

Role of T helper 17 cytokines in the tumour immune inflammation response of patients with laryngeal squamous cell carcinoma

WEI-JING TANG¹, LEI TAO¹, LI-MING LU², DI TANG¹ and XIAO-LIN SHI¹

¹Department of Otolaryngology, Eye, Ear, Nose and Throat Hospital, Fudan University, Shanghai 200031;

²Shanghai Institute of Immunology, Shanghai Jiaotong University School of Medicine, Shanghai 200025, P.R. China

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Abstract. Altered cytokine production can lead to immune dysfunction in patients with cancer. The present study investigated the expression of T helper (Th)17 cytokines in patients with laryngeal squamous cell carcinoma (LSCC) and their clinical significance in providing new therapeutic insights. The prevalence of Th17 cells and their receptors in patients with LSCC was studied using immunohistochemical analysis via tissue microarray technology. Flow cytometry was used to investigate the percentage of Th17 and Th1 cells in peripheral blood mononuclear cells. Furthermore, the proliferation of Th17 cells and Th17-associated cytokines, including interleukin (IL)17, IL23 and RAR-related orphan receptor γ t, was analyzed by reverse transcription-quantitative polymerase chain reaction. The results revealed that the prevalence of Th17 cells in patients with LSCC was elevated in their primary tumors, as well as in peripheral blood, compared with that in healthy controls. It was further demonstrated that Th17 cells could be induced and expanded in the tumor microenvironment through cytokines produced by the tumor cells. In conclusion, Th17 cells have a substantial impact on the carcinogenesis of LSCCs, and could serve as a potential therapeutic target to modulate the anti-tumor response in these carcinomas.

Introduction

Laryngeal carcinoma is the 11th most common form of cancer in humans worldwide (1). Pathologically, >95% of all laryngeal malignancies are squamous cell carcinomas (SCCs) (2). It has been reported that >75% of cases of laryngeal SCC (LSCC)

are attributable to cigarette smoking and alcohol consumption. Cigarette smoking is associated with an increased risk of LSCC compared with that of individuals who have never smoked, and heavy alcohol intake is an independent risk factor of LSCC (3). LSCC remains difficult to treat; furthermore, treatment can cause severe long-term side effects. For patients who are not cured by surgery and/or chemo-radiotherapy, there are few effective treatment options (4). Thus, targeted therapies and predictive biomarkers are urgently required in the perspective of LSCCs to improve the management and minimize the treatment toxicity, as well as to allow the selection of patients who are likely to benefit from both non-selective and targeted therapies (5).

Previous studies have confirmed that tumor immune inflammation is important in the tumor microenvironment (6). Cluster of differentiation (CD) 4⁺ T cells are essential organizers of cell-mediated immunity, participating in each stage of the immune response. It has been confirmed that naïve (uncommitted) CD4⁺ T cells can be induced to different specific lineages according to the local cytokine, including towards T helper (Th)1, Th2, Th17 and T regulatory (Treg) cells (7). Interleukin (IL)17-producing Th17 cells, which are different from Th1 and Th2 cells, have been described as serving critical roles in inflammation and autoimmune diseases, as well as in cancer development (8-11). Considering these facts, it could be concluded that pro-inflammatory Th17 cells may have extensive effects on LSCC pathogenesis and anti-tumor response.

The present study revealed that patients with LSCC have elevated levels of Th17 cells in their primary tumors and peripheral blood compared with those in healthy controls. In addition, the LSCC microenvironment was identified as a strong Th17-cell inducer.

Materials and methods

Study subjects. A total of 70 tumors and 70 adjacent control tissues (pathologically confirmed normal mucosa) were obtained from patients with LSCC undergoing surgery. Peripheral blood was obtained from another 36 patients with LSCC and from 16 healthy individuals, who served as controls. The patients were registered for treatment at the Eye, Ear, Nose and Throat Hospital of Fudan University (Shanghai, China) from September 2013 to January 2015. For LSCC samples, patients with a known history of LSCC were enrolled, and malignancy was assessed by a pathological examination of the biopsies. The tumor staging of

Correspondence to: Professor Lei Tao, Department of Otolaryngology, Eye, Ear, Nose and Throat Hospital, Fudan University, 83 Fenyang Road, Shanghai 200031, P.R. China
E-mail: doctortaolei@163.com

