The BTLA and PD-1 signaling pathways independently regulate the proliferation and cytotoxicity of human peripheral blood γδ T cells

Hyun J. Hwang1,2 | Jae J. Lee1 | Sung H. Kang3 | Jin K. Suh3 | Eun S. Choi3 | Seongsoo Jang4 | Sang-Hyun Hwang4 | Kyung-Nam Koh3 | Ho J. Im3 | Nayoung Kim1

1Asan Institute for Life Sciences and Department of Convergence Medicine, Asan Medical Center, University of Ulsan College of Medicine, Seoul, Republic of Korea
2Asan Medical Institute for Convergence Science and Technology, Asan Medical Center, Seoul, Republic of Korea
3Department of Pediatrics, Asan Medical Center, University of Ulsan College of Medicine, Seoul, Republic of Korea
4Department of Laboratory Medicine, Asan Medical Center, University of Ulsan College of Medicine, Seoul, Republic of Korea

Correspondence
Nayoung Kim, Asan Institute for Life Sciences and Department of Convergence Medicine, Asan Medical Center, University of Ulsan College of Medicine, Seoul 05505, Republic of Korea.
Email: naykim@amc.seoul.kr

Ho J. Im, Department of Pediatrics, Asan Medical Center, University of Ulsan College of Medicine, Seoul 05505, Republic of Korea.
Email: hojim@amc.seoul.kr

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Abstract
Background: B- and T-lymphocyte attenuator (BTLA) and programmed cell death-1 (PD-1) inhibit γδ T cell homeostasis and activation. This study aimed to determine whether BTLA and PD-1 signaling pathways were convergent or independent in human peripheral blood γδ T cells. Herein we demonstrate that the signalings of BTLA and PD-1 regulated proliferation and cytotoxicity of human γδ T cells, respectively.

Methods: Human peripheral blood γδ T cells were cultured with inactivated Jurkat cells in the presence of interleukin-2 and zoledronate (Zol) for 14 days. Flow cytometry was performed to evaluate the phenotypes and functions of γδ T cells.

Results: The proliferation of the γδ T cells was increased when PBMCs were cocultured with inactivated herpes virus entry mediator (HVEM)low Jurkat cells. The cytotoxicity of the expanded γδ T cells was not affected by coculture with inactivated HVEMlow Jurkat cells and was further increased in the

Abbreviations: AV, annexin V; BTLA, B- and T-lymphocyte attenuator; CFSE, carboxyfluorescein succinimidyl ester; FBS, fetal bovine serum; GVHD, graft versus host disease; HSCT, hematopoietic stem cell transplantation; HVEM, herpes virus entry mediator; IPP, isopentenyl pyrophosphates; ITIM, immunoreceptor tyrosine-based inhibitory motif; ITSM, immunoreceptor tyrosine-based switch motif; mAb, monoclonal antibody; NBP, aminobisphosphonate; PB, peripheral blood; PBMC, peripheral blood mononuclear cells; PBS, phosphate-buffered saline; PD-1, programmed cell death-1; PD-L1, PD-ligand 1; PI, propidium iodide; SHP, src homology region 2 domain-containing phosphatase; WT, wild-type; Zol, zoledronate.

[Correction added on 19 January 2021, after first online publication: Ho J. Im was designated as corresponding author]

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1 | INTRODUCTION

γδ T cells are innate-like immune cells that participate in the early phase of infection and they kill tumor cells directly by degranulation of granzymes and perforin.1 γδ T cells constitute 1%–10% of total T cells. They have unique characteristics that distinguish them from αβ T cells, which make up the majority of T cells.2 Unlike αβ T cells, γδ T cells do not require MHC molecules for antigen recognition.3 Human Vγ9Vδ2 T cells, which are the majority of circulating γδ T cells, require cell-to-cell interactions to induce their reactivity. Vγ9Vδ2 T cells proliferate and are activated by recognizing phospho-antigen directly.1

