Abstract. The present study was carried out to assess the percentage of T helper 17 (Th17) lymphocytes in blood and tissue and IL-17A serum concentrations in patients with epithelial ovarian tumors. Two isoforms, IL-17A and IL-17F, as well as IL-21 and IL-22, were simultaneously investigated. The study group consisted of 60 women affected by epithelial ovarian tumors (benign, borderline and malignant) and 20 women without ovarian pathology as a control group. The evaluation of the percentage of Th17 cells secreting IL-17A, IL-17F, IL-21 and IL-22 in peripheral blood and tumor tissues was performed using flow cytometry applying a Th17 cytokine staining panel. The blood serum concentration of IL-17A was determined using ELISA. We found no statistically significant differences in the subpopulations of Th17 lymphocytes, either in peripheral blood or in ovarian tissues, following comparison of the women with and without ovarian pathology as a control group. The evaluation of the percentage of Th17 cells secreting IL-17A, IL-17F, IL-21 and IL-22 in peripheral blood and tumor tissues was performed using flow cytometry applying a Th17 cytokine staining panel. The blood serum concentration of IL-17A was determined using ELISA. We found no statistically significant differences in the subpopulations of Th17 lymphocytes, either in peripheral blood or in ovarian tissues, following comparison of the women with and without ovarian pathology. Negative correlations were found between the percentage of CD4+/IL-17A+F+ (rs=0.8, p=0.02) and CD4+/IL-17+ (rs=-0.78, p=0.03) in the tissue and IL-17A in blood serum in the group of patients with borderline ovarian tumors. A negative correlation was also found between IL-17A and the percentage of CD4+/IL-21+ in peripheral blood (rs=-0.48, p=0.03) in the group of patients with ovarian cancer. The increased percentage of Th17 cells in tissue was not correlated with the overall survival of the ovarian cancer patients. In conclusion, we showed that more Th17 cells secreted IL-17A and IL-21 in the tissue of borderline ovarian tumors and less IL-17A in serum. We also observed that in peripheral blood of the patients with ovarian cancer, there was a higher percentage of Th17 lymphocytes and a lower concentration of IL-17A in serum indicating a negative correlation. An increased percentage of Th17 cells in ovarian tissue does not influence the time of survival of patients with ovarian cancer.

Introduction

Th17 lymphocytes are a helper cell population that play a crucial role in the immune response. The name of these cells originates from the secretion of cytokine IL-17 by these cells (1). They diverge from innate CD4+ T cells in the presence of transforming growth factor-β (TGF-β) and IL-6 (1). IL-17A and IL-17F are the major cytokines secreted by Th17 cells. They are highly homogeneous, bind to the same receptor and have similar biological activities (2). These cytokines are also involved in the development of antibacterial and anti-inflammatory responses (2). In contrast, IL-17F is a weak inducer of the expression of proinflammatory cytokines and it is produced by a wide spectrum of immune cells (2). Th17 lymphocytes also secrete interleukin (IL)-21 and IL-22. IL-21 enhances the expression of perforin, granzyme B, IFN-γ, CXCR3 peripheral NK cells and CD8+ lymphocytes, which reinforces an antitumor response (3). The expression of IL-22 inhibits TGF-β (4,5). The protective role of IL-22 in inflammatory diseases is associated with the induction of the expression of β-defensins and lipocalin-2 (6,7).

In the present study, we assessed the proportion of Th17 cells secreting IL-17A, IL-17F, IL-21 and IL-22 in peripheral blood and in the microenvironment of benign, borderline (BOT) and malignant epithelial ovarian tumors. We examined the relationship between the percentage of CD4+/IL-17A+F+, CD4+/IL-21+ and CD4+/IL-22+ Th17 cells in the peripheral blood and ovarian tissues and the IL-17A serum concentrations. In the present study, we investigated not only the two isoforms of IL-17A and IL-17F, but also IL-21 and IL-22. To
clarify the percentage of Th17 cells we conducted a study to
determine whether Th17 cells and IL-17A may be applied
as prognosticators in patients affected by ovarian carcinomas.

