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The Activation Pattern of Blood Leukocytes in Head and Neck Squamous Cell Carcinoma Is Correlated to Survival

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Abstract

Head and neck squamous cell carcinoma (HNSCC) is known to cause substantial immunosuppression. The present study was designed to characterize blood leukocyte activation in HNSCC and to investigate if the individual activation pattern could be related to tumor progress and survival. The leukocyte activation profile of HNSCC patients and healthy controls was assessed with flow cytometry. HNSCC patients displayed increased numbers of monocytes, neutrophils and total leukocytes as well as an enhanced neutrophil/lymphocyte ratio. In addition, patients had a higher percentage of CD69⁺, CD71⁺ and CD98⁺ T cell subsets and NK cells, and a reduced expression of L-selectin in CD14highCD16⁺ monocytes and neutrophils when compared to controls. These changes could be correlated to both tumor burden and spread to lymph nodes. Among the cancer patients an increased neutrophil/lymphocyte ratio, a low neutrophil and CD14high CD16⁺ monocyte activation state and an elevated CD4/CD8 ratio were related to poor survival. In contrast, a high percentage of CD98⁺ Th cells appeared to be associated with a better outcome. Taken together, the present data indicate that HNSCC causes activation of blood leukocytes and that the individual activation pattern can be linked to prognosis.

Introduction

Head and neck squamous cell carcinoma (HNSCC) is aggressive in nature. It induces production of cytokines and growth factors that regulate the expression of genes controlling growth, survival, and chemosensitivity [1,2]. Such dysregulation of the inflammatory response is believed to perpetuate the malignant phenotype. In addition, HNSCC tumors have the ability to produce immunosuppressive mediators that affect the immune function of the host. Local interactions between the tumor and infiltrating leukocytes are also suggested to cause immunological alterations, e.g., an increased number of activated T cells [1,2,3,4,5,6]. However, reports concerning leukocyte activation are not unequivocal. Some authors have found reduced leukocyte numbers among HNSCC patients, whereas others have not [3,4,7]. The present study was designed to investigate the role of leukocytes and their activation in HNSCC. To this end, peripheral blood from newly diagnosed, still untreated patients was compared to blood obtained from healthy age- and gender-matched controls. An attempt to link these changes to tumor burden, lymphatic spread and survival was also made.

Materials and Methods

Ethical Statement

The study was approved by the Ethics Committees of Karolinska Institutet and Lund University and a written informed consent was obtained from all participants.

Patients

In total, 20 patients (14 males and 6 females) diagnosed with HNSCC were sampled before initiation of treatment along with 20 healthy controls (12 males and 8 females). The median age of the patients was 69 years (range 52–87) and of the controls 70 years (range 51–89). The control individuals were closely matched to the cancer patients regarding age, gender, medication, smoking and alcohol consumption. Neither control subjects nor HNSCC patients had autoimmune disorders, ongoing immune modulating medication or a previous history of malignant diseases. The clinical tumor (T) and lymph node (N) classification of the cancer patients at the time of inclusion are shown in Table 1. The patients’ different tumor locations were as follows: 4 epipharyngeal, 2 esophageal, 6 tonsillar, 2 gingival, 1 laryngeal, 3 hypopharyngeal, 1 tongue, and finally 1 dermal location.
Table 1. Clinical tumor (T) and lymph node (N) classification of the HNSCC patients.

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¹x – Tumor size not determined.

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Blood sampling

Two blood samples were collected from each individual; 4 ml in a test tube containing EDTA for leukocyte differential count analysis performed on a Coulter® LH750/GenS cell counter (Beckman Coulter, Marseille, France), and another 4 ml in a tube containing buffered tri-sodium citrate for flow cytometry analysis. To ensure identical blood sampling the procedure was carried out in the morning after at least 30 min rest.