Professor Li-Ming Lu, Shanghai Institute of Immunology, Shanghai Jiaotong University School of Medicine, 280 South Chongqing Road, Shanghai 200025, P.R. China
E-mail: lulunew2003@163.com

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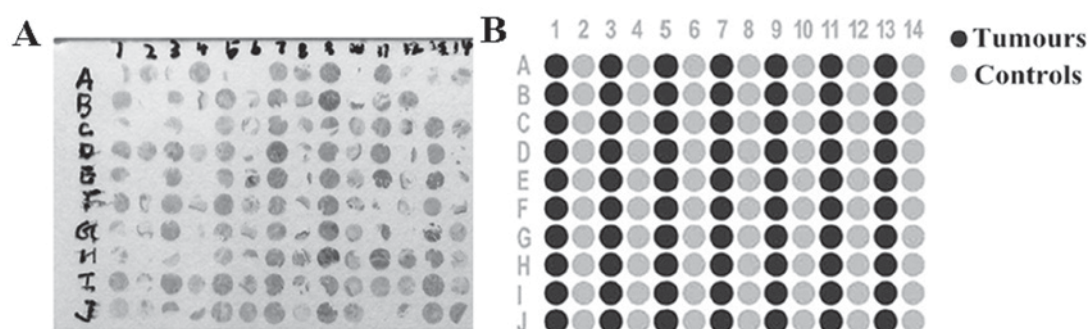


Figure 1. Multitumor TMA. (A) Section containing 70 tumors and 70 adjacent tissues samples. The diameter of each tissue spot is 0.6 mm. (B) Schematic layout of the TMA section. TMA, tissue microarray.

the patients was performed in accordance with the American Joint Committee on Cancer tumor-node metastasis (TNM) classification (12). All donors participated on a voluntary basis and provided written informed consent. The protocol was approved (no. KJ2008-01) by the ethics committee of the Eye, Ear, Nose and Throat Hospital of Fudan University (Shanghai, China) and was in agreement with this institutions ethical guidelines.

Tissue microarray (TMA) construction. TMA technology is a method where a large number of tissue samples are placed on a microscopic glass slide, which facilitates the transition from basic research to clinical applications (13). Briefly, the most representative tumor and adjacent tissue samples were selected in pairs. The corresponding areas of each donor paraffin blocks were perforated using a trephine needle (Quick Ray[®]; Unitma Co., Ltd., Seoul, Korea) with a size of 1.5 mm. The 1.5-mm-sized tissue cores were transferred and embedded into the recipient block with 140 empty 1.5-mm-sized holes. All study specimens were obtained with both tumor and adjacent tissue samples from each donor block. Multiple 4- μ m-thick sections were cut with a microtome and transferred to poly-L-lysine-coated slides (#22247-1; Hannotech Co., Ltd., Dongguan, China). TMA blocks were constructed, each containing one sample from all tumors and adjacent tissues (Fig. 1).

Immunohistochemical staining. Two TMA blocks with 70 tumors and 70 adjacent tissues samples were used for immunostaining, followed by standard procedures for the avidin-biotin-peroxidase method (14). The following anti-human monoclonal antibodies were used: Rabbit anti-IL17 (#bs-2140R; BIOSS, Beijing, China) and rabbit anti-IL17A receptor (IL17R; #bs-2606R; BIOSS). Briefly, the color reaction was developed with a 3,3'-diaminobenzidine solution at 20°C for 5 min, and antibodies were diluted to 1:200 and incubated at 37°C for 1 h. The cells were counterstained with hematoxylin.

The slices were evaluated by two pathologists without knowledge of the clinical outcome. The percentage of immunoreactive cells was graded on a scale of 0 to 4, as follows: No staining was scored as 0; 1-10% of stained cells was scored as 1; 11-50% of stained cells was scored as 2; 51-80% of stained cells was scored as 3; and 81-100% of stained cells was scored as 4. The staining intensities were graded from 0 to 3, as follows: 0 was defined as negative; 1 as weak; 2 as moderate; and 3 as strong. The raw data were converted into an

immunohistochemical score (IHS) by multiplying the quantity and intensity scores. An IHS score of 9-12 was considered to represent strong immunoreactivity (+++); 5-8 was considered as moderate (++); 1-4 was considered as weak (+); and 0 was considered as negative immunoreactivity (-). On the final analysis, the cases that had an IHS <1 were considered as negative, and those with an IHS \geq 1 were regarded as positive.