Materials and methods

Research subjects and samples. The study group consisted
of 60 women. The patients were subdivided into 3 groups:
a group of 24 women with malignant epithelial ovarian
tumors (cystadenocarcinoma), 25 women with benign ovarian
tumors (cystadenoma), and 11 women with serous borderline
tumors (BOTs). The control group consisted of 20 women
without ovarian pathology, undergoing surgery due to urinary
incontinence. All women with ovarian cancer had stage
III or IV tumors according to FIGO (International Journal of
Gynecology and Obstetrics, January, 2014). The study samples
included blood serum, peripheral blood, ovarian tissues without
pathology and tissues of malignant and benign tumors of the
ovary. The study was approved by the Bioethics Committee
of the Medical University of Lublin (KE-0254/90/2011). The
patients gave their written consent before they participated
in the present study.

Isolation of mononuclear cells from peripheral blood
(PBMCs). Immediately after blood was taken from the anticu-
bital vein, peripheral blood mononuclear cells (PBMCs) were
isolated by density gradient centrifugation using Gradiisol L
formulation at a specific density of 1.077 g/ml (Aqua Medica,
Łódź, Poland) for 20 min at 700 x g. The pellet containing
the PBMCs was washed twice in phosphate-buffered saline
(PBS) and evaluated for the number (using Neubauer chamber)
and viability (trypan blue staining-0.4% trypan blue solution;
Sigma-Aldrich, Munich, Germany). Viability of <95% was a
disqualifying criteria for conducting further research.

Isolation of mononuclear cells infiltrating tumor and
healthy tissues. During surgery, fragments of ovarian tumor
not containing necrotic areas (the size of 1 cm³) or healthy
ovarian tissue were collected and minced with a scalpel. The
minced tissue was suspended in 30 ml of RPMI-1640 medium
(Biochrom, Holliston, MA, USA) and subjected to digestion
in a mixture containing 1 mg/ml collagenase type IA, 1 mg/
ml DNase type I, 0.1 mg/ml hyaluronidase (all from Sigma-
Aldrich) at 37°C for 60 min, and constantly vortexed. After
digestion, the suspension was filtered through a strainer
(70 µm; BD Biosciences, San Jose, CA, USA) and centrifuged
for 5 min at 700 x g. The cell suspension was washed twice in
RPMI-1640 medium.

Evaluation of the percentage of Th17 cells. The evaluation of
the percentage of Th17 cells (secreting IL-17A, IL-17F, IL-21
and IL-22) in the PBMCs, healthy tissue and tumor tissue was
performed by flow cytometry using a Th17 cytokine staining
panel according to the manufacturer's recommendations using
a FACSCanto (both from BD Biosciences) (Fig. 1).

Establishment of PBMC culture and ovarian tissues (tumor
or control) and stimulation with ionomycin. A 24-h culture of
the PBMCs and ovarian tissue was set-up. The medium was

Figure 1. Example of flow cytometric analysis of Th17 cell subpopulations infiltrating the tumor tissue. Based on the parameters of the FSC, showing the cell
size and SSC, showing the granularity-lymphocytic cells, the gate was set. In the second step of analysis, CD4⁺ lymphocytes were gated, including an analysis
of the expression of IL-17A, IL-17F, IL-21 and IL-22. In addition, the gates of CD4⁺/IL-17A⁺/IL-17F⁺, CD4⁺/IL-17A⁺/IL-17F⁻ and CD4⁺/IL-17A⁻/IL-17F⁺ were
analyzed for coexpression of IL-21 and IL-22.
prepared consisting of 97% RPMI-1640 (Biochrom) supplemented with 2 mM L-glutamine, 2% human albumin (ZLB Bioplasma, Bern, Switzerland), and antibiotics in an amount of 100 U/ml penicillin and 100 mg/ml streptomycin (both from Sigma-Aldrich). Cultivation was carried out in 6-well plates in a 5-ml culture medium. For each patient, two cultures were established, one from the PBMCs and the other from the tumor cells or normal ovarian tissue. The culture was conducted in an incubator under standard conditions (5% CO2, 95% humidity, 37°C) for 4 h. To individual wells, ionomycin at a concentration of 1 μg/ml and PMA at a concentration of 25 ng/ml were added to stimulate cells for the production of cytokines and brefeldin at a concentration 10 μg/ml (both from Sigma-Aldrich) in order to inhibit the activity of the endoplasmic reticulum, leading to retain the cytokines within the cell.

