Peritoneal resident macrophages in mice with MLL-AF9-induced acute myeloid leukemia show an M2-like phenotype

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Background: Acute myeloid leukemia (AML) is a devastating disease with a poor prognosis. Innate and adaptive immunity is closely related to the progression of leukemia. Macrophages within the leukemic microenvironment have a tendency toward a leukemia-permissive phenotype. However, the characteristics of macrophages in leukemia, including their kinetics, gene expression, and functional roles have not been fully illuminated.

Methods: In the current study, the characteristics of peritoneal resident macrophages, which were large peritoneal macrophages (LPM), from mice with mixed lineage leukemia (MLL)-AF9-induced AML were investigated. AML-associated large macrophages (AML-LPM) were gated as F4/80+ MHC-II− by flow cytometry. To further investigate the relationship between the leukemic microenvironment and macrophage characteristics, RNA sequencing was performed. Meanwhile, apoptosis, killing ability, and phagocytic function of peritoneal resident macrophages in MLL-AF9-induced AML were assessed.

Results: The results suggested that AML microenvironment was found to affect the kinetics and morphology of peritoneal resident macrophages. The results of RNA sequencing suggested that the gene expression of AML-LPMs differed significantly from that of normal LPMs. The AML microenvironment also had effects on the apoptosis, killing ability, and phagocytic function of peritoneal resident macrophages.

Conclusions: These data suggest that peritoneal resident macrophages in mice with AML induced by MLL-AF9 show an M2-like phenotype. The reversal of macrophage polarization in the leukemic microenvironment may potentially enhance the immunotherapeutic effect in AML.

Keywords: Peritoneal resident macrophages; large peritoneal macrophages (LPM); leukemic microenvironment; MLL-AF9-induced acute myeloid leukemia; M2-like phenotype

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Introduction

Macrophages are immune cells that can be found in almost every organ and tissue in the body (1). Macrophages present as different phenotypes and play diverse roles depending on the organ or tissue, and the phenotype and function were regulated by microenvironmental signals (2). Gene expression characteristics analysis of macrophages in a steady state revealed heterogeneity among macrophages in different tissues. Significant differences in gene expression patterns existed among the four classical macrophage populations, which was closely related to heterogeneity in organ-specific microenvironments (3). Macrophages enter into body tissue in their undifferentiated form and differentiate into resident macrophages. Macrophages in different tissues show significant differences in morphology, phenotype, and function, and heterogeneity also exists among macrophages in the same tissue microenvironment (4). Studies suggest that there are two monocyte subsets in the blood, which have distinct functions and might differentiate into different types of macrophages (5).

Different subsets of resident macrophages exist in certain tissues and serosal cavities, such as the lung, spleen, and abdominal cavity (6). In the spleen, at least three types of macrophage subsets can be found: red medulla macrophages, metalloid macrophages, and limbic macrophages. Meanwhile, in the peritoneal cavity, two types of macrophages are present, small peritoneal macrophages (SPM) and large peritoneal macrophages (LPM), both of which are peritoneal resident macrophages (5). These two macrophage subgroups are involved in the fight against pathogens related to intraperitoneal infection, as well as inflammation resist and the maintenance of intraperitoneal homeostasis. The SPM and LPM subgroups differ from other intraperitoneal cells. Both have been found to express the typical molecular markers CD11b and F4/80, but at different levels (5). F4/80 is expressed at much higher levels in LPMs than in SPMs. MHC-II-type molecules can be found at high levels in SPMs but are not expressed in LPMs. SPMs and LPMs are derived from different sources. SPMs are derived from hematopoietic stem cells and are supplemented by circulating monocytes, while LPMs are derived from precursor cells of the yolk sac, have no relation to hematopoietic precursor cells, and can self-renew. SPMs and LPMs show differences in function due to their different origins and phenotypes. In Lavin et al.’s study, neither SPMs nor LPMs could produce a large number of pro-inflammatory or anti-inflammatory cytokines under normal physiological conditions (7). In acute inflammatory states [such as lipopolysaccharide (LPS) abdominal stimulation], most SPMs secrete cytokines such as nitric oxide (NO), tumor necrosis factor (TNF-α), macrophage inflammatory protein 10 (MIP-1α), and regulated upon activation normal T cell expressed and secreted factor (RANTES). LPMs are also able to secrete cytokines such as keratinocyte chemoattractant (KC), granulocyte colony-stimulating factor (G-CSF), and Monocyte chemotactic protein-1 (MCP-1) (5). SPMs and LPMs play different roles in inflammation, with SPMs having more M1-type macrophage properties in inflammatory infection and LPMs being more like M2-type macrophages (7). Recent studies have shown that LPMs can induce B-1 cells to secrete IgA through the GALT-independent pathway, and that LPMs are resupplied to the abdominal cavity during inflammation and participate in the process of inflammatory regression (8). Macrophages play a key role in the host anti-infection process and are able to absorb and degrade microorganisms. Both SPMs and LPMs display phagocytic function, albeit with some differences. Studies have shown that LPMs have significantly higher phagocytic function than SPMs. SPMs and LPMs play an important role in maintenance of intra-abdominal homeostasis, anti-infection/inflammation, phagocytosis, and other processes.

