Programmed death of leukocytes through Fas and TNFR1 in malnourished patients with pancreatic cancer

Programowana śmierć leukocytów indukowana przez receptory śmierci Fas i TNFR1 u niedożywionych chorych z rakiem trzustki

Sylwia Kędziora 1, Robert Słotwiński 1,2, Waldemar L Olszewski 1, Marzanna Zaleska 1, Gustaw Lech 1, Maciej Słodkowski 3, Ireneusz W. Krasnodębski 3

1Department of Immunology and Nutrition, Medical University of Warsaw, Poland
2Department of Surgical Research and Transplantology, Medical Research Center, Polish Academy of Sciences, Poland
3Department of General, Gastroenterological and Oncological Surgery, Medical University of Warsaw, Poland

Abstract

Aim: To evaluate the status of programmed leukocyte death (apoptosis) in malnourished pancreatic cancer patients in comparison with healthy controls.

Material and methods: The percentage of leukocytes with death receptors Fas (CD95) and TNFR1 (CD120a) was investigated in 72 pancreatic cancer patients in comparison with 30 healthy volunteers. Percentages of cells with receptors were assessed by flow cytometry.

Results: Results of the study show a significantly higher percentage of CD95+/CD3– cells and lymphocytes with TNFR1 in the group with pancreatic cancer in comparison with healthy controls.

Conclusions: Our studies show up-regulation of the pro-apoptotic signalling system in the leukocytes of patients with pancreatic cancer and a switch to apoptosis. These alterations may lead to leukocyte dysfunction and immune system suppression.

Introduction

Mortality associated with pancreaticoduodenectomy has decreased dramatically to less than 5% over the past 2 decades in high-volume centres, but persistent high morbidity rates have remained an important concern for patients, healthcare providers, and payers [1-6]. While mortality is an objective and easily quantifiable outcome parameter, morbidity is only poorly defined, and this shortcoming has severely hampered conclusive comparisons among centres and within the same institution over time [7-9]. One reason for this high morbidity is immunosuppression in pancreatic cancer [10-12], which contributes to postoperative complications. One process which is especially important for proper functioning of the immune system is apoptosis of immune cells such as leukocytes. Apoptosis or programmed cell death is a cen-
Apoptosis

A central regulator of tissue homeostasis, which causes elimination of redundant, damaged or infected cells. Apoptosis is critical in the formation of organs, limbs and other body structures, and in maintaining the function of most of the systems in an adult body. Therefore, dysregulation of the apoptotic signalling processes often leads to serious consequences, such as neurodegenerative diseases, cancer or autoimmunity [13-16]. One pathway of leukocyte apoptosis is the extrinsic pathway through death receptors such as Fas and TNFR1 (fig. 1). These receptors bind intracellular adapter molecules (TRADD and FADD), which recruit pro-caspase 8, create the death-inducing signalling complex (DISC) and activate caspase 8. Then caspase 8 activates caspase 3, leading to degradation of key intracellular substrates [17].

To evaluate the status of immune cell apoptosis, we investigated expression of death receptors Fas (CD95) and TNFR1 (CD120a) in peripheral blood leukocytes of pancreatic cancer patients in comparison with healthy controls.

Material and methods

Patients

Eighty-eight patients with pancreatic cancer were admitted to the study. According to the criteria of investigation 16 patients were excluded (patients over 75, patients with metastasis, patients with unrectactable tumours, patients treated with chemotherapy or immunotherapies, and those suffering from chronic liver or kidney diseases and diabetes). Seventy-two patients (mean age 62.1 ±8.9) were classified according to UICC (TNM classification of malignant tumours) [18, 19] – 3 patients were classified as stage I, 44 patients were at stage II, and 25 patients were at stage III. After full clinical diagnostic procedures (imaging and laboratory tests) all patients were subjected to pancreatic head resection (Whipple’s pancreaticoduodenectomy). A histopathological examination confirmed the diagnosis.

Nutritional status (loss of body mass, body mass index [BMI], albumin concentration and total lymphocyte count) was assessed. According to the ESPEN (The European Society for Clinical Nutrition and Metabolism) definition of malnutrition [20] our study included 5 patients (6.9%) with slight malnutrition (< 5% weight loss within 2 months), 43 patients (59.7%) with moderate malnutrition (5-10% weight loss within 3 months) and 24 patients (33.3%) with severe malnutrition (> 10% weight loss within 6 months). The characteristics of nutritional status of patients are shown in table I.

The control group comprised 30 healthy sex- and age-matched volunteers.