**Determination of intracellular cytokines.** The 24-h cultures were moved from the culture plate to two properly signed tubes and were washed twice in 2 ml of Flow Cytometry Staining Buffer (eBioscience, San Diego, CA, USA) after vortexing. The constant parameters used during each rinsing in this procedure consisted of: run time 5 min, 700 x g. After removal of the supernatant, 5 μl of anti-CD4 (eFluor 450; eBioscience) was added to each tube. The mixture was incubated for 20 min in darkness. After washing, the excess of antibody in 2 ml of Flow Cytometry Staining Buffer, 100 μl IC Fixation Buffer (both from eBioscience) was added to each tube in order to consolidate. Mixing/vortexing, this mixture was also incubated for 20 min in darkness. Then, it was washed twice with 2 ml of the permeabilization buffer (eBioscience). Preparing the buffer involved a 10-fold dilution with PBS. After removing the supernatant, the cells were resuspended in 100 μl of permeabilization buffer and separated to previously prepared cytometry tubes into control and test samples. Thereafter, 5 μl of the antibodies was added to the test samples: anti-IL-17A (FITC), anti-IL-17F (PE), anti-IL-21 (eFluo 660), anti-IL-22 (PerCP-eFluor 710). After vortexing, all samples were incubated for 20 min in darkness. They were washed twice. After the cells were suspended in the Flow Cytometry Staining Buffer, cytometric analysis was carried out. Analysis was performed using flow cytometric 8-channel BD FACSCanto II (BD Biosciences, San Diego, CA, USA). Measurements were performed using BD FACSDiva software.

**Evaluation of the IL-17A cytokine level.** The concentration of IL-17A was determined by ELISA. Quantikine® Human IL-17 ELISA kit (cat. no. D17O0; R&D Systems, Minneapolis, MN, USA) was applied. The procedure was performed according to the manufacturer's instructions, and an automatic Victor3 reader (PerkinElmer, Inc., Waltham, MA, USA) was utilized.

**Statistical analysis.** Non-parametric and Kruskal-Wallis tests were used to verify the differences between the studied groups, post hoc (Dunn) to assess the internal differences between 2 groups, Wilcoxon matched pairs signed-rank for comparing the value of the parameter pairs and Spearman's rank correlation coefficient and its validity to assess the correlation between the two parameters. Kaplan-Meier analysis was used to compare survival curves depending on the range of the percentage of CD4+/IL-17A and concentration of IL-17A. Overall survival was defined as the interval between the date of surgery and the last follow-up or date of death. P-values <0.05 were considered significant. Statistical analysis was performed using 10.0 PL Statistics for Windows (StatSoft, Inc., Tulsa, OK, USA).

**Results**

**Study group.** There was no differences in age (p=0.4), body mass index (BMI) (p=0.053), gravidity (p=0.46) or the concentration of leukocytes in the peripheral blood (p=0.29) of all the study groups (Table I).