Antibodies and reagents

The monoclonal antibody (mAb) combinations used for flow cytometry analysis are presented in the Table S1. The following mAbs detecting various surface antigens were purchased from Beckman Coulter: CD3-EC (clone UCHT1), CD4-PCy5(13B8.2), CD11c-RPE (BU15), CD14-PCy5 (RM052), CD16-EC (3G8), CD16-PCy5 (3G8), CD25-EC (B1.49.9), CD56-PCy5 (N901), CD62L-RPE (DREG56), CD64-PCy5 (22), CD69-RPE (TP1-55-3), CD69-EC (TP1-55-3), CD71-FTTC (YD)1-2.2, chemoattractant receptor-homologous molecule expressed on T helper (Th)2 cells (CRTH2)-RPE (BM16) and human leukocyte antigen (HLA)-DR-PCy5 (Immu357). CD8-FITC (44D7) was obtained from Serotec (Oxford, UK), whereas for Seahorse box (Phos)3-FITC (PCH(101) and CD123-FTTC (6H6) were from eBioscience (San Diego, CA). CD8-FITC (DR25) and CD14-FTTC (DUC4) were purchased from DakoCytomation (Glostrup, Denmark), while CD71-PCy5 (M-A712) and lineage cocktail (Lin; FTTC-conjugated mAbs directed against CD3 (SK7), CD14 (MP9), CD16 (3G8), CD19 (SJ25C1), CD20 (L27) and CD56 (NCAM16.2)) were from BD Bioscience (San Jose, CA). An RPE-conjugated mAb against blood dendritic cell antigen 2 (BDCA2; AC144) was from Miltenyi Biotec (Bergisch Gladbach, Germany). The following isotype controls were used: mIgG-FITC, mIgG1-RPE (P3) and mIgG1-PCy5 (P3) from eBioscience, and mIgG2b-EC (MPC-11) from Beckman Coulter.

Flow cytometry analysis

Blood from 20 HNSCC patients and 20 healthy controls was analyzed on a Coulter Epics XL flow cytometer (Beckman Coulter) and leukocytes were gated based on forward and side scatter properties (Figure S1). Events in the range 40,000-200,000 were collected depending on the occurrence of the investigated leukocyte population, and analyzed with Expen32 analysis software (Beckman Coulter). To ensure flow cytometric standardization, the voltage settings were updated daily using FlowSet calibration beads (Beckman Coulter). All mAbs were titrated before use, and staining intensity was controlled on a weekly basis. For extracellular staining, 50 μl blood was incubated with mAbs for 20 min. Erythrocytes were lysed with 600 μl 0.1% (v/v) formic acid for 3–5 seconds and the ionic strength was rendered iso-osmotic by addition of 200 μl 51 mM Na2CO3, 0.20 M Na2SO4 and 0.22 M NaCl. Intracellular staining of Foxp3 was performed using the IntraPrep™ Permeabilization Reagent Kit, according to instructions of the manufacturer (Beckman Coulter). For both extra- and intracellular staining, cells were washed in PBS and resuspended in PBS containing 1% formaldehyde prior to analysis.

Statistics

Statistical analysis was performed using GraphPad Prism 5 (GraphPad Software, San Diego, CA). For normally distributed data statistical analysis was determined using unpaired Student’s t-test, with Welch correction if the variances were non-homogenous. The nonparametric Mann Whitney test was used for not normally distributed unpaired data. \( \alpha \) is equal to the number of independent donors and \( \beta \)-values \( \leq 0.05 \) were considered statistically significant.

The survival function from life-time data was estimated using Kaplan-Meier analysis and a log rank test was utilized to examine the significance of the difference of the survival distribution between the groups.

Results

A leukocyte differential count indicated that the total number of leukocytes, neutrophils and monocytes was higher in HNSCC patients than among controls (Fig. 1A–C). No such difference was observed for lymphocytes, eosinophils or basophils. The neutrophil/lymphocyte ratio, suggested to be an indicator of prognosis for various types of cancer [8,9,10,11], was higher among the cancer patients (Fig. 1D). The percentage of various leukocyte subsets, CD3+ T cells, CD8+ cytotoxic T lymphocytes (CTLs), CD4+ Th cells (comprising both Th1 and Th2 cells), CD4+CD25+Foxp3 regulatory T cells (Treg), along with CD3+CD56+CD16+ NK cells, were analyzed using flow cytometry. In contrast to the differential count data, the flow cytometry analyses showed no change in the different leukocyte subsets between the HNSCC patients and the control individuals. Neither did the distribution of dendritic cell subsets (DCs), as judged by staining of CD123+BDCA2+ plasmacytoid DCs (pDCs) and Lin−CD11c+HLA-DR+ myeloid DCs (mDCs), show any differences between the two groups analyzed (data not shown).