Most macrophages are derived from hematopoietic stem cells; however, some originate from early embryonic stem cells. Educated macrophages in tissue microenvironments are known as tissue-specific macrophages (4). Tissue-specific macrophages show heterogeneity in different organs and tissues, and heterogeneity even exists among macrophages in the same tissue (9). Macrophages in the same tissue are also heterogeneous in different pathologic microenvironments. Phenotypically, educated resident macrophages show great variation, depending on the microenvironment. Phenotypic and functional heterogeneity has reportedly been found in macrophages in the blood system, nerve tissue, connective tissue, and glomerular mesangial tissue; however, little attention has been paid to peritoneal macrophage heterogeneity (7). To date, there have been few reports about the heterogeneity of peritoneal macrophages and no studies on the characteristics of peritoneal resident macrophages in leukemia. Therefore, in the present study, the dynamics and functions of peritoneal macrophages in different physiological subsets and peritoneal macrophage polarization were explored in mice with acute myeloid leukemia (AML). We present the following article in...
accordance with the ARRIVE reporting checklist (available at http://dx.doi.org/10.21037/atm-21-139).

**Methods**

**Mice and AML model**

Male C57BL/6J mice (6 to 8 weeks old) were supplied by the Animal Centre of the Institute of Hematology and Blood Diseases Hospital. All mice were maintained in a certified, specific-pathogen-free animal facility. All experiments involving animals were performed under a project license (No.: EK2020065) granted by the Animal Care and Use Committee of the institution, in compliance with institutional guidelines for the care and use of animals.

**MLL-AF9-induced AML model**

An MLL-AF9-induced mouse AML model was established as previously described (10). Briefly, lineage-negative and Sca-1-positive (Lin-Sca-1+ cells) were obtained from the bone marrow (BM) of normal C57BL/6J mice and transfected with retrovirus carrying the MLL-AF9 fusion gene (MSCV-MLL-AF9-IRES-GFP). Then, the infected cells were transplanted into lethally irradiated C57BL/6J mice. All mice developed AML. In this study, non-irradiated AML mouse model was used. A non-irradiated AML mouse model was established by injecting leukemia cells into the caudal vein of non-irradiated C57BL/6 mice (1×10^6 cells/mouse). All mice developed AML. All mice were sacrificed, and peritoneal washes were performed for further analysis.

**Cell preparation of peritoneal washes**

Peritoneal cells were obtained by peritoneal washing following standard protocol (11). The cells were resuspended in RPMI 1640 Medium (GibcoTM, Invitrogen Corporation, France) for further analysis.