**Assessment of Th17 lymphocyte subpopulation in the peripheral blood.** The number of Th17 cells in peripheral blood was decreased in the ovarian cancer patients as compared to the other study groups. We found no significant differences in the percentage of CD4+/IL-17A+/IL-17F (p=0.5) CD4+/IL-17A+/IL-F- (p=0.8), CD4+/IL-17A+/IL-F+ (p=0.32) and CD4+/IL-17A- (p=0.4) Th17 cells in the peripheral blood among the study groups (Table II). CD4+/*IL-17A* is the sum of the percentages of CD4+*/IL-17A*/IL-17F, CD4+*/IL-17A*/IL-17F- and CD4+*/IL-17A*/IL-17F* Th17 cells.

**Percentage of Th17 cells in the ovarian tumor tissue.** The amount of Th17 cells in ovarian tissue was slightly increased in the ovarian cancer patients compared to that noted in the other groups. There were no significant differences in the percentage of CD4+*/IL-17A*/IL-F (p=0.2), CD4+*/IL-17A*/IL-F- (p=0.2), CD4+*/IL-17A*/IL-F+ (p=0.7) and CD4+*/IL-17A* (p=0.5) among the group of patients (Table III). CD4+*/IL-17A* is the sum of the percentages of CD4+*/IL-17A*/IL-17F, CD4+*/IL-17A*/IL-17F- and CD4+*/IL-17A*/IL-17F* Th17 cells.

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**Table I. Demographic and clinical characteristics of the study groups.**

<table>
<thead>
<tr>
<th>Study group</th>
<th>Controls group (n=20)</th>
<th>Benign tumors (n=25)</th>
<th>Borderline tumors (n=11)</th>
<th>Ovarian cancers (n=24)</th>
<th>P-value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age (years)</td>
<td>50 (37-78)</td>
<td>55 (32-85)</td>
<td>47 (35-77)</td>
<td>55 (44-80)</td>
<td>0.400</td>
</tr>
<tr>
<td>BMI (kg/m²)</td>
<td>28 (21-36.7)</td>
<td>26 (11.9-46.7)</td>
<td>31.6 (17.9-45.7)</td>
<td>25.6 (25-35.2)</td>
<td>0.053</td>
</tr>
<tr>
<td>Gravidity (n)</td>
<td>2 (0-4)</td>
<td>2 (0-7)</td>
<td>2 (0-5)</td>
<td>2 (0-6)</td>
<td>0.460</td>
</tr>
<tr>
<td>Leukocytes 10⁷ cells/µl</td>
<td>7.2 (4.17-11.8)</td>
<td>5.8 (3-21)</td>
<td>7.2 (5-10.8)</td>
<td>7.3 (3.58-13.5)</td>
<td>0.290</td>
</tr>
</tbody>
</table>

Data are expressed as median (range). Kruskall-Wallis test was used for statistical analysis. BMI, body mass index.
Percentage of Th17 cells secreting IL-21 and/or IL-22 in the peripheral blood and ovarian tumor tissues. There were no significant differences in the percentage of CD4+/IL-21+ (p=0.5) and CD4+/IL-22+ (p=0.5) Th17 cells in the peripheral blood and ovarian tumor tissues.
blood between the groups of patients (Fig. 2). Moreover, we did not detect significant differences in the percentages of CD4+/IL-21+ (p=0.7) and CD4+/IL-22+ (p=0.8) Th17 cells in the ovarian tissue between the groups of patients (Fig. 3). CD4+/IL-21+ is the sum of the percentages of CD4+/IL-21+/IL-22- and CD4+/IL-21+/IL-22+ Th17 cells. CD4+/IL-22+ is the sum of the percentages of CD4+/IL-21-/IL-22+ and CD4+/IL-21+/IL-22+ Th17 cells.

The percentage of Th17 lymphocytes in peripheral blood and ovarian cancer tissues was next evaluated as a prognostic factor. The Kaplan-Meier survival analysis was performed in a group of patients with ovarian cancer. The patients were subdivided into 2 groups according to the median specificity and sensitivity: >1.6 or <1.6% in the peripheral blood, and >16.3 and <16.3% in ovarian cancer tissue. The result of the Mantel-Cox test was insignificant in regards to patient survival as dependent on the percentage of Th17 cells in the peripheral blood (p=0.19) and in tissue ovarian cancer (p=0.35) (Fig. 4A and B).

