Interleukin-18 Synergism with Interleukin-2 in Cytotoxicity and NKG2D Expression of Human Natural Killer Cells

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Abstract

Natural killer (NK) cells play an important role in anti-tumor immunity. Interleukin (IL)-18 is an immunoregulatory cytokine that induces potent NK cell-dependent anti-tumor responses when administrated with other cytokines. In this study, we explored the effects of combining IL-18 and IL-2 on NK cytotoxicity as well as expression levels of the NK cell receptor NKG2D in vitro. Freshly isolated PBMCs were incubated for 48 h with IL-18 and IL-2, then CD107a expression on CD3-CD56+ NK cells was determined by three-colour flow cytometry to evaluate the cytotoxicity of NK cells against human erythroleukemia K562 cells and human colon carcinoma HT29 cells. Flow cytometric analysis was also employed to determine NKG2D expression on NK cells. The combined use of IL-18 and IL-2 significantly increased CD107a expression on NK cells compared with using IL-18 or IL-2 alone, suggesting that the combination of these two cytokines exerted synergistic enhancement of NK cytotoxicity. IL-18 also enhanced NKG2D expression on NK cells when administered with IL-2. In addition, blockade of NKG2D signaling with NKG2D-blocking antibody attenuated the up-regulatory effect of combining IL-18 and IL-2 on NK cytolysis. Our data revealed that IL-18 synergized with IL-2 to dramatically enhance the cytolytic activity of human NK cells in a NKG2D-dependent manner. The results appear encouraging for the use of combined IL-18 and IL-2 in tumor immunotherapy.

Keywords: Interleukin-18 - interleukin-2 - natural killer cells - NKG2D - cytotoxicity
of IL-18 to patients with advanced cancer can generate a relatively limited anti-tumor efficacy but with no apparent toxicities (Robertson et al., 2008; Tarhini et al., 2009). Because of their divergent but complementary properties, the combination of IL-18 and IL-2 is considered as a viable strategy to induce NK cell-mediated anti-tumor responses (Son et al., 2001). However, the exact regulatory mechanism of combining IL-18 and IL-2 on NK cell function is still not well understood.

In the present study, we investigated the effects of combining IL-18 and IL-2 on cytolytic activity and NKG2D expression of human NK cells in vitro. We assayed NK cytotoxicity by multi-parameter flow cytometry using a marker, lysosomal-associated membrane protein-1 (LAMP-1 or CD107a), which can be expressed on the cell surface following degranulation. We also examined the expression of NKG2D on NK cells by using flow cytometry. We found that IL-18 synergized with IL-2 to promote the cytotoxicity and NKG2D expression of NK cells. The synergistic enhancement of NK cytolysis by both cytokines was significantly attenuated by NKG2D blockade. Therefore, we concluded that IL-18 acted synergistically with IL-2 to enhance NK cell activity at least partly via NKG2D pathway.

Materials and Methods

Cytokines and antibodies
Recombinant human IL-18 (rhIL-18) was purchased from R&D systems (Minneapolis, MN, USA) and rhIL-2 was from Shanghai Sangon Biotech Co., Ltd (Shanghai, China). Both cytokines were reconstituted in sterile distilled water and were endotoxin free. PE-conjugated mouse anti-human CD3, APC-conjugated mouse anti-human CD56, and FITC-conjugated mouse anti-human CD107a monoclonal antibodies were products of BD Pharmingen (San Diego, CA, USA). Alexa Fluor® 488-conjugated mouse anti-human NKG2D monoclonal antibody (mAb) and neutralizing antibody against NKG2D (anti-human NKG2D-blocking mAb, mouse IgG1) were purchased from R&D systems (Minneapolis, MN, USA).

PBMCs isolation and cytokine stimulation
Peripheral blood samples were obtained from healthy donors. Peripheral blood mononuclear cells (PBMCs) were isolated from heparinized venous blood in sterile by Ficoll-Histopaque density gradient centrifugation (Tianjin Haoyang, China). These cells were washed two times with RPMI 1640 medium (Hyclone, Logan, UT, USA) and were resuspended at a density of 1×10^6 cells/ml in RPMI 1640 medium containing 10% fetal bovine serum (FBS), 100 U/ml streptomycin and 100 U/ml penicillin. The cells were placed in a 96-well sterile cell culture plate (Costar, Corning, NY, USA) and stimulated with various concentration combinations of rhIL-18 (0, 100, 200 ng/ml) plus rhIL-2 (0, 100, 200 U/ml) for 48 h. Three replicate wells were used. All cultures were maintained in a humidified incubator with 5% CO_2 at 37°C.