**Flow cytometric analysis and cell sorting**

Cells of different macrophage subpopulations were obtained through cell sorting using a FACSCanto II flow cytometer (BD Biosciences), and then stained with antibodies against mouse F4/80 (APC conjugated, BM8, BioLegend) and against MHC-II (PE-conjugated, AF6-120.1, BioLegend). Peritoneal cells from normal mice were prepared as a negative control. Peritoneal cells were washed and resuspended in perlsucht bacillenemulsion (PBE) containing 0.5% fetal bovine serum (FBS) and were used to fluorescence activated cell sorting (FACS) analysis. Data were analyzed with FlowJo software (version 7.6.1). LPMs were gated as F4/80^high^ MHC-II^+, and SPMs were gated as F4/80^low^ MHC-II^−^.

**Wright-Giemsa staining**

LPMs were spun to slides and standard Wright-Giemsa staining was performed. The morphological characteristics of the LPMs were analyzed using a light microscope (AXIO Observer A1, ZEISS, Germany).

**Cell cycle analysis**

The cell cycles of peritoneal resident macrophages from leukemic and normal mice were analyzed through propidium iodide staining following standard protocol (10). Briefly, cells were fixed with ice-cold 70% ethanol and permeabilized for 11 hours. Meanwhile, RNase was used to disintegrate RNA. Finally, propidium iodide dye was added before analysis on FACS.

**cDNA synthesis and real time polymerase chain reaction**

Cells were lysed, and total RNA was extracted using the RNeasy Mini Kit (Qiagen, Valencia, CA). Reverse transcription was executed using a Super Script First-Strand Synthesis System (Invitrogen, Carlsbad, CA). Real-time polymerase chain reaction (PCR) was carried out on the Step One real-time PCR system (Applied Biosystems, Foster City, CA). The 2^−ΔΔCt^ method [ΔΔCt = (CtTARGET − CtGAPDH)_sample − (CtTARGET − CtGAPDH)_calibrator] was used to compare the expression levels of target genes. All primers of target genes are listed as previously described (11).

**RNA sequencing**

LPMs from the AML microenvironment were defined as AML-LPMs, while those from the normal physiological microenvironment were defined as normal LPMs. SPMs and LPMs were sorted from peritoneal washes on day 15 after the injection of leukemia cells, and they were also obtained from normal mice. RNA sequencing was performed by the Beijing Genomics Institute using
BGISEQ-500 following the standard protocol.

**Statistical analysis**

All results were described as the mean ± standard deviation (SD). Comparisons between two groups were analyzed by unpaired Student's *t*-test, while multiple comparisons were performed using analysis of variance. Statistical differences were considered significant when *P*<0.05.

**Results**

**Acquisition of AML-LPMs**

In this study, an AML model induced by MLL-AF9 was successfully constructed (Figure 1A). The AML onset process is divided into three phases according to time after transplantation: the early phase (7 days after transplantation), metaphase (15 days after transplantation), and late phase (23 days after transplantation) (10). Changes in the levels of GFP+ leukemia cells in the bone marrow and abdominal cavity of the mice during these phases are shown in Figure 1B. We found a high level of leukemia cell invasion in the peritoneal cavity, which might have impacted peritoneal resident macrophages (Figure 1C).

**The AML microenvironment affected the kinetics and morphology of LPMs**

With the progression of leukemia, the number of SPMs in the abdominal cavity firstly increased and then decreased, while the number of LPMs showed the opposite pattern. Neutrophils are also involved in AML progression, and the number of F4/80low neutrophils first increased and then decreased. In the AML model, the number of LPMs in the metaphase was increased (Figure 2A). A morphological comparison with sorted leukemia cells was performed through Wright-Giemsa staining. We found that the vacuolation of LPMs occurred in the middle and late stages of AML onset, and the proportion of LPMs increased with leukemia progression (Figure 2B).