Concentration of interleukin-17A (IL-17A). Our results showed statistically significant differences between the analyzed groups (p=0.001) (Fig. 5). Post hoc analysis showed
significantly higher concentrations of IL-17A in women with ovarian cancer compared to the group without ovarian pathology (p=0.00005) as well as that between patients with malignant and benign ovarian tumors (p=0.002).

Relationship between IL-17A and Th17 cell subpopulations secreting CD4+/IL-17A+, CD4+/IL-21+ and CD4+/IL-22+ in peripheral blood and in the tissues of ovarian tumors. A negative correlation was found in the percentage of CD4+/IL-21+* (r=-0.8, p=0.02) and CD4+/IL-17A+* (r=-0.78, p=0.03) in the tissue and IL-17A in blood serum of patients with borderline tumors. Moreover, a negative correlation was shown between IL-17A and the percentage of CD4+/IL-21+* in peripheral blood (r=-0.48, p=0.03) in the group of patients with ovarian cancer (Fig. 6). CD4+/IL-21+ is the sum of the percentages of CD4+/IL-21+/IL-22- and CD4+/IL-21+/IL-22+ Th17 cells. CD4+/IL-17A+ is the sum of the percentages of CD4+/IL-17A+/IL-17F-, CD4+/IL-17A-/IL-17F+ and CD4+/IL-17A+/IL-17F+ Th17 cells.

Prognostic value of IL-17A in ovarian cancer patients. The patients were divided into 2 groups according to the median value of IL-17A (5 women with >0.87 pg/ml and 19 women with <0.87 pg/ml). The result of the Mantel-Cox test was not significant (p=0.6) (Fig. 7).

Discussion

The multiparameter analysis of cytokines secreted by Th17 cells has not been previously performed. Currently we focused on the determination of the capacity of these cells to secrete IL-17 with separation into isoforms. Moreover, we assessed not only two isoforms of IL-17 (A and F), but also IL-21 and IL-22.

The phenotypes, distribution in tissues and profile of the secreted cytokines in ovarian tumors have not yet been fully explored. We found that the distribution of the different subpopulations of Th17 lymphocytes (CD4+/IL-17A+/IL-F, CD4+/IL-17A+/IL-F, CD4+/IL-17A+/IL-F, CD4+/IL-17A+/IL-17F, CD4+/IL-21+ and CD4+/IL-22+) in peripheral blood in non-malignant, borderline (BOT) and malignant tumor tissues were not significantly different in women compared to patients without ovarian pathology. There were no significant differences between
These parameters in terms of Th17 cell proportion, although, a higher concentration of Th17 cells in the tissue was observed, similarly to Kryczek et al (9). The authors demonstrated that the percentage of Th17 cells in the CD4 lymphocytes in tumor tissue was significantly higher compared to peripheral blood. The increased concentration of Th17 in a tissue and their migration to the tumor may be associated with a high expression of CXCR4, CCR6 and CD161 (9). Moreover, it is important to know the functions and interactions between IL-17F and IL-17A in the neoplastic microenvironment at both the molecular and cellular levels. Understanding these relationships may provide a new strategy for systemic anticancer therapy, particularly for ovarian cancer patients.

Kryczek et al (9) also analyzed the relationship between the percentage of Th17 cells and other immune cell subpopulations. They observed an inverse correlation between Th17 and Treg lymphocytes. Tregs have high expression of CD39 (ectonucleotidase), which converts ATP into adenosine. They suggested that the development of Th17 cells is inhibited by Tregs through the adenosinergic pathway. Ye et al (10) noted that the cells from each line of development may be interconverted in the tumor microenvironment by acquiring different functions. Th17 lymphocytes that transform into IFN-γ+FOXP3+ T cells acquire potent immunosuppressive properties. This conversion is possibly part of the tumor escape strategy against cells of the immune system. Fialová et al (11) found an increased recruitment of Th17 cells in women at the early clinical stages of ovarian cancer, while the decreased migration of Tregs was observed in patients affected by advanced disease.