Through the mevalonate pathway, aminobisphosphonates (NBPs), such as zoledronate (Zol), lead to the accumulation of isopentenyl pyrophosphates (IPP) by inhibiting farnesyl-pyrophosphate synthase. IPP is then recognized by a γδ T cell, which induces it to proliferate.4 The γδ T cells expanded and activated in response to IPP exert antitumor, antimicrobial, and antiviral responses.5–7 It has been reported that repeated and combined administration of NBPs and human Vγ9Vδ2 T cells can effectively control tumor growth in vivo.8 Human Vγ9Vδ2 T cells can directly kill leukemia cells9 without causing graft versus host disease (GVHD).10 It has been proven that the depletion of αβ T cells is a safe and effective tumor therapy.11 Clinical studies are underway to activate γδ T cells by administering Zol after αβ T and B cell-depleted hematopoietic stem cell transplantation (HSCT) to obtain an optimal therapeutic effect based on the activity of γδ T cells.12,13 Thus, γδ T-cell adoptive transfer after HSCT could be an attractive therapeutic option.14,15 It is, however, necessary to optimize methods of efficiently triggering the proliferation and activation of Vγ9Vδ2 T cells.

The presence of anti-PD-L1 mAb. These results suggest that the inactivation of the BTLA signaling pathway during expansion could help produce more γδ T cells without compromising γδ T cell function. The inhibition of BTLA or PD-1 signaling repressed phosphorylation of the src homology region 2-containing protein tyrosine phosphatase 2 and increased the phosphorylation of protein kinase B in γδ T cells. However, there were no synergistic or additive effects by a combination of BTLA and PD-1 blockade. Conclusion: These results suggest that BTLA signaling is crucial in regulating γδ T cell proliferation and function and that the BTLA and PD-1 signaling pathways act independently on the proliferation and cytotoxicity of human peripheral γδ T cells.

**KEYWORDS**
AKT, BTLA, cytotoxicity, human peripheral blood γδ T cells, PD-1, proliferation, SHP2
However, the interaction between the two inhibitory signaling pathways in γδVδ2 T cells is poorly understood. Therefore, we examined whether the BTLa and PD-1 signaling pathways are convergent or independent in γδ T cells by evaluating the proliferation, cytotoxicity, and expression of signaling molecules. The results showed that the BTLa and PD-1 signaling independently regulate the proliferation and activation of γδ T cells, suggesting that inhibition of BTLa signaling could improve the expansion of γδ T cells without compromising cytotoxicity.

2 | MATERIALS AND METHODS

2.1 | PBMC isolation and γδ T cell proliferation

Peripheral blood mononuclear cells (PBMC) or γδ T cell-depleted PBMCs from healthy adult volunteers were isolated by density gradient centrifugation using Ficoll-Paque™ (GE Healthcare). γδ T cell-depleted PBMCs were isolated using CliniMACS (Miltenyi Biotec), as previously described.13 All of the participants provided written informed consent. All of the procedures were approved by the Institutional Review Board, Asan Medical Center, Seoul, Korea (Approval No. 2015-0307). The study was performed ethically, following the Declaration of Helsinki. Isolated cells were immediately frozen in heat-inactivated fetal bovine serum (FBS) (Sigma-Aldrich) containing 10% dimethyl sulfoxide (Sigma-Aldrich) and were maintained in liquid nitrogen until use. The PBMCs were cultured at 6 × 10^5 cells/well in 24-well plates in Rosewell Park Memorial Institute (RPMI) 1640 (Corning) supplemented with 10% heat-inactivated FBS, 100 IU/ml penicillin plus 100 μg/ml streptomycin (Corning), 1 mM sodium pyruvate (Sigma), and 55 μM 2-mercaptoethanol (Gibco) in the presence of 100 U/ml recombinant human interleukin-2 (IL-2) (PeproTech) and 1 μM Zol (Selleckchem). Every 2 days, fresh IL-2 was added.

2.2 | Generation of HVEM\textsuperscript{low} Jurkat cell lines

Jurkat cells (ATCC) were cotransfected with the HVEM CRISPR/Cas9 KO plasmid (Santa Cruz) and the HVEM HDR plasmid (Santa Cruz) according to the manufacturer’s instructions. After 48 h, all transfected cells were selected with media containing 2 μg/ml puromycin (TOCRIS Bioscience) for at least 2 weeks. Before use, the reduction of HVEM expression in the selected cells was confirmed by flow cytometry. The cells expressing lower levels of HVEM were named HVEM\textsuperscript{low} Jurkat cells. Jurkat cell lines were maintained in RPMI 1640 supplemented with 10% inactivated FBS, 100 IU penicillin plus 100 μg/ml streptomycin, and 1 mM sodium pyruvate. All cells were incubated in a humidified CO\textsubscript{2} incubator.