Assay of CD107a degranulation
Freshly isolated or cytokine-stimulated PBMCs were cocultured with human erythroleukemia K562 cells or human colon carcinoma HT29 cells in a 96-well cell culture plate with an effector/target (E:T) ratio of 10:1. The final volume was 100 μl in each well. The cells were then stained with FITC-conjugated anti-CD107a mAb and incubated for 5 h in a humidified incubator with 5% CO_2 at 37°C. Monensin (eBioscience, San Diego, CA, USA) was added to each well at a final concentration of 2 μmol/l during the last 4 h of the culture to prevent the degradation of CD107a from NK cell surface and ensure the detectability of this marker after stimulation. After 5 h incubation, the cells were stained with PE-conjugated anti-CD3 mAb and APC-conjugated anti-CD56 mAb. CD107a expression on CD3 CD56^+ NK cells was determined on a FACS Calibur flow cytometer (BD Bioscience, San Jose, CA, USA). To verify the relationship between NK cytolytic enhancement and NKG2D expression on NK cells, untreated or stimulated PBMCs were pre-incubated with NKG2D-blocking mAb or isotype control IgG for 1 h before the coculture with target cells. Then, CD107a degranulation assays were performed.

Flow cytometric analysis of NKG2D expression
After different stimulations, cells were collected, washed and resuspended in 100 μl staining buffer (PBS with 0.5% BSA and 0.1% sodium azide) at 1×10^6 cells/ml. Subsequently, these cells were incubated with PE-conjugated anti-CD3 mAb, APC-conjugated anti-CD56 mAb, and Alexa Fluor® 488-conjugated anti-NKG2D mAb at 4°C in the dark for 30 min, washed twice, resuspended in 300 μl staining buffer directly for flow cytometric analysis. NKG2D expression on CD3 CD56^+ NK cells was examined on a FACS Calibur flow cytometer (BD Bioscience, San Jose, CA, USA).

Statistical analysis
Student’s t test was used to analyze whether there were significant differences in cytotoxicity and NKG2D expression of NK cells between the combined use of cytokines (IL-18 and IL-2) and either cytokine alone (IL-18 or IL-2). A value of p<0.05 was regarded as statistically significant. Statistical analyses were performed by using GraphPad Prism 5 (GraphPad Software, USA).

Results

IL-18 synergized with IL-2 to improve CD107a degranulation of human NK cells in vitro
When cytotoxicity assay was performed against K562 cells, both IL-18 and IL-2 clearly promoted the expression of CD107a on CD3 CD56^+ NK cells. Compared with using IL-18 or IL-2 alone, the combination of IL-18 (200 ng/ml) and IL-2 (100 U/ml) significantly increased the percentage of CD107a positive NK cells (24.89% versus 4.18% and 5.63%, respectively; p<0.01; Figure 1A). Similar results were also observed when using HT29 cells as targets of cytotoxicity test (Figure 1A). Besides, CD107a expression on the surface of NK cells was up-regulated by IL-18 plus IL-2 in a dose-dependent manner (Figure 1B). Although NK cells might have different sensitivity to K562 and
HT29 cells, our results revealed that IL-18 dramatically enhanced the cytolytic activity of NK cells against these two tumor cells when combined with IL-2.

**IL-18 and IL-2 did not affect the proliferation of human NK cells**

Isolated PBMCs were cultured for 48 h in the presence of IL-18 (200 ng/ml) plus IL-2 (100 U/ml), or either cytokine alone. We observed the effect of combining IL-18 and IL-2 on the expansion of NK cells in vitro by flow cytometric analysis of CD3/CD56+ cell population. As shown in Figure 2, we did not observe significant increases in the percentage of CD3/CD56+ NK cells following different cytokine stimulations. The combined use of IL-18 and IL-2 did not affect the proliferation of human NK cells.

IL-18 synergized with IL-2 to increase NKG2D expression on human NK cells

To explore the regulatory mechanism of combining IL-18 and IL-2 on NK cytolysis, we also observed the expression level of NKG2D receptor, which plays a critical role in NK-mediated cytolysis of target cells. Isolated PBMCs were incubated for 48 h with IL-18 (200 ng/ml), IL-2 (100 U/ml) or a combination of both cytokines. Flow cytometric analysis showed that the combined administration of IL-18 and IL-2 significantly enhanced the expression of NKG2D on CD3/CD56+ NK cells in comparison with either IL-18 or IL-2 alone (76.15% versus 59.88% and 65.60%, respectively; p<0.05; Figure 3A and B).

The up-regulatory effect of combining IL-18 and IL-2 on

![Figure 1](image)

**Figure 1. IL-18 Synergized with IL-2 to Increase CD107a Expression on NK Cells.** (A) Representative graphs of CD107a expression on gated CD3/CD56+ NK cells following no stimulation, stimulation with IL-18 (200 ng/ml), IL-2 (100 U/ml) or a combination of both cytokines. NK cytolysis was assayed against K562 and HT29 cells, respectively. Similar results were observed in three independent experiments. (B) The line charts represent the expression pattern of CD107a on NK cells after the culture with different dose combinations of IL-18 and IL-2. Data are given as the mean±SD from three independent experiments

![Figure 2](image)

**Figure 2. IL-18 and IL-2 Did Not Affect the Proliferation of Human NK Cells.** Flow cytometric analysis of the expansion of CD3/CD56+ NK cells following no stimulation, stimulation with IL-18 (200 ng/ml), IL-2 (100 U/ml) or a combination of both cytokines. Representative results of three independent experiments are shown here