Flow cytometry. Peripheral blood mononuclear cells (PBMCs) were isolated from 5 ml of freshly obtained peripheral blood by centrifugation (800 x g, 20°C, 20 min) on a Ficoll Hypaque density gradient (Ficoll PM 400; Sigma-Aldrich; Merck KGaA, Darmstadt, Germany). Prior to intracellular staining, the isolated PBMCs were stimulated for 5 h with 2 μ l/ml Cell Stimulation Cocktail (#00-4970; eBioscience, Inc., San Diego, CA, USA), a cocktail of phorbol 12-myristate 13-acetate (PMA; eBioscience, Inc.) and ionomycin (eBioscience, Inc.) in the presence of Protein Transport Inhibitor Cocktail (#560751; BD Biosciences, Franklin Lakes, NJ, USA). Briefly, cells were fixed and permeabilised using the BD Cytofix/Cytoperm[™] Buffer (Fixation and Permeabilization Solution; #560751; BD Biosciences) according to manufacturer's protocol. Subsequently, the isolated PBMCs were intracellularly stained with the Human Th1/Th2/Th17 Phenotyping Cocktail (560751; BD Biosciences). Flow cytometry was performed on a BD FACSCalibur (BD Biosciences) and the data were evaluated using FlowJo software version 7.6.1 (TreeStar, Inc., Ashland, OR, USA). To determine the percentage of Th17 and Th1 cells, lymphocytes were gated by plotting forward vs. side scatter followed by gating on CD4⁺ T cells. The gated cells were then analyzed for IL-17A as phycoerythrin and interferon (IFN)- γ as fluorescein isothiocyanate expression.

Reverse transcription-quantitative polymerase chain reaction (RT-qPCR). Total RNA was extracted from patients' frozen tissues using TRIzol reagent (15596-018; Thermo Fisher Scientific, Inc., Waltham, MA, USA) according to the manufacturer's protocol. Complementary DNA (cDNA) was synthesized from 1 μ g total RNA in a 20- μ l reaction system using PrimeScript RT Reagent kit (Perfect Real Time; DRR063A; Takara, Bio, Inc., Otsu, Japan). The cDNA was then diluted with sterile water and stored at -20°C. The RT procedure using the PrimeScript RT Reagent kit was performed according to the manufacturer's protocol.

Table I. Characteristic features of the patients in the current study.

Characteristics	Patients (fresh tissue) (n=70), n (%)	Patients (blood) (n=36), n (%)	Controls (blood) (n=16), n (%)
Age, years			
Mean (range)	60.63 (38-84)	59.10 (43-73)	59.50 (52-72)
Sex			
Male	69 (98.57)	36 (100.00)	16 (100.00)
Female	1 (1.43)	0 (0.00)	0 (0.00)
Site			
Supraglottic	23 (32.86)	6 (16.67)	
Glottic	40 (57.14)	26 (72.22)	
Infraglottic	7 (10.00)	4 (11.11)	
cT stage			
T ₁ +T ₂	17 (24.29)	25 (69.44)	
T ₃ +T ₄	53 (75.71)	11 (30.56)	
pN stage			
N ₀	33 (47.14)	31 (86.11)	
N ₁ +N ₂	37 (52.86)	5 (13.89)	
Clinical stage			
Early stage (I+II)	6 (8.57)	26 (72.22)	
Late stage (III+IV)	64 (91.43)	10 (27.78)	
Smoking history			
Smokers	62 (88.57)	25 (69.44)	
Non-smokers	8 (11.43)	11 (30.56)	
Alcohol consumption			
Drinkers	34 (48.57)	17 (47.22)	
Non-drinkers	36 (51.43)	19 (52.78)	