2.3 | Cell proliferation assay

PBMCs were washed and resuspended in phosphate-buffered saline (PBS) (Bioseasang). Then, the cells were labeled with cell proliferation Dye eFluor™ 670 (EF670) (Thermo Fisher Scientific) according to the manufacturer’s instructions.

For inactivation of the Jurkat cells, they were treated with 25 μg/ml mitomycin C (Sigma) for 2 h and then it was washed away three times with 1× PBS. EF670-labeled PBMCs at 3 × 10^5 cells were cultured in 96-well plates with or without the indicated monoclonal antibody and mitomycin C-treated Jurkat cells at 1.5 × 10^5 cells in the presence of 100 U/ml rhIL-2 and 1 μM Zol. Anti-human PD-L1 blocking antibody (clone 29E.2A3; Biolegend) was added at 1 μg/ml at the beginning of the culture. Fresh IL-2 was added every 2 days. In some experiments, BLTA/HVEM blocking peptides (Ac-YRVKEACGELGTGVCPE-NH\textsubscript{2}; Peptron),\textsuperscript{13} and anti-PD-1 mAb (clone EH12.2H7; Biolegend) were used. As control, scrambled peptide (Ac-ELCAPVTRKVECTYGENH\textsubscript{2}; Peptron) and isotype control (clone MOPC-21; Biolegend) were also used. The cultured cells were treated with FcR blocking reagent (Miltenyi Biotec) for 5 min and stained with phycoerythrin (PE)-conjugated anti-human TCRγδ (clone B1.1; Biolegend) and APC-H7-conjugated anti-human CD3 antibody (clone SK7; BD Biosciences) for 30 min. The EF670 dilution was analyzed by FACS Canto II (BD Biosciences) or CytoFLEX (Beckman Coulter Life Sciences) flow cytometers and FlowJo (Tree Star, Inc.) software.

2.4 | Degranulation assay

In the presence of anti-human CD107a (clone eBioH4A3; eBioscience) monoclonal antibody, ex vivo expanded PBMC were incubated with tumor cells at a 1:1 ratio of effector:target. After 1 h, 50 μM monensin (eBioscience) was added. Following overnight incubation and washing, the cells were treated with FcR blocking reagent for 5 min, and stained with fluorescein isothiocyanate (FITC)-conjugated anti-human TCRγδ (clone B1.1; eBioscience) and APC-H7-conjugated anti-human CD3 antibody (clone SK7; BD Biosciences), then analyzed by FACS Canto II or CytoFLEX flow cytometers and FlowJo software (Tree Star).
2.5 | Cell death assay

Cytotoxicity was analyzed using annexin V (AV) and propidium iodide (PI) assays. Target cells were washed and resuspended in PBS. Then, the cells were labeled with carboxyfluorescein succinimidyl ester (CFSE) by using Cell Trace Cell Proliferation Kits (Molecular Probes) according to the manufacturer's instructions. Expanded γδ T cells were resuspended in media and incubated with CFSE-labeled target cells. After 1 day, the cells were washed with AV-binding buffer (BD Biosciences). The cells were treated with FcR blocking reagent for 5 min and stained with anti-AV-APC antibody (BD Biosciences) and PI (Biolegend). Early and late apoptosis was evaluated within the CFSE positive target cells by flow cytometry with FACS Canto II or CytoFLEX. The data were analyzed by FlowJo software.

2.6 | Flow cytometry

In vitro expanded γδ T cells were harvested and stained with the following antibodies: PE-conjugated anti-human TCRγδ (clone MOPC-21; Biolegend), APC-H7-conjugated anti-human CD3 antibody (clone SK7; BD Biosciences), FITC-conjugated anti-human CD279 antibody (clone EH12.2H7; Biolegend), and APC-conjugated anti-human CD272 antibody (clone MIH26; Biolegend). For analysis of the differentiation subsets, the following antibodies were used: FITC-conjugated anti-human TCRγδ (clone B1.1; eBioscience), APC-H7-conjugated anti-human CD3 antibody (clone SK7; BD Biosciences), PE-conjugated anti-human CD45RA antibody (clone HI100; Biolegend), and APC-conjugated anti-human CD27 antibody (clone O323; eBioscience). Differentiation subsets were determined by CD45RA and CD27 expression within the γδ T cells. In some experiments, interferon-γ (IFN-γ) (clone 4S.B3; Biolegend) was intracellularly stained using BD Cytofix/Cytoperm™ (BD Biosciences). Data were acquired by flow cytometry. Flow cytometry was conducted by FACS Canto II or CytoFLEX. The data were analyzed by FlowJo software.

