Dinaciclib enhances natural killer cell cytotoxicity against acute myelogenous leukemia

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Key Points

• Dinaciclib enhances NK-cell activity against leukemia cells in preclinical AML models.
• Enhanced NK-cell activation by dinaciclib-treated AML is associated with downregulation of inhibitory NK ligand HLA-E on leukemia cells.

Introduction

Although targeting specific oncogenic pathways has improved outcomes for acute myeloid leukemia (AML), in general, survival remains poor. Dinaciclib, a cyclin-dependent kinase (CDK) 1, 2, 5, and 9 inhibitor, has potent anticancer activity against several malignancies and proapoptotic activity against AML.1,2 A major limitation for targeted cancer treatment strategies is treatment resistance, leading many to believe that multimodal combination therapy will be required for the most robust and durable treatment responses. Dinaciclib has been shown to influence the recognition and killing of cancer cells by the immune system, making it an attractive candidate for combined targeted and immune therapy for AML.3

Natural killer (NK) cells can induce a remission in 30% to 50% of patients with chemotherapy-refractory AML.4-6 NK cells respond to target cells through production of proinflammatory cytokines such as interferon-γ (IFN-γ) and tumor necrosis factor-α and the polarized release of cytotoxic granules. The balance of activating and inhibitory signals regulates NK-cell killing. Monoclonal antibodies,7 bispecific or trispecific killer engager molecules,8-10 and hypomethylating agents11 have been used to enhance NK-cell killing of cancer targets, but little is known about the therapeutic potential for combined pathway targeted and NK-cell immunotherapy.

Methods

Cell culture, NK-cell functional assays, and flow cytometry

Primary leukemic cells and healthy donor peripheral blood mononuclear cells (PBMCs) were obtained in accordance with the University of Minnesota Institutional Review Board and the Declaration of Helsinki. Human cell lines were obtained from ATCC (Manassas, VA) and validated by short tandem repeat analysis. NK cells were enriched using the EasySep Human NK-cell enrichment kit (StemCell Technologies, Vancouver, BC, Canada). The NK-cell dose for in vivo experiments was calculated based on the preenrichment PBMC NK-cell percentage. Cell culture, fluorochrome-conjugated antibodies, flow cytometry, and analysis were as previously reported.12 Dinaciclib was obtained from Selleck Chemicals (Houston, TX). AML cells were incubated with dinaciclib and then washed prior to coincubation with effector cells. For the NK-cell–killing assay, targets were cocultured with NK cells at an effector-to-target (E:T) ratio of 2:1 for 24 hours (Figure 1A). Viable target cells were enumerated by flow cytometry using Fixable Aqua Dead Cell Dye to exclude dead cells and using AccuCheck Counting Beads (Thermo Fisher, Waltham, MA). For the NK-cell activation assay, targets were cocultured with effectors at an E:T ratio of 2:1 for 4 hours as previously described.12

Human AML cell line xenograft model

Mouse studies were approved by the University of Minnesota Institutional Animal Care and Use Committee. Luciferase-transduced human HL60 mouse xenografts were established in 6- to 10-week-old NSG mice (The Jackson Laboratory, Bar Harbor, ME) as outlined in Figure 2. Roughly equal numbers of male and female mice were used for all experiments and evenly allocated to experimental groups. Dinaciclib was diluted in 20% hydroxypropyl-β-cyclodextrin and administered intraperitoneally. Recombinant human
interleukin-15 (IL15; R&D Systems, Minneapolis, MN) was administered subcutaneously. HL60 leukemic burden was determined using an IVIS 100 Imaging System (Perkin-Elmer, Waltham, MA). Human blood NK-cell numbers in mouse peripheral blood were determined by collecting 100 μL of peripheral blood by cheek bleed and enumerating human CD45+CD3−CD56+ NK cells during a 60-second collection using flow cytometry.

**Statistical analysis**

Prism software (GraphPad, San Diego, CA) was used for statistical analysis.