RT-qPCR for IL17, IL23 and RAR-related orphan receptor (ROR) γ t was performed on an Applied Biosystems 7500 Fast Real-Time PCR System (Thermo Fisher Scientific, Inc.), and the data were analyzed using the Applied Biosciences Real-Tiem PCR system 7500 software version 2.0.6 (Applied Biosystems; Thermo Fisher Scientific, Inc.). In brief, 2 μ l cDNA was added to a 20- μ l reaction mixture containing 10 μ l 2X SYBR Premix Ex Taq, 0.4 μ l forward primer (10 μ M), 0.4 μ l reverse primer (10 μ M), 0.4 μ l ROX reference dye and 6.8 μ l sterile water (all #DRR063A; Takara, Bio, Inc., Otsu, Japan). All primers were designed using the Primer Premier version 5.0 software (Premier Biosoft International, Palo Alto, CA, USA), with their specificity confirmed by the Basic Local Alignment Search Tool on the National Center for Biotechnology Information website (<http://blast.ncbi.nlm.nih.gov/Blast.cgi>). Detailed information on the aforementioned primers is as follows: IL-17, forward 5'-TCCCACGAAATCCAGGATGC-3' and reverse 5'-GGATGTTTCTCAGGTTGACCATCAC-3'; IL-23, forward 5'-TGCTCCCTGATAGCCCTGTGG-3' and reverse 5'-GCTGGGACTGAGGCTTGG AAT-3'; and ROR γ t, forward 5'-GTGGGGACAAGTCGTCTGG-3' and reverse 5'-AGTGCTGGCATCGGTTTCG-3'. The PCR conditions were: 95°C for 30 sec, followed by 40 cycles at 95°C for 5 sec and 60°C for 25 sec. All samples were processed in triplicate. The experiment was repeated for three times.

Relative gene expression was calculated using the comparative quantification cycle (Cq) method (15). The messenger RNA expression levels of the target genes were normalized by β -actin, and the $2^{-\Delta Cq}$ values were represented. For data analysis, the $2^{-\Delta Cq}$ method was used to calculate the fold-change in expression, in which ΔCq represented the difference between the Cq value of the target gene and that of β -actin ($Cq_{\text{target gene}} - Cq_{\beta\text{-actin}}$).

Statistical analysis. The data were reported as the mean \pm standard deviation or the mean \pm standard error. IL-17 and IL-17R expression in tumors and adjacent tissues was evaluated by χ^2 test. The percentage of Th17 and Th17/Th1 cells in the peripheral blood of patients with cancer compared with that of healthy controls was assessed using t-tests. Cytokines IL17, IL23 and ROR γ t in LSCC tissues compared with that of non-cancerous tissues were assessed using one-way analysis of variance. All statistical calculations were performed using SPSS version 19 (IBM SPSS, Armonk, NY, USA). $P < 0.05$ was considered to indicate a statistical significant difference.

Results

Descriptive features of the study groups. Table I shows the descriptive characteristics of the study subjects, comprising

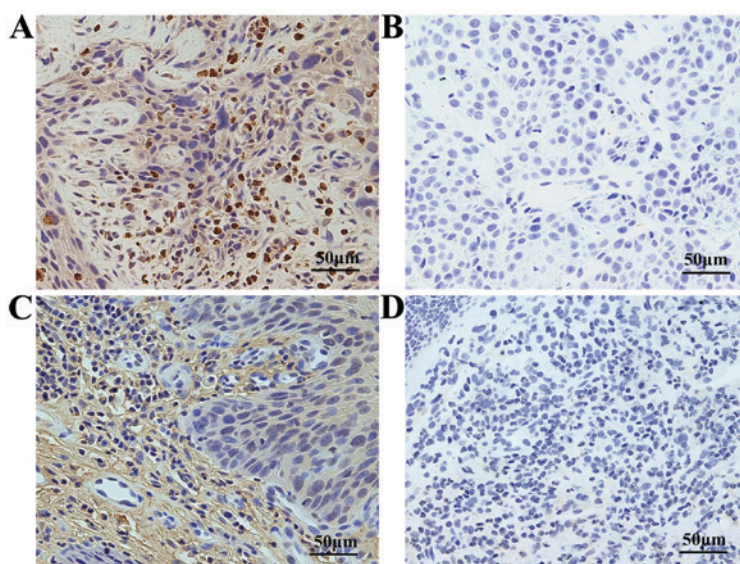


Figure 2. Positive and negative expression of IL17/IL17R immunohistochemistry in tissue microarray sections. (A) Positive and (B) negative expression of IL17. (C) Positive and (D) negative expression of IL17R. Original magnification, x20. IL, interleukin; R, receptor.