![Figure 3](image)

**Figure 3. IL-18 Synergized with IL-2 to Enhance NKG2D Expression on NK Cells.** NKG2D expression levels on NK cells determined by flow cytometry were compared following no stimulation, stimulation with IL-18 (200 ng/ml), IL-2 (100 U/ml) or a combination of both cytokines. (A) Representative graphs of NKG2D expression on gated CD3/CD56+ NK cells. (B) The bar graphs represent the expression pattern of NKG2D on NK cells after different stimulations. Data are given as the mean±SD of three independent experiments (*p<0.05 and **p<0.01)
NK cytotoxicity was NKG2D-dependent

We next blocked NKG2D pathway with anti-NKG2D antibody to investigate the role of NKG2D in the regulation of NK cytotoxicity by IL-18 and IL-2. We observed that the blockade of NKG2D receptor significantly inhibited NK cell-mediated lysis of target cells. The up-regulatory effects of combining IL-18 and IL-2 on NK cytolsis against K562 and HT29 cells were dramatically attenuated by NKG2D blockade (Figure 4). These results suggested that IL-18 synergized with IL-2 to improve the cytolytic activity of NK cells at least partly via NKG2D pathway.

Discussion

NK cells are increasingly considered as potential targets for tumor immunotherapy. Syngeneic IL-12/IL-15/IL-18-pretreated NK cells can proliferate rapidly in tumor tissue and induce sustained anti-tumor effects after adoptive transfer to tumor-bearing mice (Ni et al., 2012). Infusion of continuously growing NK92 cells into patients with lung cancer appears to have some anti-tumor capacities through promoting CIK or CD8+T cell-mediated killing of tumor cells (Son et al., 2003; Du et al., 2012). However, daily injection of IL-18 and IL-2 alone, suggesting that the cytolytic activity of NK cells was dramatically enhanced. Similar results were observed whether K562 or HT29 was used as targets of cytotoxicity assay. However, the combination of IL-18 and IL-2 did not affect the proliferation of NK cells. These data are somewhat inconsistent with a previous report from Son et al. (2001).

NKG2D is a key activating receptor expressed on the surface of almost all NK cells and is extremely important in the recognition and elimination of tumor cells (Shen et al., 2012; Zhao et al., 2014). Recent researches have demonstrated that both IL-12 and IL-15 can augment the cytolytic activity of NK cells by up-regulating the surface expression of NKG2D on NK cells (Zhang et al., 2008; Tang et al., 2013). In contrast, some cytokines, produced in response to certain danger signals, exert a marked inhibitory effect on NKG2D expression of NK cells, thereby reducing NK cytosis against target cells (Muntasell et al., 2010). Besides, IL-18/IL-2 can recover reduced NK cytotoxicity through preventing TGF-β-induced down-regulation of NKG2D (Song et al., 2006).

On the basis of these studies, we hypothesized a possible relationship between NKG2D expression and NK cytolytic enhancement by IL-18 plus IL-2. We found that IL-18 synergized with IL-2 to increase NKG2D expression on NK cells and that blockade of NKG2D with anti-NKG2D mAb dramatically attenuated the up-regulatory effect of combining IL-18 and IL-2 on NK cytolsis. However, we also observed that NK cytolsis against target cells was not completely eliminated by NKG2D blockade. It might be explained by the existence of other NK cell receptors mediating NK cell activity. These results demonstrated that the synergistic enhancement of NK cytotoxicity by IL-18 plus IL-2 was at least partly dependent on the up-regulation of NKG2D expression.

Cytokine therapy has been widely considered as a promising strategy for cancer immunotherapy. For instance, IL-21 and IL-7 could exert potent anti-tumor capacities through promoting CIK or CD8+T cell-mediated killing of tumor cells (Rajbhandary et al., 2013; Yuan et al., 2014). In the present study, we point out that there is a potential model for tumor immunotherapy based on the improvement of NK cell activity by IL-18 plus IL-2. Studies indicate that the combination of IL-18 and IL-2, or a fusion protein IL-18/IL-2 can induce effective anti-tumor responses in several tumor-bearing mice models (Son et al., 2003; Du et al., 2012). However, daily injection of IL-18 and IL-2 also leads to severe pulmonary injury in normal mice (Segawa et al., 2011), suggesting that improper combined use of IL-18 and IL-2 may produce undesired physiological consequences. Therefore, further study is needed to evaluate the efficiency and toxicity of these cytokines in clinical application.

Figure 4. NK Cytolytic Enhancement by IL-18 Plus IL-2 was Dependent on the Up-regulatory Expression of NKG2D. NK cytolsis against K562 (A) and HT29 (B) cells was determined by flow cytometric analysis of the percentage of CD107a positive NK cells after pre-incubation of isolated or stimulated cells with anti-NKG2D mAb. Data represent the mean±SD of three independent experiments. *P<0.01 versus IgG isotype control group.