2.7 | Detection of phosphoprotein by flow cytometry

Jurkat cells were preincubated with anti-human PD-L1 blocking antibody (Biolegend) for 30 min in the presence of FcR blocking Ab and then treated with 100 μM pervanadate, an inhibitor of phosphatases, which was used as a phosphorylation inducer. In the presence of IL-2 and Zol, PBMC and tumor cells were coincubated for 20 min and immediately fixed in a 4% paraformaldehyde solution. After washing with PBS containing 2% FBS, the cells were permeabilized with BD™ Phosflow Perm Buffer III (BD Biosciences) for 30 min on ice. Then, they were incubated with anti-pSHP-1 (clone L99-921; BD Biosciences), anti-pAKT (Clone M89-61; BD Biosciences), anti-pERK1/2 (ab212153; Abcam), anti-pSHP-1 (ab192669; Abcam) antibody or IgG1 isotype control for 1 h, following FcR blocking. For pSHP-1 staining, PE-conjugated goat antirabbit IgG Ab (ab724655; Abcam) was treated. Thirty minutes before washing the cells, FITC-conjugated TCRγδ (clone B1.1, eBioscience) and APC-H7-conjugated CD3 antibody (clone SK7; BD Biosciences) were additionally added. Flow cytometry was conducted by CytoFLEX (Beckman Coulter Life Sciences). The data were analyzed by FlowJo (Tree Star, Inc.) software to understand the phosphorylation status of SHP-1/2, ERK1/2, and AKT.

2.8 | Statistical analyses

Growth curves were analyzed by Friedman test, non-parametric repeated-measures analysis of variance using InStat (GraphPad Software). Other results were analyzed using paired two-tailed t tests by GraphPad Prism Ver. 6.0 software (GraphPad Software). When the p value was less than .05, the difference was considered to be significant.

3 | RESULTS

3.1 | BTLA signaling plays a critical role in the proliferation of γδ T cells

We found that the proliferation of γδ T cells in the peripheral blood (PB) from healthy donors induced by IL-2 and Zol was significantly impaired by coincubation with mitomycin C-inactivated wild-type (WT) Jurkat cells, which express HEM (Figures 1A and S1). The differences between the control group, which was treated only with IL-2 and Zol (IL-2 + Zol(−)), and the groups coincubated with WT Jurkat in the absence or presence of anti-PD-L1 were statistically significant at Day 10. To determine whether blocking BTLA and/or PD-1 signaling could improve the proliferation of the γδ T cells, PBMC were cultured with HEMlow or WT Jurkat cells in the presence or absence of anti-PD-L1 signaling. The HEMlow Jurkat cells were produced using a CRISPR/Cas9 system to delete the HEM gene. As expected, HEM expression was downregulated in HEMlow Jurkat cells relative to mock-transfected
Blocking BTLA/HVEM interactions increase γδ T cell proliferation. (A) αβ T cell-depleted PBMCs were cultured with inactivated Jurkat cells with or without anti-human PD-L1 blocking antibodies in the presence of IL-2 and zoledronate(−). The gating strategy is shown in Figure S2. The frequency of γδ T cells (CD3⁺TCRγδ⁺) was determined by flow cytometry. The absolute numbers of γδ T cells (CD3⁺TCRγδ⁺) was calculated at the indicated days. Data were shown as mean ± SEM of six independent experiments (donor no. n = 6, except n = 3 for isotype control group). Friedman test was performed (*p < .05). (B) After EF670-labeling, PBMCs were stimulated as described above. The top plots were only treated with IL-2 and zoledronate(−) (Zol) and the lower plots were cultured with inactivated leukemic cells as indicated on the left. Cells were harvested at 6 or 8 days after culture, and the proliferation was determined by EF670 dilution by flow cytometry. Representative histograms are shown out of three independent experiments. (C) Experiment was performed as above, except that BTLA/HVEM blocking peptides and anti-PD-1 mAb were used instead of HVEMlow Jurkat cells and anti-PD-L1 Ab(−), IL-2- and Zol-treated. BTLA, B- and T-lymphocyte attenuator; HVEM, herpes virus entry mediator; IL-2, interleukin-2; mAb, monoclonal antibody; mlgG1k, isotype control; PBMC, peripheral blood mononuclear cell; PD-1, programmed cell death-1; PD-L1, programmed death-ligand 1
WT Jurkat cells (Figure S1). The expression of PD-L1 was not changed by the deletion of HVEM in Jurkat cells (Figure S1, lower panel).