Table II. Positive results of IL17/IL17R immunohistochemistry in tumors and controls.

Molecule	Tumors, n (%)	Controls, n (%)	P-value
IL17	57 (81.43)	26 (37.14)	<0.05
IL17R	64 (91.43)	49 (70.00)	<0.05

IL, interleukin; R, receptor.

116 patients with LSCC and 16 healthy controls. Pathologically, all patients had SCC. Of a total of 116 patients, 115 patients (99.14%) were males and only 1 (0.86%) was female. The control group had a similar sex distribution, including 16 males (100.00%).

Immunohistochemical observations. To study the expression of IL17 and IL17R *in vivo*, the tumors and adjacent tissues were stained for IL17 and IL17R using immunohistochemistry (Fig. 2). Differential IL-17 and IL-17R expression in tumors and adjacent tissues was confirmed.

Overall, positive staining for IL17 was noted in 57 (81.43%) of 70 tumors with the following scores: 0, 13 (18.57%) of 70; 1-4, 16 (22.86%) of 70; 5-8, 27 (38.57%) of 76; and 9-12, 14 (20.00%) of 70 samples. Among the controls, positive staining for IL17 was noted in 26 (37.14%) of 70 samples. IL17R positive staining was observed in 64 (91.43%) of 70 cases. These included 0, 6 (8.57%) of 70; 1-4, 22 (31.43%) of 70; 5-8, 28 (40.00%) of 70; and 9-12, 14 (20.00%) of 70 cases. Among the controls, positive staining for IL17R was noted in 49 (70.00%) of 70 samples (Table II).

Elevated levels of Th17 cells in the peripheral blood of patients with LSCC. The prevalence of Th17 cells in the peripheral blood of 36 patients with LSCC was compared with that of

16 healthy controls to address whether there was an increased prevalence of Th17 cells in patients with LSCC. PBMCs were isolated and stimulated with PMA and ionomycin for 5 h in the presence of Protein Transport Inhibitor, and Th17 and Th1 cells were next quantified by flow cytometry. Th17 cells were identified as a CD4⁺IL17⁺ cell population. Th1 cells were identified as a CD4⁺IFN- γ ⁺ cell population. As shown in Figs. 3 and 4A, there was a statistically significantly higher percentage of Th17 cells in the peripheral blood of patients with cancer ($1.6860 \pm 0.1866\%$, $n=36$) compared with that of healthy controls ($0.4963 \pm 0.7862\%$, $n=16$; $P<0.05$).

Th17/Th1 cells are increased in the peripheral blood of patients with LSCC. Th17/Th1 cells, which simultaneously produce the Th17 cytokine IL17 and the Th1 cytokine IFN- γ (16), accounted for $\sim 3.714\%$ in healthy individuals ($3.714 \pm 0.3487\%$; $n=16$). In the peripheral blood of patients with LSCC, this subpopulation was significantly increased ($21.03 \pm 1.750\%$; $n=36$; $P<0.05$; Fig. 4B).

Upregulation of intracellular cytokines IL17, IL23 and ROR γ t in Th17 cells. RT-qPCR revealed that the Th17-associated intracellular cytokines and transcription factors IL17, IL23 and ROR γ t in LSCC tissues were upregulated compared with their levels in matched adjacent non-cancerous tissues (Fig. 5). This indicated that Th17 cells expanded in the LSCC tissue cell population, and may be involved in LSCC development and metastasis.

Discussion

Tumors grow in a complex and active microenvironment. Within or nearby the tumor nests, lymphocytes as well as endothelial, stromal and innate cells are present, which interact with each other to form the tumor microenvironment (17). Inflammation is causally associated with cancer development through processes that involve genotoxicity, aberrant tissue repair, proliferative responses, invasion and metastasis (17).