**AML microenvironment affects LPM gene expression**

To further investigate the effects of different microenvironments on macrophages, the gene expression profiles of LPMs from two groups were analyzed. The results suggested that gene expression of LPMs in the leukemic microenvironment differed significantly from that of normal LPMs. Differentially expressed genes (DEGs; with fold change (FC) ≥2, false discovery rate (FDR) ≤0.001) were used to further confirm our observation. A total of 1,532 differentially expressed genes were found in AML, including 907 up-regulated genes and 625 down-regulated genes (Figure 3A,B). To explore the distinctive features of AML-LPMs, the 1532 DEGs were subjected to Kyoto Encyclopedia of Genes and Genomes (KEGG) pathway analysis. Figure 3C shows the top 15 enriched KEGG pathways; of note, these included the cell cycle and p53 signaling pathways. Gene Ontology (GO) analysis of the DEGs in AML was also performed. The top 10 significantly enriched GO entries are listed in Figure 3D. The results of verification by reverse transcription PCR were consistent with those of RNA sequencing (Figure 3E).

**AML microenvironment affects the immunologic function of LPMs**

A nitric oxide fluorescent probe was used to detect the active nitrogen content of LPMs in different microenvironments, comparing the killing ability of macrophages. The active nitrogen content of AML-LPMs was significantly lower than that of normal LPMs, which was different from the gene expression of iNOS (Figure 4A). Macrophages in different microenvironments were used to phagocytize FITC-GFP medium was the main type of AML-LPM, but FITC-GFPlow and FITC-GFPhigh were the main types of normal LPMs (Figure 4B). There were significant differences in phagocytic function between AML-LPMs and normal LPMs. Propidium iodide staining revealed more G2/M/S phase cells among AML-LPMs, showing that AML-LPMs have the potential for proliferation; however, normal LPMs did not have this function (Figure 4C).
Figure 1 Isolation of LPMs (large peritoneal macrophages) in the MLL-AF9-induced AML model. (A) Schematic diagram of the MLL-AF9-induced AML model construction; (B) changes in GFP+ leukemic cells during the pathogenesis of MLL-AF9-induced AML (acute myelogenous leukemia); (C) AML-LPMs were obtained using a flow-sorting technique. ***, P<0.001.
Figure 2 The kinetics and morphology of peritoneal resident macrophages in mice with MLL-AF9-induced AML. (A) The changes in each subgroup of F4/80+ cells in MLL-AF9-induced AML mice; (B) the morphological characteristics of Wright-Giemsa-stained LPMs from different stages of leukemia onset. *, P<0.05; **, P<0.01. MLL, mixed lineage leukemia; AML, acute myeloid leukemia; LPMs, large peritoneal macrophages.

Discussion

The peritoneal cavity selectively attracts and maintains specialized immune cells such as macrophages, which influence the immune microenvironment of the peritoneal cavity (12). There is some phenotypic and functional evidence demonstrating that the two distinct subsets of macrophages show significant heterogeneity in the mouse peritoneal cavity (5). For instance, they differ in size, as determined by flow cytometric light-scatter measurements. Previous research revealed that SPMs and LPMs in the peritoneal cavity of mice can express F4/80 and CD11b, which had a phagocytic function in vivo. However, the present study found there to be significant differences in the expression of several surface markers between SPMs and LPMs, especially in the expression of MHC-II, which is expressed by SPMs but not by LPMs. Significant differences were also found in the expression of F4/80, which is highly expressed by LPMs but not SPMs. Under steady state conditions, LPMs, which originate independently from hematopoietic precursors, maintain their physiological numbers and retain their ability to proliferate in the peritoneal cavity of mice; conversely, under inflammatory conditions, SPMs, which appear to originate from circulating monocytes, increase in number remarkably (13). Yang et al. reported that leukemia cell proliferation led to the aggravation of microenvironment deterioration (14). In the leukemic microenvironment, LPMs could self-renew and maintained the dynamic balance of cell proliferation in the abdominal cavity, and peaked in number during the early stage. At the same time, the leukemic microenvironment affected the morphology of LPMs. Moreover, studies have
shown that macrophage subsets also affect other peritoneal cell populations such as T cells and dendritic cells directly, which may play an important role in the immune state of the peritoneal cavity (5,15).