HNSCC patients had a higher value of the total T cell population positively stained for the early activation marker CD69 and the proliferation marker CD71 compared to healthy individuals (Fig. 2A and B). Further analyses of various T cell subsets revealed that an increased frequency of CTLs from cancer patients (Fig. 1D). The percentage of various leukocyte subsets, CD3+ T cells, CD8+ cytotoxic T lymphocytes (CTLs), CD4+ Th cells (comprising both Th1 and Th2 cells), CD4+CD25+Foxp3 regulatory T cells (Treg), along with CD3+CD56+CD16+ NK cells, were analyzed using flow cytometry. In contrast to the differential count data, the flow cytometry analyses showed no change in the different leukocyte subsets between the HNSCC patients and the control individuals. Neither did the distribution of dendritic cell subsets (DCs), as judged by staining of CD123+BDCA2+ plasmacytoid DCs (pDCs) and Lin−CD11c+HLA-DR+ myeloid DCs (mDCs), show any differences between the two groups analyzed (data not shown).

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CD14\textsuperscript{high}CD16\textsuperscript{+} monocytes exhibited a low mean fluorescence intensity (MFI) of L-selectin (Fig. 2J, K) in HNSCC patients. No other discrepancies were observed.

When the size of the primary tumor, as either T1/T2 or T3/T4, was related to the individual leukocyte activation pattern the T3/T4 group presented a higher number of Th cells and a lower percentage of CTL than the T1/T2 group (Fig. 3A and B). Accordingly, the CD4/CD8 ratio, often used as a prognostic indicator [12,13], was higher among the T3/4 patients (Fig. 3C).

In addition, the number of CD69 expressing monocytes as well as the CD14\textsuperscript{high}CD16\textsuperscript{+} and CD14\textsuperscript{high}CD16\textsuperscript{+} subgroups, was increased among those with a more advanced disease. This group had a higher frequency of CD14\textsuperscript{dim}CD16\textsuperscript{+} monocytes positive for L-selectin (Fig. 3D–G). It is also worth noticing that patients with lymphatic spread (N+) exhibited a higher neutrophil/lymphocyte ratio than patients without metastases (N0). The N+ patients also displayed an elevated percentage of CD69\textsuperscript{+} and CD71\textsuperscript{+} T cells, a higher number of Th cells expressing CD71 and a higher CD71 density (MFI) on their NK cells (Fig. 4A–E).

To evaluate the prognostic value of the leukocyte markers, described above, the patient’s status at the time of cancer discovery, before any treatment was given, was related to the total survival during a 24 months observation period. The HNSCC group was divided according to the mean values of the different parameters, characterized as either higher than mean or lower than mean. The group with a high neutrophil/lymphocyte ratio had a worse outcome than the lower mean ratio group. A higher number of activated neutrophils and CD14\textsuperscript{high}CD16\textsuperscript{+} monocytes, determined by a lower CD62L expression, and an elevated number of CD96\textsuperscript{+} Th cells predicted a better survival (Fig. 5A–D). The CD4/CD8 ratio is an index with disputed prognostic value [3,13,14,15,16]. Here we show that patients with a high ratio were found to have a reduced survival (Fig. 5E). Generally, the prognostic value of the leukocyte factors seemed to be most pronounced during the first 12 months after diagnosis (Fig. 5A–E).

**Discussion**

The present study suggests that the HNSCC patients display increased numbers of total leukocytes, neutrophils and monocytes, and accordingly a higher neutrophil/lymphocyte ratio than control subjects. The cancer patients also exhibit an enhanced percentage of activated T cell subsets and NK cells, as determined by an increase in the density of CD69, CD71 and CD98. In addition, CD14\textsuperscript{high}CD16\textsuperscript{+} monocytes and neutrophils from HNSCC patients have a lower expression of L-selectin. The activation frequency of total T cells, Th cells, NK cells and monocyte populations seems to correlate with tumor burden and lymphatic spread in such a way that a high state of activation can be found among patients with a more severe disease. It also appears as if the neutrophil/lymphocyte ratio as well as the activation status of neutrophils, CD14\textsuperscript{high}CD16\textsuperscript{+} monocytes and Th cells, at the time of cancer diagnosis, is linked to the patient’s life expectancy.

High numbers of neutrophils and monocytes were observed in HNSCC patients using differential count analysis. No such differences were found with flow cytometry. This disparity could be related entirely to the analysis methods used. It needs to be emphasized that classification of leukocytes by differential count...
depends on volume, conductivity and light scattering, whereas
flow cytometry simply distinguishes leukocyte subsets by their cell
surface markers. The difference in results may also be affected by
the way data is presented as either absolute number or as
percentage. In line with this, Kuss et al. has shown a reduced
absolute number of total T cells, Th cells and CTLs in patients
with HNSCC without any corresponding changes in the
percentage [7].