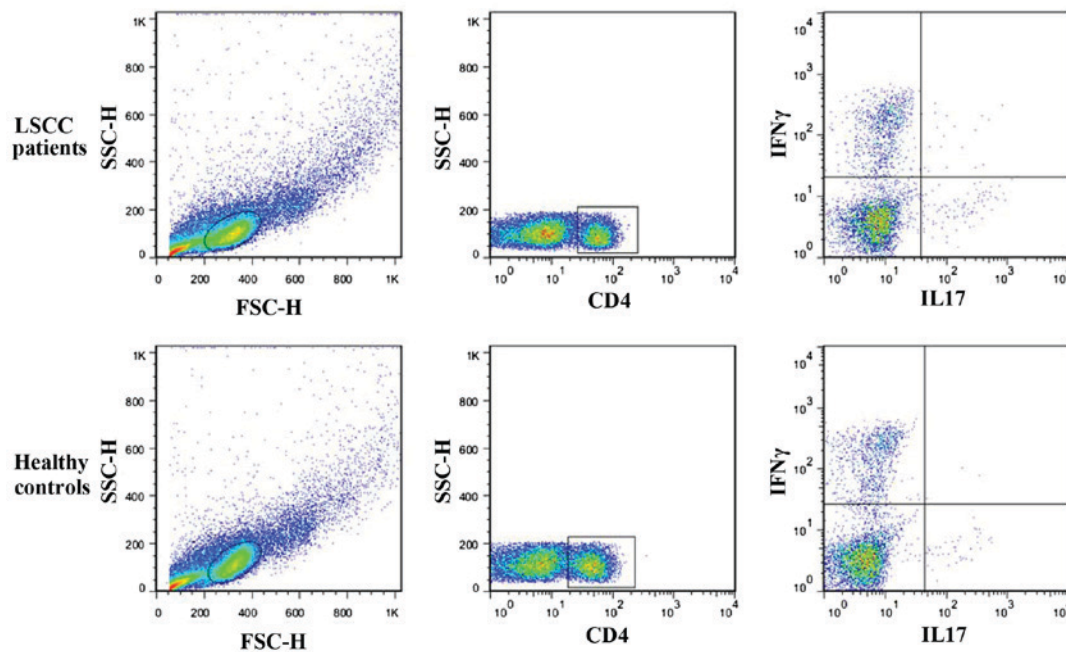


Figure 3. Levels of Th17 and Th1 cells in peripheral blood mononuclear cells from patients with LSCC and healthy controls. SSC, side scatter; FSC, forward scatter; Th, T helper; LSCC, laryngeal squamous cell carcinoma; CD, cluster of differentiation; IL, interleukin; IFN, interferon; H, height.

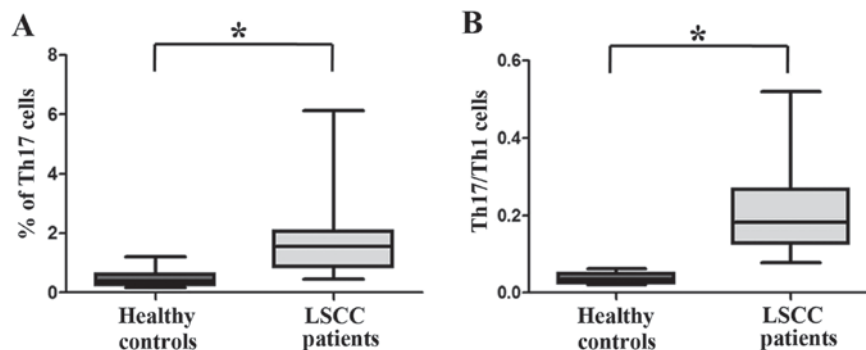


Figure 4. Prevalence of Th17 and Th17/Th1 cells in the peripheral blood of patients with LSCC and healthy controls. (A) Comparison of the prevalence of Th17 cells in the peripheral blood of healthy controls and patients with LSCC (* $P<0.05$). The data are expressed as the frequency of Th17 cells in the lymphocyte population. The box plots show the median (middle line), 25th and 75th percentiles (box), and the extreme values (whiskers). (B) Prevalence of Th17/Th1 cells in the peripheral blood of patients with LSCC compared with that in healthy controls with (* $P<0.05$). The box plots show the median (middle line), 25th and 75th percentiles (box), and the extreme values (whiskers). Th, T helper; LSCC, laryngeal squamous cell carcinoma.

