenza virus-infected A549s. A549 cells (Target, T) were infected (vA549) with influenza H1N1 PR/8 virus at an MOI of 2 and then cultured with or physically separated from the purified PAM-expanded $\gamma 9V\delta 2$ T cells (Effector, E) at E/T ratios of 20:1 for 6 h. The percentages (mean $\pm$ s.e.m.) of dead A549 cells among target cells ($CD3^-$ population), identified as $CD3^-$ EthD2 $^+$, for four different experiments are shown. * P <0.05. MOI, multiplicity of infection; PAM, aminobisphosphonate pamidronate.

culture were significantly reduced after 24, 48 and 72 h of coculture of virus-infected A549 cells and $\gamma 9V\delta 2$ -T cells, compared to virus-infected A549 cells alone. In contrast, there was no significant decrease in virus titers when $\gamma 9V\delta 2$ -T cells were separated from virus-infected A549 cells (Figure 4). These data suggest that the inhibition of influenza virus replication by $\gamma 9V\delta 2$ -T cells requires cell-to-cell contact.

Mechanisms underlying the cytotoxicity of $\gamma 9V\delta 2$ -T cells

As PAM-expanded $\gamma 9V\delta 2$ -T cells expressed high or medium levels of NKG2D, Fas, FasL and TRAIL, we further determined whether their relevant ligands were expressed in influenza

Figure 4 Influenza virus replication in A549 cells was inhibited by $\gamma 9V\delta 2$ -T cells. A549 cells (Target, T) were infected (vA549) with influenza H1N1 PR/8 virus at an MOI of 2 and then cultured with or physically separated from the purified PAM-expanded $\gamma 9V\delta 2$ T cells (Effector, E) at E/T ratios of 20:1 for the indicated time. The virus titers of supernatants were determined by $TCID_{50}$ on MDCK cells. Data are the mean $\pm$ s.e.m. of $TCID_{50}$ titers of four separate experiments. MOI, multiplicity of infection; PAM, aminobisphosphonate pamidronate.

virus-infected A549 cells. As shown in Figure 5a, A549 cells expressed high levels of NKG2D ligands (stress-inducible MHC class I-related proteins A and B, MAC A/B),²⁷ Fas and TRAIL receptors (DR4 and DR5),²⁸ but very low levels of FasL. The expressions of MIC A/B and DR5 were upregulated in A549 cells after influenza virus infection. In contrast, there were no significant changes in expressions of Fas, FasL and DR4 in A549 cells after influenza virus infection.

To further determine whether NKG2D, Fas–FasL and TRAIL pathways were involved in the cytotoxicity of $\gamma 9V\delta 2$ -T cells, neutralizing antibodies for NKG2D, FasL and TRAIL were used. As shown in Figure 5, blockades of NKG2D, TRAIL or FasL significantly inhibited the cytolytic activities of $\gamma 9V\delta 2$ -T cells against influenza virus-infected A549 cells. These results indicate that NKG2D, TRAIL and Fas–FasL pathways are

Figure 5 Mechanisms underlying the cytotoxicity of $\gamma 9V\delta 2$ -T cells. (a) Phenotype of influenza virus-infected A549 cells. A549 cells were infected with influenza H1N1 PR/8 virus (MOI=2) and then cultured for 5 h. The expression level of surface MICA/B, Fas, FasL and DR4, DR5 on the virus infected-A549 cells were analyzed by flow cytometry. Data shown here are representative of four separate experiments. (b) PAM-expanded $\gamma 9V\delta 2$ -T cells were cocultured with influenza PR/8 virus-infected A549 cells at a ratio of 10:1 for 5 h. The perforin inhibitor CMA, granzyme B inactivator Bcl-2, anti-FasL, anti-TRAIL anti-NKG2D blocking antibodies, or their relevant isotype control (mouse IgG1, mlgG1), were used. The cytotoxicity was analyzed by flow cytometry as the percentage of EthD-2 $^+$ cells in the $CD3^-$ population and calculated as % inhibition relative to those without any treatment. The data shown as the mean $\pm$ s.e.m. are representative of four independent experiments. ** P <0.01 compared to their relevant isotype control. CMA, concanamycin A; FasL, Fas–Fas ligand; MOI, multiplicity of infection; PAM, aminobisphosphonate pamidronate; TRAIL, tumor-necrosis factor-related apoptosis-inducing ligand.

involved in the killing of influenza virus-infected lung alveolar epithelial cells by V γ 9V δ 2-T cells.

To confirm the involvement of cytolytic granule release in the killing of virus-infected A549 cells by V γ 9V δ 2-T cells, the perforin specific inhibitor CMA and granzyme B inactivator Bcl-2 were used. As shown in Figure 5, the cytolytic activities of V γ 9V δ 2-T cells against influenza virus-infected A549 cells were significantly abrogated after Bcl-2 treatment or CMA treatment. These results demonstrate that the perforin–granzyme B pathway is involved in the cytotoxicity of V γ 9V δ 2-T cells against influenza virus-infected lung alveolar epithelial cells.

DISCUSSION

In this study, using influenza virus-infected lung alveolar epithelial cells as the model, we have demonstrated for the first time that PAM-expanded V γ 9V δ 2-T cells can efficiently kill virus-infected lung alveolar epithelial cells and thus contribute to virus clearance. Our study suggests a novel therapeutic approach of using PAM to activate and expand human V γ 9V δ 2-T cells against influenza virus infection.

We showed that similar to IPP,^{25,26} PAM is a potent stimulator for the activation and expansion of human V γ 9V δ 2-T cells from PBMC in the presence of IL-2. Upon PAM stimulation, resting V γ 9V δ 2-T cells can be activated, as evidenced by the upregulated expressions of CD68, NKG2D, FasL and TRAIL. Importantly, during 20 days of stimulation by PAM and IL-2, V γ 9V δ 2-T cells can be largely expanded by approximately 93-fold, suggesting that PAM could be used for large-scale expansion of functional $\gamma\delta$ -T cells *in vitro* for adoptive immunotherapy in influenza virus infections.