![Schematic diagram of extrinsic pathway of apoptosis.](image)

**Fig. 1.** Schematic diagram of extrinsic pathway of apoptosis. (A) Binding of TNF-α to receptor TNFR1 allows binding of the intracellular adapter molecules TRADD and FADD and recruitment of pro-caspase 8 leading to formation of the complex DISC. Activation of caspase 8 and activation of caspase 3 initiate apoptosis. (B) Binding of FasL to receptor Fas allows binding of the intracellular adapter molecule FADD and recruitment of pro-caspase 8 leading to formation of the complex DISC. Activation of caspase 8 and activation of caspase 3 initiate apoptosis.

**Ryc. 1.** Schemat zewnętrznego szlaku apoptozy. (A) Przyłączenie TNF-α do receptora TNFR1 pozwala na przyłączenie białek adapterowych TRADD i FADD oraz prokaspazy 8 i utworzenie kompleksu DISC. Aktywacja kaspazy 8 prowadzi do aktywacji kaspazy 3 i inicjuje apoptozę. (B) Przyłączenie FasL do receptora Fas pozwala na przyłączenie białka adapterowego FADD oraz prokaspazy 8 i utworzenie kompleksu DISC. Aktywacja kaspazy 8 prowadzi do aktywacji kaspazy 3 i inicjuje apoptozę

TNF-α – czynnik martwicy nowotworu, TNFR1 – receptor-1 TNF-α, Fas – CD95, FasL – CD95L, ligand receptora Fas, TRADD – białko adapterowe z domeną śmierci (DD) wiązające się z receptorem TNFR1, FADD – białko adapterowe z domeną śmierci (DD) wiązające się z receptorem Fas, DISC – kompleks sygnałowy indukujący śmierć
Flow cytometry

The peripheral blood of each patient was collected for flow cytometry analyses. For staining of surface proteins, 100 µl heparinised blood samples were incubated with 20 µl of the following monoclonal antibodies: CD3 PE, CD95 FITC (Becton Dickinson Biosciences) and CD120a PE (Beckman Culter) and their appropriate isotype-matched control antibodies. After 30 min of incubation at room temperature, red cells were lysed by the addition of FACS Lysing Solution (Becton Dickinson Biosciences) to the tubes. After another 10 minutes, samples were washed in PBS with 1% bovine serum azide (NaN₃) and fixed with 1% paraformaldehyde (Sigma, Chemical Co).

The stained cells were acquired using the LSR instrument (Becton Dickinson) to detect two fluorescence parameters and two scatter parameters. The data were analysed using Cell Quest Pro Software, version 9.2 (Becton Dickinson) and expressed as the percentage of lymphocytes with positive surface protein expressions (means ± SD).

Statistics

Statistical analysis was performed using the StatSoft Statistica v.7.0 program. To evaluate the statistical significance of the difference in percentage of cells with death receptors between the group with pancreatic cancer and the control group Mann-Whitney test with Bonferroni correction was used. Significance was set at \( p < 0.05 \).

Results

The percentage of some leukocytes with receptors involved in programmed lymphocyte death is altered in pancreatic cancer patients as compared to healthy controls. The percentage of CD95+/CD3– cells was significantly higher in pancreatic cancer patients as compared to the healthy control group (\( p = 0.009 \)) (fig. 2, 4a). On the other hand, the percentage of CD95–/CD3+ cells was lower in pancreatic cancer patients as compared to the healthy control group, but there was no significance.

The percentage of lymphocytes, monocytes and granulocytes with receptor TNFR1 (CD120a) was higher in pancreatic cancer patients as compared to the healthy control group. However, there were significant differences only for lymphocytes with TNFR1 between the pancreatic cancer group and the control group (fig. 3, 4b).

Discussion

Our results show significantly higher expression of Fas (CD95) on CD3– cells and TNFR1 (CD120a) on lymphocytes in patients with pancreatic cancer as compared to healthy controls, which corresponds to the overexpression of pro-apoptotic proteins on leukocytes. These alterations revealed up-regulation of the pro-apoptotic signalling system in the leukocytes of patients with pancreatic cancer and switch to apoptosis.