The increased number of neutrophils and monocytes seen in the
HNSCC patients are supported by previous reports showing
similar findings in other types of cancer [8,17,18]. The increase
might reflect an increased inflammation and an enhanced
infiltration of immature neutrophils and monocytes from the bone
marrow as a consequence of an increased leukocyte turnover.
Accordingly, the high neutrophil-lymphocyte ratio, that was
observed in the HNSCC patients, indicates an ongoing systemic
inflammation [19]. Furthermore, a high neutrophil-lymphocyte
ratio appears to be synonymous with a reduced survival rate,
which suggests that a high systemic inflammation is linked to the
patient’s life expectancy. This is in accordance to recent reports
suggesting this ratio to be a potential prognostic factor in several
forms of cancer [8,9,10,11,19]. The role of neutrophils in cancer
has recently gained attention. They are thought to be pro-
tumorigenic by secretion of pro-angiogenic substances and
suppression of the adaptive immune system [20,21,22]. In
contrast, there are studies reporting an anti-tumorigenic role for
these cells. Gregory and Houghton have suggested that activated
neutrophils can elicit antitumor activity [22,23]. In support of this,
the present data demonstrates that HNSCC patients with a higher
frequency of activated neutrophils have a better survival.

CD14<sup>hi</sup>CD16<sup>+</sup> monocytes are known to produce IL-10. The
enhanced activation of these monocytes seen among the HNSCC
patients with a higher number of total T cells, Th cells and CTLs
was surprising. However, these monocytes might have until now
unknown anti-tumor functions that might contribute to an
increased survival. The present data indicate that, with the
exception of the more mature monocyte population
CD14<sup>dim</sup>CD16<sup>+</sup>, patients with an increased tumor burden

Figure 2. Blood from HNSCC patients (n=20) and controls (n=20) was incubated with different Abs and analyzed with flow

cytometry. (A, B) Abs against CD3, CD69 and CD71 were used to distinguish activated total T cells. (C, D) To identify activated cytotoxic T
lymphocytes (CTLs) CD69, CD71 and CD8 Abs were used. (E, F, G) CD4<sup>+</sup> T helper (Th) cells were discriminated by CD4 Abs, and the activation status
was established using CD69, CD71 and CD98 Abs. (H) Th2 cells were detected by CD4 and CRTH2 Abs, and the frequency of CD98<sup>+</sup> Th2 cells was
determined. (I) CD3, CD16, CD56 and CD69 Abs were used to discriminate CD69<sup>+</sup> CD3<sup>+</sup> CD16<sup>+</sup> CD56<sup>+</sup> natural killer (NK) cells. (J) Neutrophils were
determined as CD16<sup>+</sup> granulocytes, and the mean fluorescence intensity (MFI) of CD62L<sup>+</sup> cells was calculated. (K) Staining of CD14, CD16 and CD62L
was used to discriminate activation of the monocyte population CD14<sup>hi</sup>CD16<sup>+</sup>. The MFI of CD62L<sup>+</sup> CD14<sup>hi</sup>CD16<sup>+</sup> monocytes was established. *

p<0.05; ** p<0.01; *** p<0.001.
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generally exhibit a more active monocytes profile. It has previously been reported that there is an increased activation of immature macrophages in cancer patients [26]. This indicates that the activated monocytes in patients with a more advanced disease might be of a more immature phenotype.

The tumor appears to affect the frequency of CTLs and Th cells, as demonstrated by an increase in Th cells and a decrease in CTLs as well as a higher CD4/CD8 ratio, and thereby inhibiting the anti-tumorigenic CTL response and promoting a pro-tumorigenic environment. In accordance, previous reports have demonstrated an increase in the percentage of T cells and Th cells in addition to a decrease in CTLs in patients with an advanced tumor disease [4]. A high CD4/CD8 ratio was also connected to poor survival. The prognostic value of this ratio has been disputed by several studies showing contradictory results [3,13,14,15,16]. The present data supports the notion that this ratio indicates that patients with a high anti-tumorigenic response have a better chance of survival.

The high number of activated T cell subsets and NK cells among the HNSCC patients and specifically in patients with a verified lymph node metastasis reveals an enhanced immune activation that corresponds well with the increased neutrophil-lymphocyte ratio. Both indicate an accelerated systemic immune activity and an enhanced leukocyte turnover. In agreement, increased levels of CD69+ T cells have been shown in patients with advanced HNSCC [4]. The increase in Th2 activation might also be the result of the tumors to promote a pro-tumorigenic immune response. The increased activation of the Th2 cells was only accompanied by an increased expression of CD98 and not CD69 or CD71. It is not inconceivable that this observation reflects the fact that the microenvironment of the tumor attracts individual cells. The increased Th2 cell activation is well in analogy with previous studies reporting high levels of Th2 cytokines in serum, including IL-4, IL-6, IL-10 and granulocyte-monocyte colony-stimulating factor (GM-CSF) [1,27]. The increased frequency of CD98+ Th cells that was presently found to be linked to survival.