Miyahara et al (12) suggested that a low concentration of Th17 cells, and a high concentration of Tregs is associated with elevated concentrations of TGF-β cytokines in the tumor. They found that tumor cells secrete large amount of the latent form of TGF-β (inactive). However, the level of the active form of TGF was very low due to its short half-life. In addition, TGF-β may be present in the free form and in the form of membrane-bound Tregs. Increased percentage of Tregs and reduced Th17 cells in the tumor supported the view that these cells mutually regulate the presence of other cytokines in the tumor microenvironment (8,11). However, the molecular mechanisms underlying the formation and mutual regulation of Treg and Th17 cells in the tumor microenvironment remains unknown.

There is increasing evidence suggesting that Th17 cells protect against cancer development in several manners. Firstly, tumor-infiltrating Th17 cells express several effector cytokines, similar to those detected in patients affected by infectious diseases. This suggests that Th17 cells infiltrating the tumor may be functionally similar to T effector cells. In accordance with this possibility, Th17 cells are negatively correlated with the presence of Tregs and are positively correlated with effector cells secreting IFN-γ, cytotoxic CD8+ and NK cells, in the same tumor microenvironment. Secondly, Muranski et al reported a protective role of Th17 cells during carcinogenesis (13). Transgenic T cells with a Th17 phenotype after treatment with TGF-β and IL-6 induced tumor eradication in mice. Furthermore, mice deficient in IL-17 showed accelerated tumor growth and lung metastases in several cancer models. Tumor-infiltrating Th17 lymphocytes do not produce granzyme B and perforin; therefore, Th17 cells do not act as intermediaries in direct cytotoxic activity against tumor cells (3). Instead, Th17 cells recruit other effector cells of the immune system. According to this hypothesis, IL-17 and INF-γ derived from Th17 cells synergistically induce the production of CXCL9 and the chemokine CXCL10 by tumor cells, which in turn promotes the migration of effector T lymphocytes into the tumor microenvironment (3). CXCL9 and CXCL10 levels were found to be directly correlated with the number of CD8+ tumor-infiltrating and NK cells. Th17 lymphocytes stimulate tumor cells to secrete CCL20, which recruits dendritic cells into the tumor microenvironment. These data strongly support the view that Th17 cells play an indirect role in antitumor immunity through the promotion of effector T cells, NK and dendritic cells (7).

There is strong evidence demonstrating the role of IL-17 secreted by Th17 cells in the promotion of carcinogenesis. IL-17 induces IL-6 production by tumor cells and stromal cells. IL-6 activates STAT3, which increases the level of genes facilitating tumor progression and the development of metastasis (7). Tartour et al (14) reported that transfection with IL-17 of human cervical cancer cells increased tumor growth when transplanted into nude mice. In addition, mice lacking IL-17 demonstrated reduced tumor growth of B16 melanoma and MB49 bladder cancer, suggesting a role of IL-17 in promoting tumor growth (14).

Kryczek et al (9) also demonstrated a correlation between the percentage of Th17 cells and patient survival time. In patients whose peritoneal fluid contained high concentrations of IL-17, the average survival time was 78 months. In turn, patients with decreased levels of IL-17 in peritoneal fluid lived significantly shorter (27 months). Opposite data were published by Lan et al (15) who found that high IL-17 expression was correlated with improved progression-free survival in advanced stage ovarian cancer patients. No significant difference was observed in overall survival between the high and low IL-17 expression groups (15). In the present study, the decreased percentage of Th17 cells (CD4+/IL-17+) in the tumor microenvironment did not correlate with a reduced survival time of patients affected by ovarian cancer, probably due to advanced stage of the ovarian cancer group.