While apoptosis is an essential biological process for normal development and maintenance of tissue homeostasis, it is also involved in a number of pathological conditions including tissue injury, degenerative diseases, immune diseases and cancer [21]. Whether activated by membrane-bound death receptors [22] or by stress-induced mitochondrial perturbation with subsequent cytochrome c release [23], the activation of downstream caspases leads to stepwise cellular destruction by disrupting the cytoskeleton, shutting down DNA replication and repair, degrading chromosomal DNA and finally disintegrating the cell into apoptotic bodies [24]. Several mechanisms of activation of apoptosis in different physiological or pathological conditions in cells have been proposed and studied intensively [25]. Numerous factors, especially death receptors Fas (CD95) and TNFR1 (CD120a), have been suggested to play an essential role in the apoptotic process.
The extrinsic pathway of apoptosis can be induced through members of the TNF/TNFR superfamily such as receptor 1 for TNF-α (TNFR1/CD120a) and receptor Fas (CD95). The receptors and ligands in this superfamily have unique structural attributes that couple them directly to signalling pathways for cell proliferation, survival, and differentiation. Thus, they have assumed prominent roles in the generation of tissues and transient microenvironments. Most TNF/TNFR are expressed in the immune system, where their rapid and potent signalling capabilities are crucial in coordinating the proliferation and protective functions of pathogen-reactive cells [32]. Receptors TNFR1 and Fas take part in the initial stage of apoptosis, which is a reversible phase. Only the execution process (connected with execution caspases, cytoplasmic endonuclease, cytoplasmic proteases, degradation of nuclear and cytoskeletal proteins) is irreversible and definitely indicates cell death [27].

Our findings demonstrate high expression of TNFR1 in the lymphocytes of patients with pancreatic cancer, which may suggest an increased level of apoptosis of peripheral blood lymphocytes in patients with pancreatic cancer by the extrinsic pathway. Other studies have shown a high level of TNFR1 in the lymphocytes of patients after heart transplants [28] and HIV infected patients [29]. Also in patients with head and neck cancer the peripheral blood mononuclear cells had a significantly higher proportion of apoptotic cells than these cells of controls. The authors suggest that activated TNFR1+ and also Fas+ cells are pre-programmed to die [30].

Our data also demonstrate high expression of CD95+ in the CD3– cells of patients with pancreatic cancer, which may also suggest an increased level of programmed death in these cells. Other studies have shown decreased expression of Fas (CD95) on peripheral blood lymphocytes but increased Fas (CD95) expression on tumour-infiltrating lymphocytes in patients with renal cell carcinoma in comparison with healthy volunteers [31]. However, other studies suggest that peripheral blood mononuclear cells of kidney transplant recipients are more susceptible to activation-induced cell death and that the Fas-FasL pathways is involved in this process [32]. Similar results were obtained in patients with non-small cell lung cancer. The proportion of blood CD95+ lymphocytes was significantly higher when compared with healthy people. Moreover, this proportion was higher in smokers when compared with non-smokers [33]. Enhanced expression of Fas receptor on lymphocytes was also observed in gastric mucosa in children during Helicobacter pylori infection [34].

Our results show overexpression of death receptors on leukocytes in patients with pancreatic cancer as compared with healthy controls. These alterations revealed up-regulation of the pro-apoptotic signalling system in the leukocytes of patients with pancreatic cancer and the possibility to switch to apoptosis. This suggests that patients with pancreatic cancer may show an increased level of leukocyte dysfunction and consequently higher immune system suppression. This can further lead to a greater number of infectious complications, organ failure and a higher death rate, especially after pancreatic resection [35-38]. A possible explanation of inappropriate changes in expression of death receptors on leukocytes in patients with pancreatic cancer is that these patients are mostly malnourished. The frequency of malnutrition in patients with pancreatic cancer ranges from 80% to 85% [39]. This is malnutrition which, apart from reduced skeletal, cardiac and respiratory muscle function, poor wound healing and loss of non-fat body mass, causes impairment of immune functions as a response to under-nutrition of immune cells [40]. To improve the nutritional status of immune cells it may be required to introduce pre-operative immune-enhancing diets as a routine procedure after flow cytometry examination. It may lead to some changes in the expression of apoptotic proteins and perhaps a decrease or even elimination of the differences in the expression of these proteins between patients with pancreatic cancer and healthy people.

In conclusion, our studies show up-regulation of the pro-apoptotic signalling system in the leukocytes, mainly lymphocytes, of patients with pancreatic cancer and a possibility to switch to apoptosis as compared with healthy people. These alterations may lead to leukocyte dysfunction and immune system suppression. This prob-

**Fig. 3.** Percentage of leukocytes with receptor TNFR1 (CD120a) in pancreatic cancer patients in comparison with healthy controls

**Ryc. 3.** Odsetek leukocytów z receptorem TNFR1 (CD120a) u chorych z rakiem trzustki w porównaniu z grupą kontrolną
Problem can be resolved by introducing pre-operative immune-enhancing diets in patients with pancreatic cancer as a routine procedure after immunological examination. Further research needs to be undertaken to examine this issue.

Acknowledgements
This work was supported by Projects No. 2 PO5B 05928 and N N402 306836 funded by the Ministry of Science and Higher Education.

References