A concern for $\gamma\delta$ -T cell-based immunotherapy is whether PAM-expanded V γ 9V δ 2-T cells can traffic to the lung, the primary infection site, during an influenza infection. Indeed, more recently, we have shown that the *in vitro* PAM-expanded V γ 9V δ 2-T cells can migrate to the lung and control influenza disease in immunodeficient mice.¹¹ In addition, in a humanized mouse model, we further demonstrated that PAM can activate and expand V γ 9V δ 2-T cells *in vivo*, and then control human and avian influenza virus infections.¹¹ Therefore, PAM could be an alternative option for the treatment of influenza virus infection by targeting V γ 9V δ 2-T cells.

The antiviral mechanisms of V γ 9V δ 2-T cells against different viruses are different. For examples, human V γ 9V δ 2-T cells have cytolytic activities against CMV- and herpes simplex virus-infected cells in an HLA-unrestricted manner *in vitro*.^{12,13,29} In addition to killing HIV-infected cells, V γ 9V δ 2-T cells can also block HIV entry through the coreceptor CCR5 by releasing certain CCR5-ligand chemokines.^{17,30} For the hepatitis C virus, V γ 9V δ 2-T cells can induce non-cytolytic inhibition of virus replication through the secretion of IFN- γ .²⁵ Previously, we also demonstrated that IPP-expanded V γ 9V δ 2-T cells can inhibit human influenza H1N1 virus replication by releasing IFN- γ .²⁰ Here, we further found that PAM-expanded V γ 9V δ 2-T cells can kill influenza virus-infected lung alveolar epithelial cells. Thus, our results indicate that PAM-expanded

V γ 9V δ 2-T cells can control influenza virus infection through both cytolytic and non-cytolytic mechanisms.

NKG2D is a potent costimulatory receptor of the cytotoxic functions of human natural killer and V γ 9V δ 2-T cells.³¹ It can trigger V γ 9V δ 2-T cells to release cytolytic granules by recognition of the NKG2D ligand.³² MICA/B, as the ligand for NKG2D, was found to be upregulated in lung alveolar epithelial cells after influenza virus infection in the current study. In addition, we showed that almost all the PAM-expanded V γ 9V δ 2-T cells expressed NKG2D, and the cytotoxicity of V γ 9V δ 2-T cells against influenza virus-infected lung alveolar epithelial was significantly blocked by NKG2D neutralizing antibody, indicating the killing of influenza virus-infected cells by V γ 9V δ 2-T cells requires NKG2D activation and recognition.

Previously, we found that the TRAIL pathway was not involved in the killing of influenza virus-infected macrophages.¹⁰ However, here we found that the blockade of TRAIL by its neutralizing antibody significantly decreased the cytolytic activity of V γ 9V δ 2-T cells against influenza virus-infected lung alveolar epithelial cells. This may be due to the different expression levels of TRAIL receptors; we found that influenza virus-infected lung alveolar epithelial cells expressed high levels of DR4 and DR5, the receptors of TRAIL, whereas virus-infected macrophages only expressed very low levels of DR4 and DR5 (data not shown).

It has been shown that the Fas/FasL-mediated pathway was also involved in the killing of virus-infected macrophages by V γ 9V δ 2-T cells.^{10,33} Our data showed that both influenza virus-infected lung alveolar epithelial cells and PAM-expanded V γ 9V δ 2-T cells expressed high levels of Fas or FasL, and the blockade of the Fas/FasL pathway by FasL neutralizing antibody significantly inhibited the cytolytic activity of V γ 9V δ 2-T cells against influenza virus-infected cells. These findings indicate that the Fas/FasL-mediated pathway is also involved in the killing of influenza virus-infected lung alveolar epithelial cells by V γ 9V δ 2-T cells.

We confirmed that similarly to their cytotoxic function in natural killer and CD8⁺ T cells, perforin and granzyme B facilitated the killing of influenza virus-infected lung alveolar epithelial cells in V γ 9V δ 2-T cells. These findings are consistent with previous studies in tumors and other virus-infected cells.^{12,13,17} Importantly, this cytolytic response required direct cell-to-cell contact.

Of note, here, we also found that PAM-expanded V γ 9V δ 2-T cells can kill the lung alveolar epithelial cell line, A549. This is not surprising, as A549 cells are immortalized human alveolar epithelial cells and express high levels of MIC A/B, Fas, FasL, DR4 and DR5. Importantly, comparing this to A549 cells without virus infection, PAM-expanded V γ 9V δ 2-T cells had significantly higher cytotoxic activity against influenza virus-infected A549 cells. Indeed, the expression of MIC A/B and DR5 was also upregulated in A549 cells after influenza virus infection. Whether these upregulated molecules, or even some other undefined molecules, contribute to their higher cytotoxic activity against virus-infected cells will need to be determined in further studies.

In conclusion, we have demonstrated for the first time that PAM-expanded human V γ 9V δ 2-T cells can recognize and kill

influenza virus-infected lung alveolar epithelial cells and thus contribute to virus clearance. The cytotoxic activity of PAM-expanded V γ 9V δ 2-T cells is dependent on cell-to-cell contact and requires NKG2D activation. The pathways of perforin–granzyme B, TRAIL and Fas–FasL are also involved in their cytotoxicity. Our study provides insight into the effector functions of V γ 9V δ 2-T cells against influenza A virus in the human body. Targeting these cells by commercially available PAM could potentially offer an alternative treatment for influenza virus infection.

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