Role of cluster of differentiation 39 (CD39) in the immunosuppressive status associated Acute myeloid leukemia (AML)

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Abstract

Background: CD39 is an integral membrane protein (ectoenzyme) that phosphohydrolyzes ATP in a Ca²⁺-and Mg²⁺-dependent fashion to yield AMP, Can be viewed as immunological switch, and is expressed in spleen, thymus, lung, and placenta, and in these organs it is associated primarily with endothelial cells and immune cell populations, such as B cells, NK cells, dendritic cells, Langerhans cells, monocytes, macrophages, mesangial cells, neutrophils, and regulatory T cells. Aim of the study is to assess the expression of CD39 on CD4⁺ CD25³high T-reg cells in patients with acute myeloid leukemia to identify its role in immunosuppressive status of AML patients. Methods: Flow cytometric analysis of T-reg cells using CD4 CD25 and CD39 in 60 AML patients and 15 apparently healthy controls with matched age and sex was done. Results: There was a significant difference between cases and controls as regards expression of CD39 on T-helper cells. A highly significant elevation of expression of CD39 has been obtained on T-reg cells in AML patients compared with Controls with increased expression of CD39 on T-reg cells than on helper T cells. Conclusion: CD39 plays a significant role in immunosuppressive status of AML patients particularly with expressing the receptor of α chain of IL-2 (CD25)

Key words: Acute Myeloid Leukemia, CD39, Regulatory T Cells

Introduction:

Acute myeloid leukemia is a complex and heterogeneous hematopoietic tissue neoplasm characterized by rapid expansion of immature myeloid cells in bone marrow and other organs (1). The affected cells undergo an uncontrolled proliferation and impaired differentiation program. Typically, the cells are blocked at various maturation steps and are resistant to cell death (2).

Regulatory T cells are Foxp3⁺CD25⁺CD4⁺ T cells that constitutively express the receptor of α chain of IL-2 (CD25) with diverse immunosuppressive functions. Subsets of T-reg cells include natural T-reg cells (nTregs) that are thymus-derived but undergo further expansion in peripheral tissues, and induced T-reg cells (iTregs) that are converted from conventional T cells (Tcons) in the periphery (3).

Both subsets have been shown to suppress autoreactive lymphocytes and thus to limit the magnitude of innate and adaptive immune responses. Accordingly, impaired T-reg function aggravates autoimmune diseases while T-reg mediated immunosuppression may inhibit pathogen clearance and promote chronic infection. In addition to controlling autoimmunity, T-reg cells have been ascribed to have a role as mediators of cancer-related immunosuppression (4).

T-reg cells suppress effector cells and antigen presenting cells either in a contact-dependent or -independent manner whereas nTregs can use both mechanisms, iTregs induce immunosuppression through cytokines, including IL-4, IL-10, or TGF-β. Cell-to-cell interactions can occur between T-reg cells and T-effs with or without APCs. T-reg-to-Teff contact results in the suppression or apoptosis of T-effector cells. On contact, formation of gap junctions occurs between these 2 T cells. cAMP transferred through the gap from T-reg cells to T-effs suppresses the proliferation of T-effector cells by decreasing IL-2 production and phosphodiesterase 3 inhibitors, which increase cAMP, also increase T-reg cells (5).

In patients with AML, T-reg cells were found to more efficiently hydrolyze adenosine

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triphosphate to adenosine compared with Tregs in healthy control patients. It can be speculated that this might increase T-reg cells cAMP levels in patients with AML and might explain the finding that T-reg cells in AML patients are more suppressive than T-reg cells in control patients. T-reg cells can also suppress IL-2 mRNA in responding T cells. Depleting IL-2 from the microenvironment through its highly expressed CD25 is another mechanism to competitively decrease IL-2 available for T-effs (6).

Current treatments for acute myeloid leukemia (AML) have not been changed for several decades and have not resulted in satisfactory outcomes. Modulating the immune system may improve survival in patients with AML (6). Studies indicate that defects in antileukemic effector cells in patients with AML can contribute to the development and persistence of the disease (7). Some Studies (7) have implicated that immune suppressive regulatory T cells (T-regs) contribute to a defective antileukemic immune response.

CD39 is an integral membrane protein that phosphohydrolyzes ATP, and less efficiently ADP, in a Ca<sup>2+</sup>- and Mg<sup>2+</sup>-dependent fashion, to yield AMP and is expressed in spleen, thymus, lung, and placenta, and in these organs it is associated primarily with endothelial cells and immune cell populations, such as B cells, natural killer (NK) cells, dendritic cells, Langerhans cells, monocytes, macrophages, mesangial cells, neutrophils, and regulatory T cells (8).

CD39 (with CD73) degrade ATP, ADP, and AMP to adenosine. They can be viewed as “immunological switches” that shift ATP-driven pro-inflammatory immune cell activity toward an anti-inflammatory state mediated by adenosine (9).