Figure 3. Blood from HNSCC patients was incubated with various Abs, followed by flow cytometry analysis. The HNSCC patients were divided according to the size of the primary tumor (stage T1/T2 versus T3/4; n = 9 and n = 11, respectively). (A) T helper (Th) cells were defined as CD4+ lymphocytes, whereas (B) cytotoxic T lymphocytes (CTLs) were identified by CD8, and (C) the CD4/CD8 ratio was calculated. (D) CD14 and CD69 were used to discriminate activated monocytes. (E) CD69 expressing CD14highCD16− monocytes were identified. (F) The monocyte population CD14highCD16+ was recognized and the number of CD69+ cells was determined. (G) In addition, the percentage of CD62L+ CD14dimCD16+ monocytes was established. * p<0.05; ** p<0.01.

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could reflect an amplified Th1 anti-tumor activity, since no association was found for activated Th2 cells. NK cells have cytolytic functions and are believed to play a role in tumor immune surveillance [28]. It is therefore tempting to speculate that the higher percentage of activated NK cells...

Figure 4. Blood was obtained from HNSCC patients and divided according to presence of regional lymphatic node metastases (N0 versus N+; n = 10 and n = 10, respectively). (A) The neutrophil/lymphocyte ratio was determined by leukocyte differential count analysis. (B, C) Blood was incubated with CD3, CD69 and CD71 Abs, and the percentage of activated total T cells was established with flow cytometry. (D) Abs against CD4 and CD71 were used to identify activated T helper (Th) cells. (E) The mean fluorescence intensity (MFI) of CD62L⁺CD3⁻CD56⁻CD16⁺ natural killer (NK) cells was calculated. * p≤0.05.

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Figure 5. A Kaplan-Meier survival analysis following diagnosis of HNSCC patients, before start of treatment, appeared to be related to (A) the neutrophil/lymphocyte ratio, (B) expression of CD98 among the Th cells, (C) CD62L among neutrophils, and (D) CD14⁺CD16⁺ monocytes, and (E) the CD4/CD8 ratio. The patients were divided according to the mean values of the parameters analyzed, characterized as either higher or lower than mean. * p≤0.05.

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presently observed reflects an enhanced immune surveillance activity against the tumor. However, it must be considered that NK cells also have been attributed to impair anti-tumor functions in HNSCC [6,29].

To conclude, the present study demonstrates that there is an increased systemic inflammation in HNSCC patients. This was for instance determined by an increased activation of leukocytes. It is also clear that a higher frequency of activation is found among patients with a more severe disease. Although, the number of patients investigated is limited and the cancer population is heterogeneous the results are clear cut indicating that the leukocyte activation state at the time of diagnosis can be of prognostic value for survival. Further, it signifies a tight link between the immune phenotype and the anti-tumor immune response and hence the survival of the patient.

Supporting Information

Table S1 Antibody panel used for flow cytometry analysis. 3Th cells = T helper cells; 2CTLs = cytotoxic T lymphocytes; 3NK cells = natural killer cells; 1pDCs = plasmacytoid dendritic cells; 5mDCs = myeloid dendritic cells; 6Linage cocktail = CD3-, CD14-, CD16-, CD19-, CD20-, CD56-FITC.

References


Figure S1 Blood from HNSCC patients (n = 20) and controls (n = 20) was incubated with various Abs and analyzed with flow cytometry. Lymphocytes, monocytes and granulocytes were distinguished based on FS and SS plotting. From these cell populations, CD3+ T cells, CD8+ cytotoxic T lymphocytes (CTLs), CD4+ T helper (Th) cells, CD4+CD102+CD19+ Th2 cells, CD5+CD56+CD16+ natural killer (NK) cells, CD16+ neutrophils, CD14+ monocytes, CD14highCD16+ monocytes, CD14highCD16+ monocytes and CD14dimCD16+ monocytes were discriminated. Staining of CD62L, CD69, CD71 and CD98 was used to determine the activation of the different leukocyte subsets. (TIF)

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Author Contributions

Conceived and designed the experiments: CRM AMK SB LOC. Performed the experiments: CRM. Analyzed the data: CRM. Contributed reagents/materials/analysis tools: RU SB KR LOC. Wrote the paper: CRM. LOC. Supervised total project: LOC.

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