**Results and discussion**

We first tested the effects of dinaciclib on NK-cell–mediated cytotoxicity against human AML cells. AML cell lines were cultured in RPMI-1640 with 10% fetal bovine serum containing 1% penicillin/streptomycin and treated with dinaciclib or control vehicle for 24 hours. Cells were then washed to eliminate the drug and then cocultured with primary NK cells at an E:T ratio of 2:1 for 24 hours. Viable AML target cells were enumerated by flow cytometry. Compared with the vehicle control group without NK cells, treatment with 20 nM dinaciclib alone resulted in 74.9% and...
76.1% target killing whereas NK cells combined with 20 nM dinaciclib killed 86.3% and 91.3% of the targets for KG-1 and THP-1 cells, respectively. The NK-killing index (percentage) was determined by comparing the viable leukemic cell number after treatment with drug alone to drug plus NK cells. Treatment with 20 nM dinaciclib increased the NK-killing index compared with vehicle control (KG-1, 45.6% vs 14.5% \( P < .05 \); THP-1, 63.7% vs 31.7% \( P < .01 \)) (Figure 1B). To confirm that increased killing of leukemia targets was due to NK-cell–mediated cytotoxicity, we measured NK-cell activation. Dinaciclib treatment enhanced NK-cell degranulation compared with control treatment as measured by CD107a expression (17.7% vs 8.3%; \( P = .002 \)) and inflammatory cytokine (IFN-γ) production (17.3% vs 0.4%; \( P = .01 \)) (Figure 1C).

To further investigate the translational potential for our findings, we evaluated the ability of dinaciclib to enhance NK-cell killing of primary patient-derived human AML targets. Consistent with the effects on human AML cell lines in vitro, dinaciclib also potently induced NK-cell degranulation in response to primary AML targets in vitro (Figure 1D). Furthermore, in a mouse xenograft model that recapitulates the key features of clinical NK immunotherapy for AML, dinaciclib enhanced the activity of primary human NK cells against human HL-60 AML targets (Figure 2B). Although NK cells failed to reduce the AML burden in untreated leukemic mice, NK-cell treatment in dinaciclib-treated mice significantly reduced the leukemic burden. The reduction in leukemic burden after dinaciclib was associated with a trend toward higher peripheral blood NK-cell numbers (Figure 2C), suggesting that dinaciclib-treated AML cells may promote expansion and/or survival of adoptively transferred NK cells. Together, these findings lend credence to the clinical potential for, and supports further investigation of, this novel combined targeted and immunotherapy approach.

To explore potential mechanisms driving enhanced NK-cell activation against dinaciclib-treated AML targets, we evaluated the effects of dinaciclib on NK-cell ligand expression on leukemia cells. Dinaciclib treatment led to decreased expression of inhibitory ligands (Figure 3A), possibly through its inhibitory effects on CDK9-mediated transcription. Notably, HLA-E expression was markedly decreased on primary patient-derived AML blasts after dinaciclib treatment (Figure 3B). These data suggest that downregulation of HLA-E,
which inhibits the NK-cell function through interaction with NKG2A on NK cells, contributes to enhanced NK-cell cytotoxicity. Therefore, we compared the effects of dinaciclib treatment to NKG2A blockade on NK-cell activation. Leukemic cells were preincubated with an NKG2A-blocking (z199 clone; Beckman Coulter, Brea, CA) or isotype control antibody at 1 μg/mL for 1 hour before coculture with effector cells. The NKG2A-blocking antibody enhanced NK-cell expression of CD107a and IFN-γ after coculture with vehicle-treated leukemic cells but not dinaciclib-treated AML targets (Figure 3C). Furthermore, the NK-cell activation induced by...
Dinaciclib was comparable to the effects of the NKG2A-blocking antibody. Taken together, these findings suggest a key role of dinaciclib-mediated downregulation of HLA-E on leukemia targets in enhancing NK-cell cytotoxicity.

To our knowledge, this is the first report supporting the therapeutic potential for targeting oncogenic signaling to alter the expression of NK ligands on AML cells and thereby sensitize leukemic targets to NK immunotherapy. The use of primary leukemic blasts and NK cells as well as a mouse AML xenograft model that closely mimics the clinical use of adoptive NK-cell therapy for AML patients supports the translational relevance of our findings. By demonstrating that dinaciclib can sensitize AML targets to NK-cell–mediated cytotoxicity, our study introduces a new therapeutic paradigm for AML and supports further investigation into this promising new treatment approach.

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Authorship

Contribution: H.D.Y., J.S.M., and C.E.E. designed the study; H.D.Y. and D.K.S. performed the experiments; H.D.Y., M.F., J.S.M., and C.E.E. analyzed the data; H.D.Y. and C.E.E. wrote the manuscript; and all authors edited and approved the manuscript.

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