**Aim of the work:**

The study is designed to assess the expression of CD39 on CD4<sup>+</sup>CD25<sup>high</sup> T-reg cells in patients with acute myeloid leukemia to identify its role in immunosuppressive status of AML patients

**Methods:**

This Case-Control study was conducted at Clinical Pathology, Medical Oncology and Hematology Departments, faculty of medicine, Al-Hussein hospital in the period from November 2016 to January 2018.

The study was carried out on 60 patients with acute myeloid leukemia and 15 apparently healthy controls with matched age and sex.

All patients provided informed consents. The patients were 29 females and 31 males with age range from 17 to 83 years.

**All subjects were subjected**

1. Full history taking and clinical examination.
2. Laboratory investigations which includes:
   - Complete blood count (CBC) including P.B smear.
   - Bone marrow aspiration with examination of Leishman-stained smears laying stress on BM blasts % and blast morphology (for patients only).

**Sample collection**

Two ml of venous blood were obtained from each patient and healthy volunteers by sterile syringes under aseptic conditions to be used for CBC and flow cytometry.

**Methodology**

1. **Complete blood count (CBC)** using Sysmex SF 3000, Roche Diagnostic GmbH, and Mannheim, Germany.

2. **Detection of T-reg cells using CD4 CD25 and CD39 by flow cytometry** using FAC scan.

a) **Principle** In this technique the cells are electronically gated using light scatter parameters (forward scatter reflecting the cell size, and the side scatter reflecting the internal structure of the cell). Antibodies specific for various cellular antigens can be labeled with different fluorochromes that can absorb and emit laser light, allowing simultaneous multicolor flow
cytometric analysis of two or more cell-associated antigens (\textsuperscript{10}).

b) Monoclonal antibodies

The T-reg cells markers were Fluorescein isocyanate (FITC)-conjugated anti-human CD39 \{eBioscience, USA\}, phycoerythrin (PE)-conjugated anti-human CD25 \{eBioscience, USA\} and Peridinin chlorophyll protein complex (PErCP) - Conjugated anti-human CD4.

c) Methods

i) The surface staining was done by adding 10 ul of each monoclonal antibody (anti-CD 4 PerCP, anti –CD 25 PE & anti- CD 39 FITC) to 100 ul of whole anti coagulated blood into the same tube.

ii) The tubes were vortexed and incubated in the dark at room temperature for 30 minutes.

iii) 500 ul of lysing solution was added to the tubes.

iv) The tubes were vortexed and incubated in the dark at room temperature for 10 minutes.

v) The tubes were centrifuged at 1500 rpm for 5 minutes and the supernatant was discarded.

vi) 2 ml of Phosphate buffered saline (PBS) were added to each tube and mixed thoroughly.

vii) The tubes were vortexed and centrifuged at 1500 rpm for 2 minutes and the supernatant was discarded.

viii) Cells were suspended in 500 ul PBS to be ready for analysis.

b) Data acquisition and analysis were performed on software \{cell quest software (BD, San jose, USA)\}.

i) Lymphocytes were gated via their forward and side scatter properties and Treg cells were identified based on their expression of CD4 and CD25.

ii) Positive and negative populations of cells were determined using un-reactive isotype matched monoclonal antibodies as control for background staining.

iii) To discriminate between CD25\textsuperscript{high} T-regs and CD25\textsuperscript{low} activated effectors-memory T cells, we used CD25 expression on non CD4\textsuperscript{+} cells as an internal control. They only express intermediate levels of CD25 (CD25\textsuperscript{low}), whereas CD4\textsuperscript{+} T cells express CD25 with high (CD25\textsuperscript{high}) or intermediate (CD25\textsuperscript{low}) intensities. Only CD4\textsuperscript{+} cells expressing CD25 with higher intensities than the internal control cells were included in the gate for CD25\textsuperscript{high} cells. The gate for CD25\textsuperscript{low} cells was set to include cells expressing CD25 at levels above those of the isotype control but at lower expression levels than the CD25\textsuperscript{high} cells.

iv) Assess the expression of CD39 on CD 4\textsuperscript{+} CD25\textsuperscript{high} T cells.

Statistical methodology:

IBM SPSS statistics (V. 19.0, IBM Corp, USA, 2010) was used for data analysis. Data were expressed as Mean ± standard deviation (SD) for quantitative parametric measures in addition to median for quantitative non-parametric measures and both number and percentage for categorized data.

- The level of significance was stated at 0.05 to be accepted.
- P value > 0.05 was considered non significant result.
- P value < 0.05 was considered significant result.
- P value <0.01 or <0.001 was considered a highly significant result.

Statistical tests used in the study:

1- Student t test: used for Comparison between two independent mean groups for parametric data.
2- **Chi-square test**: to study the association between each 2 variables or comparison between 2 independent groups as regards the categorized data.