Tumors modulate the inflammatory environment through the secretion of soluble growth factors and chemoattractants, which render inflammatory cells suppressive against anti-cancer T cell responses (17). $CD4^+$ Th cells, as a highly heterogeneous population, serve critical roles in tumor immunological responses (18). Initially, immunologists considered that $CD4^+$ T cells mainly included two types, namely Th1 and Th2 (19); however, at least four distinct $CD4^+$ T cell subsets have been confirmed to exist, specifically Th1, Th2, Th17 and Treg cells (19).

LSCCs produce various immunosuppressive and tumor-promoting cytokines, leading to an impaired anti-tumor response (20). In previous cytogenetic studies, it was observed that multiple signaling pathways were activated in LSCC, including the p53, vascular endothelial growth factor, epidermal growth factor receptor, transforming growth factor (TGF)- β and nuclear factor- κ B signaling pathways (21). In addition, it has

been well established that inflammation is closely connected to LSCC development due to the induction of chronic inflammation caused by exposure to irritants in inhaled air, particularly cigarette smoke. Such irritants enhance the accumulation of viruses and airborne microbes, thereby promoting tumor growth (22). Considering these facts, it can be concluded that pro-inflammatory Th17 cells have extensive effects on LSCC pathogenesis and anti-tumor response.

It was previously shown that Th17 cells are involved in tissue inflammation by inducing the release of various cytokines, including IL6, IL21, IL23, IL1 β and TGF- β , by neighboring tumor cells, tumor-derived fibroblasts and antigen-presenting cells (23). Human Th17 cells mainly release the pro-inflammatory cytokine IL17, and one important role of IL17 appears to be the regulation of local inflammation through the upregulation of other pro-inflammatory cytokines and chemokines (24). In this regard, it is reasonable to propose

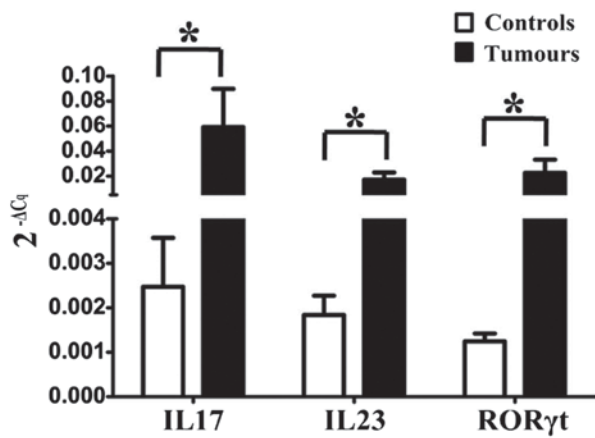


Figure 5. Reverse transcription-quantitative polymerase chain reaction analysis of the Th17-associated intracellular cytokines and transcription factors IL17, IL23 and RORγt in tumors and controls. *P<0.05. IL, interleukin; ROR, RAR-related orphan receptor; Cq, quantification cycle.

the impact of Th17 cells on cancer pathogenesis and progression. In gastric cancer, there was both an elevation of Th17 frequency in peripheral blood and tumor-draining lymph nodes (25). Furthermore, in ovarian cancer, it was shown that Th17 cells were increased in tumor tissue, but not in peripheral blood (15). However, the function of Th17 cells and their influence on LSCC development in humans remain unknown.

Several studies demonstrated that human Th17 cells produce IL17A and exhibit RORγt expression (26,27). These cells also express IL17R, which is also expressed by Th1 cells (26,27). Human Th17 cells appear to exclusively originate from a small subset of T-cell precursors, which constitutively express RORγt and IL-17R, and develop into Th17 cells in response to IL1β and IL23 in the apparent absence of TGF-β (28). Furthermore, even established Th17 cells can be induced to produce IFN-γ in addition to IL17A (Th17/Th1 cells), suggesting a common developmental association between Th17 and at least a subset of Th1 cells (28). Notably, numerous studies have reported increased expression of IL17/Th17 cells in different cancers, including colorectal (29), prostate (30), gastric (31), lung (32), ovarian (33), oral (34) and head and neck (35) carcinomas.