The inhibited proliferation of the γδ T cells in coculture with WT Jurkat cells was rescued by HVEMlow Jurkat cells (Figure 1A; p = .065). The gating strategy to identify γδ T cells was depicted in Figure S2. HVEMlow HL-60 cells also increased γδ T cell proliferation, compared with WT HL-60 cells (Figure S3A,B). Blocking PD-1/PD-L1 signaling increased the number of γδ T cells only in the presence of HVEMlow Jurkat cells. The treatment with anti-PD-L1 mAb alone failed to increase γδ T cell proliferation when BTLA/HVEM signaling was intact. These results were confirmed by EF670 dilution, as seen in Figure 1B. By Day 8, EF670low/− proliferating cells were 97.5% when treated only with IL-2 and Zol. Deletion of HVEM in Jurkat cells rescued the proliferating cells over 95% at Day 8, compared with 57.3% when cultured with WT Jurkat cells. The averages of the results were summarized in Figure S4. The treatment of BLTA/HVEM blocking peptides during culture increased γδ T cell proliferation which was inhibited by WT Jurkat cells, and anti-PD-1 Ab alone did not increase it (Figure 1C). Taken together, these results suggest that reduction of BLTA/HVEM signaling is a prerequisite for the optimal proliferation of Vδ2 γδ T cells, whereas PD-1/PD-L1 signaling has a limited role in their proliferation.

PD-1 expression on γδ T cells was assessed to determine whether the reduction of BLTA/HVEM or PD-1/PD-L1 signaling during the ex vivo expansion altered PD-1 expression on the γδ T cells. Stimulation with IL-2 and Zol for 10 days significantly upregulated BLTA and PD-1 expression on the γδ T cells, and reduction of BLTA and PD-1 signaling did not change BLTA and PD-1 expression on the γδ T cells (Figure S5).

### 3.2 Blocking BLTA signaling reduces the more differentiated γδ T cell population

It has been suggested that the more differentiated T cells are, the less antitumor efficacy they have,35 and thus the subsets of expanded γδ T cells were analyzed by flow cytometry. As seen in Figure 2, coinoculation with HVEMlow Jurkat cells reduced the proportion of terminally differentiated CD27−CD45RA+ γδ T cells significantly, compared with that incubated with WT Jurkat cells. Coincubation with WT Jurkat cells slightly increased the number of CD27+CD45RA+ naïve T cells, and coinoculation with HVEMlow Jurkat cells in the presence of PD-L1 blockade increased the number of CD27+CD45RA− central memory T cells, although the differences were not statistically significant. These results imply that inhibiting BTLA signaling pathway during expansion could be beneficial for cancer immunotherapy by preventing terminal differentiation of γδ T cells.

### 3.3 The antitumor activity of γδ T cells is not affected by blocking BTLA signaling

To evaluate the effect of a blockade of BTLA/HVEM and PD-1/PD-L1 signaling on the effector function of γδ T cells, PBMCs were treated with anti-human PD-L1 mAb while cocultured with HVEMlow Jurkat or WT Jurkat cells in the presence of IL-2 and Zol. We then examined the antitumor activity of ex vivo expanded γδ T cells using PI and AV in tumor cells. The cells expanded under the conditions shown in Figure 4 were incubated with the same number of CFSE-labeled Jurkat cells for 1 day (Figure 3A). PBMC proliferated by IL-2 and Zol treatment effectively induced apoptosis in all Jurkat cell lines (Figure 3B,C). Likewise, γδ T cells cultured with WT Jurkat cells or HVEMlow Jurkat cells caused apoptosis of the target cells comparable to the PBMC control group. Contrary to expectations, anti-PD-L1 mAb treatment did not increase both the early and late apoptosis of the target cells, despite increased expression of CD107a (Figure 4).