3- **Mann-Whitney test**: was done for quantitative variables for not normally distributed and p-value < 0.05 was considered significant.

4- **Pearson correlation (r) test**: to study the possible association between each two variables among each group for non-parametric data and p-value < 0.05 was considered significant.

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**Figure (1):** Histogram (1): Flow cytometric detection of CD39 on T-reg cells and T-helper cells in AML patients showing increased expression of CD39 on T-reg cells (98.2%) than on T-helper cells (5.83%).

**Figure (2):** Histogram (2): Flow cytometric detection of CD39 on T-reg cells and T-helper cells in apparently healthy control showing decreased expression of CD39 on T-reg cells (5.18%) and on T-helper cells (7.3%).

**Results:**

Table (1): Demographic data of AML patients and control group.

<table>
<thead>
<tr>
<th></th>
<th>Cases n=60</th>
<th>Controls n=15</th>
<th>t-test</th>
<th>P value</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Age (years)</strong></td>
<td>17 – 83</td>
<td>18 – 70</td>
<td>0.45</td>
<td>0.65</td>
</tr>
<tr>
<td><strong>X±SD</strong></td>
<td>43.8±14.1</td>
<td>41.9±16.5</td>
<td></td>
<td></td>
</tr>
<tr>
<td><strong>Sex</strong></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Male</td>
<td>31</td>
<td>18</td>
<td>0.0237</td>
<td>0.981</td>
</tr>
<tr>
<td>Female</td>
<td>29</td>
<td>7</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

This table (1) shows:
Role of cluster of differentiation 39 (CD39) in the immunosuppressive status associated Acute…

➢ No significant difference between the two studied groups as regards age.

➢ No significant difference between the two studied groups as regards sex

Table (2): Comparison of CD39*CD4+ (%) (CD39 on T-helper cells) and CD39*CD4*CD25+ (%) (CD39 on T-reg cells) between AML patients and controls.

<table>
<thead>
<tr>
<th></th>
<th>Cases n=60</th>
<th>Controls n=15</th>
<th>M</th>
<th>P-value</th>
</tr>
</thead>
<tbody>
<tr>
<td>CD39CD4 (%)</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Range</td>
<td>2.8-98.02</td>
<td>5.1-26.27</td>
<td>4.38</td>
<td>0.036</td>
</tr>
<tr>
<td>X±SD</td>
<td>25.1±21.5</td>
<td>12.8±5.9</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Median</td>
<td>18.15</td>
<td>12.1</td>
<td></td>
<td></td>
</tr>
<tr>
<td>CD39CD4CD25(%)</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Range</td>
<td>16.14-100.</td>
<td>5-23.53</td>
<td></td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>X±SD</td>
<td>58.7±22.4</td>
<td>10.98±5.3</td>
<td>34.9</td>
<td></td>
</tr>
<tr>
<td>Median</td>
<td>57.69</td>
<td>9.4</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

This table (2) shows: a comparative study revealing a significant difference between cases and control as regards expression of CD39 on T-helper cells. But there is a highly significant elevation of expression of CD39 on T-reg cells in AML patients compared with Controls.

Table (3) Correlation between Blast cells no. in peripheral blood and CD39*CD4+ (%)

<table>
<thead>
<tr>
<th></th>
<th>r</th>
<th>P</th>
<th>Sig</th>
</tr>
</thead>
<tbody>
<tr>
<td>Blasts vs. CD39*CD4+</td>
<td>0.13</td>
<td>&gt;0.05</td>
<td>NS</td>
</tr>
</tbody>
</table>

This table (3) shows Correlation between Blast cells count in peripheral blood and expression of CD39 on T-helper cells and it reveals no significant correlation.

Table (4) Correlation between Blast cells no. in peripheral blood and CD39*CD4*CD25+ (%)

<table>
<thead>
<tr>
<th></th>
<th>r</th>
<th>Sig</th>
</tr>
</thead>
<tbody>
<tr>
<td>Blasts vs. CD39<em>CD4</em>CD25+</td>
<td>-0.06</td>
<td></td>
</tr>
</tbody>
</table>

This table (4) shows Correlation between Blast cells count in peripheral blood and expression of CD39 on T-reg cells and it reveals an inverse correlation not reached to the significant level.

Table (5) Correlation between Blast cells (%) in B.M and CD39*CD4+ (%)

<table>
<thead>
<tr>
<th></th>
<th>r</th>
<th>P</th>
<th>Sig</th>
</tr>
</thead>
<tbody>
<tr>
<td>Blasts vs. CD39*CD4+</td>
<td>0.08</td>
<td>&gt;0.05</td>
<td>NS</td>
</tr>
</tbody>
</table>

This table (5) shows Correlation between Blast cells (%) in Bone marrow and expression of CD39 on T helper cells and it reveals no significant correlation.