AML-LPMs and LPMs were respectively analyzed using KEGG pathway analysis, and the results showed that although macrophages that were educated by the leukemic microenvironment shared common characteristics, such as the molecular function of binding, educated macrophages exhibited significant differences, including in changes in cellular composition and biological processes. In most solid tumors, the majority of tumor-associated macrophages (TAMs) are M2-like macrophages. In this paper, we found that AML-LPMs expressed M2-associated genes, thus differing from normal LPMs. As a malignant tumor of the blood system, leukemia has many similarities with solid tumors in pathogenesis but has its own unique characteristics (9,16). As in polarized M2 macrophages,
anti-inflammatory molecules, such as IL-10, arginase-1, and mannose (CD206), were expressed at high levels in AML-LPMs; however, unlike M2-polarized macrophages, they did not significantly express transforming growth factor-β (TGF-β). M2-polarized macrophages have been confirmed to perform an immunosuppressive role in vivo (17). Al-Matary et al. revealed that the number of M2-like TAMs was remarkably increased in the BM of AML patients compared to normal controls (18). Some results have suggested that TAMs perform a crucial role in leukemia progression. Researchers also found that TAMs were polarized to an M2-like phenotype in bone marrow by the leukemic microenvironment, which subsequently played an immunosuppressive role in the BM and spleens of leukemia-bearing mice (17,18). Conversely, bone marrow-derived macrophages (BMDMs) from leukemia sustained the in vitro proliferation of AML cells, which is apparently higher than that of heathy mice (18,19). LPMs also interact with the leukemia microenvironment to accelerate disease progression. A recent study showed
that an increase in M2-like TAMs was associated with poor prognosis in patients with AML (14). Some genes, such as M2-like markers including CD206 and IL-10, which were also studied in pancreatic cancer (20), have been confirmed to be highly expressed in AML-LPMs. Differences in cytokine expression between different pathological microenvironments may be an important factor contributing to the heterogeneity of TAMs (20,21).

Macrophages play an important role in pathogenic bacteria killing and phagocytosis. Nitric oxide secreted by macrophages, is involved in killing and inhibiting exogenous microorganisms. Intracellular NO from macrophages is synthesized through the NOS pathway and regulated by iNOS gene, and its intracellular NO content is closely related to the anti-inflammatory and killing functions of macrophages (22). NO synthesis can be induced by IFN-γ, and this induction process often requires the presence of a second signal (TNF-α or LPS), using IFN-γ or TNF-α alone to induce only low doses of NO with NO direct cytotoxic effects (8,23,24). Studies have shown that in the AML microenvironment, there was significant difference in intracellular NO content between AML-LPMs and normal LPMs, in other words there had some difference in killing function (Figure 4A). These may be caused by immunosuppression, which is closely related to immune reprogramming of immune cells. Immunosuppression is related to the effect of TAMs on other immune cells, such as T cells, through different signaling pathways, which results in the formation of a pro-tumorigenic microenvironment (25). Comparison of macrophage phagocytosis showed that AML-LPM and LPM phagocytic GFP+ particles also showed significant differences, which may be attributable to significant differences in the innate immune-related functions of macrophages in different microenvironments. CD47, which is expressed on macrophages, acts as an inhibitor in phagocytosis (25). Anti-CD47 antibody-based approaches may enhance phagocytosis of leukemia cells by LPMs (26). Comparison of cell cycles of different LPMs between leukemic and normal microenvironment showed cell proportion had significant differences in G0/G1 and G2/M stages, with more LPM proliferation observed in the leukemic microenvironment, which was also consistent with the results of RNA sequencing. The expression level of the pro-apoptotic gene P53 was higher in normal LPMs than in AML-LPMs, while that of the anti-apoptotic BCL2 gene was not significantly different between the two groups.

In conclusion, the heterogeneity of macrophages resident in the same tissue is closely related to different pathological microenvironments (27). Leukemia is similar to malignant solid tumors but has some differences. During the progression of leukemia, changes in the microenvironment of M2-like TAMs may improve their killing and phagocytic innate immune functions (28,29). Changing the polarization state of leukemia-associated macrophages is expected to be a new immunotherapeutic strategy to treat leukemia.

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**Footnote**

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