The present study revealed that patients with LSCC have elevated levels of Th17 cells in their primary tumors and peripheral blood compared with those exhibited by healthy controls. The LSCC microenvironment was identified as a strong Th17-cell inducer. First, histopathological characterization was performed, and immunohistochemistry staining was used to detect IL17 and IL17R expression in patients with LSCC. Overexpression of both IL17 and IL17R was observed in tumors compared with the expression detected in adjacent tissues. Next, whether Th17 cells have any functional implications for LSCC development was evaluated. The frequencies of Th17 and Th1 cells in the peripheral blood of patients with LSCC were investigated, and a higher percentage of Th17 cells was detected in the peripheral blood of patients with LSCC compared with that in healthy controls. No association was observed between TNM staging and Th17 cell frequency, which suggests that Th17 cells were consistently elevated in patients with LSCC, independently of tumor stage. This may

represent a link between inflammation and cancer; however, the exact mechanisms by which elevated levels of pro-inflammatory Th17 cells and the resultant secretion of cytokines contribute to inflammatory processes in cancer remain to be elucidated.

The present study demonstrated that Th17 cells can be induced by PBMCs of patients with LSCC. Upon incubation of isolated naïve CD4⁺ T cells in LSCC PBMCs *in vitro*, a significantly elevated number of Th17 cells was detected. Therefore, it can be proposed that the LSCC microenvironment is able to induce a Th17 lineage commitment. It was also observed that Th1 cells were downregulated in peripheral blood. It is known that Th1 cytokines, such as IFN-γ, are diminished in LSCC (36), whereas IL17 is upregulated; thus, this may be the mechanism responsible for Th17 and Th1/Th17 cell modulation in LSCC. However, the outcome of these Th1/Th17 cells under tumor influence (whether they simply attenuate their IFN-γ secretion or change into another T-cell population) remains to be investigated. Our hypothesis is that Th1 cells are functionally modulated by the tumor microenvironment and are converted into Th17 cells.

RT-qPCR demonstrated that the levels of Th17-associated intracellular cytokines and transcription factors (including IL17, IL23 and RORγt) of LSCC tissue were upregulated. Since it was shown that IL23 and RORγt lead to Th17 expansion (37), as well as to selective enrichment of IL17-producing cells by modulating the proliferation of memory T cells (38), the present study was able to show that IL23 and RORγt levels in LSCC lead to a strong enhancement of Th17 cell expansion directly at the tumor site. There have been different reports about the influence of Th17 on tumor progression, a number of which depict a positive influence of Th17 on tumor proliferation (39,40), while others describe an inhibitory influence on tumor growth (41,42). Ciree *et al* (43) demonstrated that Th17 cells are upregulated in the T-cell lymphomas mycosis fungoides and Sézary syndrome, and may act as a tumor growth-promoting or -inhibiting factor. In addition, the authors observed an association between Th17 expression and polymorphonuclear neutrophil infiltration. Muranski *et al* (44) reported that Th17 cells in a mouse model were able to eradicate melanomas. This association was affirmed by Garcia-Hernandez *et al* (45); in their description, neutrophils were attracted to the tumor microenvironment by a Th17-dependent mechanism, and indicated that depletion of neutrophils resulted in a diminished capacity to control tumor growth. In addition, Honorati *et al* (46) reported an increased susceptibility of osteosarcoma cells to natural killer cells under the influence of Th17.

Altogether, the present study demonstrated that Th17 cells are highly present in LSCC. However, this appears to act as a double-edged sword: On one hand, Th17 cells accelerate metastasis and appear to be, therefore, beneficial to tumors; on the other hand, Th17 cells appear to be beneficial to the host due to their proliferation-reducing activity. Therefore, it is important to study the function of Th17 cells in malignant diseases in depth, and to attempt to elucidate their mechanism of action and their modulation by the tumor microenvironment. The present results raise a further issue to investigate, namely whether Th17 cells express different molecules or secrete different cytokines in patients with LSCC compared with those in healthy individuals. Future studies should

consider whether Th17 cells can recognize tumor cells and if they are at all able to impair tumor growth or metastasis *in vivo*; in addition, their exact potential mechanism of action must be elucidated.

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