Table (6) Correlation between Blast cells (%) in bone marrow and expression of CD39 on Reg T cells and it reveals an inverse correlation not reached to the significant level.

<table>
<thead>
<tr>
<th></th>
<th>R</th>
<th>p</th>
<th>Sig</th>
</tr>
</thead>
<tbody>
<tr>
<td>Blasts vs. CD39<em>CD4</em>CD25+</td>
<td>-0.12</td>
<td>&gt;0.05</td>
<td>NS</td>
</tr>
</tbody>
</table>

Discussion:

Current treatments for acute myeloid leukemia (AML) have not been changed for several decades and have not resulted in satisfactory outcomes. Modulating the immune system may improve survival in patients with AML. T-reg cells have been ascribed to have a role as mediators of cancer-related immunosuppression. Studies indicate that defects in antileukemic effector cells in patients with AML can contribute to the development and persistence of the disease. Studies have implicated that immune suppressive regulatory T cells (T-regs) contribute to a defective antileukemic immune response.

CD39 is an integral membrane protein expressed in spleen, thymus, lung, and placenta, and in these organs it is associated primarily with
endothelial cells and immune cell populations, such as B cells, natural killer (NK) cells, dendritic cells, Langerhans cells, monocytes, macrophages, mesangial cells, neutrophils, and regulatory T cells. This ectoenzyme phosphohydrolyzes ATP, and less efficiently ADP, in a Ca^{2+}- and Mg^{2+}-dependent fashion, to yield AMP (9).

CD39 (with CD73) degrade ATP, ADP, and AMP to adenosine. They can be viewed as “immunological switches” that shift ATP-driven pro-inflammatory immune cell activity toward an anti-inflammatory state mediated by adenosine (9).

In patients with AML, T-reg cells were found to more efficiently hydrolyze adenosine triphosphate to adenosine compared with Tregs in healthy control patients. It can be speculated that this might increase T-reg cells cAMP levels in patients with AML and might explain the finding that T-reg cells in AML patients are more suppressive than T-reg cells in control patients. T-reg cells can also suppress IL-2 mRNA in responding T cells. Depleting IL-2 from the microenvironment through its highly expressed CD25 is another mechanism to competitively decrease IL-2 available for T-effs (6).

The present study was carried out on two groups: the patient group included 60 patients with de novo AML who aged from 17 to 83 years with mean ± SD (43.8±14.1). There were 29 females and 31 males. The control group included 15 apparently healthy individuals aged from 18 to 70 years old with mean ± SD (41.9 ± 16.5). There were 7 females and 8 males.

Regarding a comparative study of CD39 expression on T-helper and T-reg cells in patients suffering from de novo AML& apparently normal controls revealed a significant difference between cases and control as regards expression of CD39 on T- helper cells. However there was a highly significant elevation of expression of CD39 on Reg. T- cells in AML patients compared with Controls with increased expression of CD39 on reg T cells than on helper T cells suggesting role of CD 39 in immunosuppressive status of AML patients.

These results were in accordance with a previous study (11) that revealed a significant difference between AML patients and controls as regards expression of CD39 on Reg. T- cells.

Studying expression of CD39 on T-reg cells in head and neck squamous cell carcinoma (12) the results revealed a significant difference between head and neck squamous cell carcinoma (HNSCC) and controls.

In another Study (13) CD39 was found to be expressed and active on CLL cells. The level of CD39 activity correlates with the stage of the disease. A decrease in CD39 activity was correlated with worsening of disease. These findings suggest that CD39 may be involved in the pathogenesis and prognosis of CLL.

Our study revealed no significant correlation between Blast cells count in peripheral blood and expression of CD39 on T helper cells. On the other hand an inverse correlation existed though non significant level between Blast cells count in peripheral blood and expression of CD39 on T-reg cells, As well no significant correlation existed between Blast cells (%) in Bone marrow and expression of CD39 on T helper cells though an inverse correlation not reached to the significant level existed between Blast cells (%) in bone marrow and expression of CD39 on T-reg cells.

Regarding the age and gender , the results are comparable to that of other studies (14) (15) (2) who demonstrated that median age of the patients was 51, 48 and 42 years; respectively and male to female ratio was 285/215, 70/53 and 48/34; respectively.

In another study (16), the results revealed that distribution of affected males and females is nearly equal, at all ages, The incidence of AML is higher in males than in females with male to female ratio of 1:1:1.0. In contrary to the results (17) which stated that AML is more common in males than females with a ratio of 1.18:0.84 (the same findings announced by American cancer society estimates 2014).

In conclusion, the present study shows that CD39 is an immunosuppressive though not a good prognostic marker in AML. However the prognostic value has to be verified by follow up of cases on a larger scale. AML is more common in males than in females.

References:
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