The next step of the research was to assess the proportion of Th17 lymphocytes secreting IL-21 and IL-22. The percentages of CD4+/IL-17+/IL-21+, CD4+/IL-21+/IL-22+, CD4+/IL-21+/IL-22+, CD4+/IL-21+ and CD4+/IL-22+ Th17
cells in peripheral blood and tissue did not significantly differ among the groups. Moreover, a significant correlation was found between CD4+/IL-21+ in peripheral blood and IL-17A in the group of women with ovarian cancer. IL-21 is produced predominantly by Th17 and NKT cells. In cooperation with TGF-β it induces the differentiation of T cells towards the Th17 phenotype. It also participates in the mutual regulation of Th17 and Treg cells. In certain circumstances, IL-21 may exert anti-inflammatory effects due to its ability to inhibit dendritic cell maturation and stimulation of IL-10 (15). In this context, IL-21 stimulates an immune response against tumor cells and promotes a CD8+ T cell response against viruses (16,17). Immunosstimulatory activity of IL-21 may be possibly applied as a potential immunotherapeutic agent for the treatment of human cancers. Neutralization of IL-22 may reduce metastasis, chemotherapy, and inflammation associated with cancer. Given that IL-2 BP is a specific natural antagonist of IL-22, it may be a prime candidate as an anti-IL-22 therapy. Anti-TNFα drugs such as adalimumab, etanercept and infliximab transiently decrease IL-22 expression, likely since Th22 cells depend on TNF for differentiation. The anti-IL-6 antibody tocilizumab may also suppress the differentiation of both Th1 and Th22 cells. A neutralizing antibody against IL-12p40, ustekinumab, is able to target both IL-12 and IL-23 and therefore prevent the differentiation of Th1, Th17 and Th22 cells, eliminating sources of IL-22 (18-20). However, the nature and clinical relevance of IL-22+ cells is poorly defined in patients affected by ovarian cancer.

Xiang et al (21) reported that IL-17 contributed to ovarian cancer malignancy by promoting the self-renewal of CD133+ cancer stem cells and IL-17. They showed by recombinant human IL-17 stimulation and IL-17 transfection that the growth and sphere formation capacities of CD133+ ovarian cancer stem cells were significantly enhanced in a dose-dependent manner increasing the tumorigenesis capacity in nude mice. These data suggest that the IL-17 signaling pathway may serve as a therapeutic target for patients with ovarian cancer. Kryczek et al (22) also revealed a relationship between a key transcription factor (STAT3) and an important epigenetic marker (H3K79) in determining cancer stemness and tumorigenesis.

Our previous data revealed that the percentage of Th17 cells in peritoneal fluid (PF) corresponds with the severity of endometriosis (23). The percentage of Th17 lymphocytes in PF was significantly higher in patients with moderate/severe endometriosis compared to patients with a minimal/mild form of the disease (23). Continuing with our research we demonstrated a higher percentage of Th17 cells in tissue than in peripheral blood and a negative correlation between CD4+/IL-17+ lymphocytes and the concentration of IL-17A in patients with ovarian cancer, thus that IL-21 could adjust Th17 differentiation. The generation of Th17 cells in the conventional manner is attenuated by blocking IL-21. This cytokine is capable of acting on Th17 cells in an autocrine manner in response to antigen stimulation (24).

In conclusion, we showed that more Th17 cells secreted IL-17A and IL-21 in the tissues of borderline ovarian tumors and less IL-17A in serum. We also observed that in peripheral blood of patients with ovarian cancer, a higher percentage of Th17 lymphocytes was negatively correlated with a lower concentration of IL-17A in serum. An increased percentage of Th17 cells in ovarian tissue did not influence the survival time of patients with ovarian cancer.

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References