The expanded γδ T cells were challenged by HVEMlow Jurkat or WT Jurkat cells, and then, CD107a expression was measured to detect the degranulation of the ex vivo expanded γδ T cells. Ex vivo expanded γδ T cells in all five groups expressed similar levels of CD107a in response to HVEMlow Jurkat and WT Jurkat cells. Inhibitory signaling by the WT and the HVEMlow Jurkat cells did not affect the degranulation of γδ T. However, the expression of CD107a on γδ T cells cocultured with HVEMlow Jurkat cells was significantly upregulated by the additional inhibition of PD-L1 signaling. Taken together, PD-L1 signaling, rather than HVEM signaling, regulated the degranulation of γδ T cells. The experiments with HVEMlow HL-60 cells also had no significant effect on degranulation (Figure S6). Collectively, the reduced HVEM signaling did not alter the target cell death by γδ T cells during the culture as well as the reaction period.

γδ T cells could produce cytokines, such as IFN-γ, upon stimulation. Thus we assessed intracellular IFN-γ by flow cytometry (Figure 4D,E). γδ T cells were cultured with inactivated WT or HVEMlow Jurkat cells in the presence or absence of anti-PD-L1 Ab for 2 weeks, and the cells were stimulated with WT Jurkat cells in the
FIGURE 2  Blocking the BTLA/HVEM interaction reduces the terminally differentiated γδ T cell subset. (A) PBMC from healthy donors expanded in the presence of IL-2 and zoledronate (Zol) for 14 days. Representative dot plots showing the phenotype of baseline or in vitro expanded cells (left panels) and the CD45RA and CD27 expression of the γδ T (CD3⁺TCRγδ⁺) cells (right panels). Representative data are shown out of three to four independent experiments using distinct donors. (B) PBMC were activated in the indicated culture condition for 14 days. The frequency of differentiation subsets in the in vitro expanded γδ T cells. The following subtypes were used for analysis: T naive (CD45RA⁺CD27⁺), T_CM (CD45RA⁻CD27⁺), T_EM (CD45RA⁻CD27⁻), and T_TMEMRA (CD45RA⁺CD27⁻). Data were shown as mean ± SEM of three to four independent experiments using distinct donors per groups. BTLA, B- and T-lymphocyte attenuator; HVEM, herpes virus entry mediator; IL-2, interleukin-2; PBMC, peripheral blood mononuclear cell; T_CM, central memory T; T_EM, effector memory T; T_TMEMRA, terminally differentiated T. Paired t test was used (**p < .01 and p = .0093).
presence of BTLA/HVEM blocking peptides and/or anti-PD-L1 Ab to measure intracellular IFN-γ. Blocking PD-1 signaling clearly increased IFN-γ production, whereas that of BTLA had only minor effects. In summary, the reduction of BTLA signaling during expansion did not affect the function of γδ T cells.

3.4 | BTLA and PD-1 signaling are independent of each other in γδ T cells

BTLA and PD-1 control cell proliferation and apoptosis through downstream molecules, such as AKT, ERK, and SHP-2, which mediate T cell activating and
Blockade of the BTLA/HVEM signaling does not affect the degranulation of in vitro expanded γδ T cells. (A) Representative histograms showing CD107a expression in the γδ T cells. Gray lines indicate the isotype control. (B) Cells were expanded in the indicated culture condition for 14 days. In the presence of anti-human CD107a mAb, in vitro expanded γδ T cells with the indicated inactivated cells (on top) were incubated overnight with WT Jurkat (upper panel) or HVEM<sup>low</sup> Jurkat (lower panel) cells. Representative data are shown out of three to four independent experiments using distinct donors. (C) The percentages of CD107a<sup>+</sup> γδ T cells in the anti-PD-L1 treated group were normalized by dividing by that of the untreated group. Data were shown as mean ± SEM of three to four independent experiments using distinct donors. Paired t test was used (*p < .05). Each P value is .0190, .0345, and .0103 from left to right. (D) The gating strategy to identify intracellular IFN-γ is depicted. (E) Expanded γδ T cells with the indicated inactivated cells (on top) were incubated overnight with WT Jurkat cells in the presence of BTLA/HVEM blocking peptides and anti-PD-L1 Ab or adequate controls for IFN-γ intracellular staining. (F) The percentages of IFN-γ<sup>+</sup> γδ T cells were summarized. N = 2. BTLA, B- and T-lymphocyte attenuator; HVEM, herpes virus entry mediator; IFN-γ, interferon; mAb, monoclonal antibody; PD-L1, programmed death-ligand 1; WT, wild-type.
inhibitory signalings, respectively. Thus, we next investigated the effects of immediate BTLA/HVEM and PD-1/PD-L1 signaling on the phosphorylation of AKT, ERK, and SHP-2 in resting γδ T cells. PBMCS were incubated with WT Jurkat or HVEM<sup>low</sup> Jurkat cells and/or anti-PD-L1 mAb in the presence of IL-2 and Zol for 20 min, and the expression of the phosphorylated protein was evaluated in γδ T cells by flow cytometry.

FIGURE 5 (See caption on next page)
We observed that the phosphorylation of AKT was increased slightly by blockade of the PD-1/PD-L1 interaction, and a reduction of BTLA/HVEM signaling in addition to it further increased pAKT, which was statistically significant in the percentages as well as MFI (Figures 5C and 5F). Likewise, coincubation with HVEM low Jurkat cells also increased pAKT, compared with that of WT Jurkat cells. The activation of ERK was not changed in the presence of IL-2 and Zol (Figures 5D and 5G). The induction of pSHP-2 was slightly repressed by anti-human PD-L1 treatment in MFI, and inhibition of the BTLA/HVEM interaction reduced the percentage of pSHP-2+ γδ T cells. However, BTLA/HVEM and PD-1/PD-L1 signaling had no synergistic or additive effects on the phosphorylation of SHP-2 (Figures 5E and 5H).

4 | DISCUSSION

Previous research on the individual signals of BTLA and PD-1 in circulating γδ T cells has been conducted, but there are few reports on the interactions of these signals. In this study, the effects of the BTLA/HVEM and PD-1/PD-L1 inhibitory signaling pathways were evaluated on the proliferation and effector functions of human peripheral γδ T cells, in particular Vγ9Vδ2 T cells. We found that BTLA/HVEM blockade could improve the efficient expansion of human peripheral γδ T cells without compromising γδ T cell function. These results could be utilized for the development of γδ T cell therapy for leukemia patients. In addition, the results suggest that the BTLA and PD-1 signaling pathways independently act on the proliferation and cytotoxicity of γδ T cells.

First, we found that culture with inactivated leukemia cells hampered the expansion of human peripheral γδ T cells severely. This may explain the insufficient elimination of tumor cells by γδ T cells in vivo in patients with leukemia, where BTLA on PB γδ T cells can come into contact with HVEM on leukemic cells. Coincubation with WT Jurkat cells slightly increased the proportion of CD27+CD45RA+ naïve T cells, which are known to be associated with GVHD. The proliferation capacity of γδ T cells was upregulated by blocking the HVEM/BTLA interaction without reducing their cytotoxicity, whereas inhibition of PD-1/PD-L1 signaling induced degranulation of γδ T cells. We chose to inhibit the BTLA and PD-1 ligands on the target cells, rather than BTLA and PD-1 directly because we were concerned that the direct blockade of BTLA and PD-1 might unexpectedly affect the proliferation and function of γδ T cells during expansion. However, BTLA also binds to the HCMV UL144 protein to limit lymphocyte-induced inflammation. Potential interactions of BTLA with other ligands might explain the partial inhibition of downstream molecules in this study. In addition, we cannot exclude the possibility that Fc receptors on the γδ T cells might play a stimulating role in response to anti-PD-L1 mAb. Thus, we confirmed the results with BTLA/HVEM blocking peptides and anti-PD-1 mAb and with HVEM low HL-60 cells, which also improved γδ T cell expansion.

In previous reports, both PD-1 and BTLA inhibit the proliferation of T cells. Moreover, BTLA negatively controls the homeostasis of γδ T cells and inhibits phospho-antigen-induced γδ T cell proliferation. Consistent with previous reports, HVEM-deleted Jurkat cells enhanced the proliferation capacity of γδ T cells more than WT Jurkat cells. However, an anti-PD-L1 blockade alone does not have a proliferative effect on γδ T cells. Taken together, the expansion of γδ T cells is tightly regulated by BTLA rather than PD-1.

It should be mentioned that the deletion of the HVEM gene did not affect endogenous PD-L1 expression on Jurkat cells and HL-60 cells and that blocking the BTLA and PD-1 signaling pathways did not affect each other’s expression on γδ T cells either. Although γδ T cells can...
transiently upregulate PD-L1 expression, PD-L1 do not have signaling domains in its short cytoplasmic tail. Thus, it is unlikely blocking PD-L1 on γδ T cells could affect the results in this study. In one study, the expression of BTLA was downregulated by IL-2 and bromohydrin pyrophosphate after 4 days, whereas the expression of PD-1 was upregulated. However, we observed a significant increase in the expression of BTLA and PD-1 in γδ T cells expanded with IL-2 and Zol 10 days. It has previously been reported that the degranulation and cytotoxicity of PD-1+ γδ T cells were reduced in response to PD-L1-expressing target cells, but they were recovered by anti-PD-L1 treatment. Similarly, we observed that inhibition of sustained PD-1 signaling increased CD107a degranulation of γδ T cells. However, it was insufficient to cause enhanced cell death in the target cells.

PD-1 upregulates the expression of SHP-2 in tumor-infiltrating lymphocytes, and inhibition of PD-1 signaling suppresses the phosphorylation of SHP-2. Likewise, pSHP-2 was slightly less induced by inhibition of PD-1 signaling against WT Jurkat cells in our system. In addition, SHP-2 is associated with the phosphorylation of BTLA by TCR engagement. We observed that the specific inhibition of the BTLA/HVEM and PD-1/PD-L1 pathways in γδ T cells reduced phosphorylation of SHP-2 in percentages or MFI, but the combined inhibition did not show any additional effect. Reduction of the BTLA/HVEM or PD-1/PD-L1 signal in γδ T cells inhibited the induction of pSHP-2 in response to Jurkat cells, whereas phosphorylation of AKT was increased by the reduction of both BTLA/HVEM and PD-1/PD-L1 signals. As immune checkpoint receptors, BTLA and PD-1 signaling appear to play independent roles in inhibiting γδ T cell proliferation and cytotoxicity, respectively, and they had no additive effect on each other. In a recent report, BTLA and PD-1 regulate T cell signaling differentially and only partially through SHP-1 and SHP-2, that supports our study. In Terms of activating signaling, dual inhibition of BTLA and PD-1 increased pAKT, compared with that of BTLA single blockade, implying the importance of BTLA signaling in γδ T cells.

In conclusion, BTLA/HVEM inhibitory signaling is a crucial factor in proliferation in ex vivo expanded Vδ2 γδ T cells. On the contrary, PD-1/PD-L1 signaling could provide partial enhancement in proliferation and cytotoxicity. These results indicate the importance of BTLA signaling in γδ T cells and suggest that simultaneous suppression of both signals could effectively improve immunotherapy with γδ T cells. Furthermore, the results of this study could lead to a better understanding of the interactions between BTLA/HVEM and PD-1/PD-L1 signaling in tumor microenvironments.

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CONFLICT OF INTERESTS
The authors declare that there are no conflict of interests.

AUTHOR CONTRIBUTIONS
Ho J. Im and Nayoung Kim conceived, designed, and discussed the research. Eun S. Choi, Sung H. Kang, and Kyung-Nam Koh provided samples and analyzed the results. Hyun J. Hwang, Jae J. Lee, Seongsoo Jang, Sang-Hyun Hwang, and Nayoung Kim conducted the research and analyzed the results. Kyung-Nam Koh, Sang-Hyun Hwang, and Nayoung Kim obtained the grants. Hyun J. Hwang and Nayoung Kim wrote the manuscript.

DATA AVAILABILITY STATEMENT
The data that support the findings of this study are available from the corresponding author upon reasonable request.

ORCID
Nayoung Kim http://orcid.org/0000-0001-8598-0068

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**SUPPORTING INFORMATION**
Additional Supporting Information may be found online in the supporting information tab for this article.

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