New system for predicting the outcome of secondary peritonitis

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Abstract: Despite improvements in the methods of diagnostics, surgical interventions and intensive care, the problem of treating patients with diffuse peritonitis remains relevant. Diffuse peritonitis is a major contributor to mortality in all urgent care settings and the second leading cause of sepsis in critically ill patients. At the same time, even in developed countries, the number of patients with peritonitis does not tend to decrease, and mortality rates remain high, reaching 90-93% with the development of abdominal sepsis and toxic shock syndrome. One of the ways to reduce mortality in peritonitis is the use of objective systems for prognosis of the peritonitis outcome, allowing to compare the results of patient treatment and to choose the optimal treatment tactics for each particular patient.

The objective — To develop a new system for predicting the outcome of secondary peritonitis (survival or death) focused on the criteria of abdominal sepsis and multiple organ dysfunction syndrome (associated or not associated with peritonitis), and to analyze its accuracy versus the most common comparable systems.

Material and Methods — Our study was based on analyzing the treatment outcomes in 352 patients with secondary diffuse peritonitis. On admission, sepsis was diagnosed in 15 (4.3%), and toxic shock in 4 (1.1%) patients. The main causes of death were purulent intoxication and/or sepsis (51 cases or 87.9%), cancer intoxication (4 cases or 6.9%), and acute cardiac failure (3 cases or 5.2%). We analyzed the effectiveness of several systems of predicting the peritonitis outcomes: the Mannheim’s Peritoneal Index (MPI), World Society for Emergency Surgery Sepsis Severity Score (WSES SSS), Acute Physiology and Chronic health Evaluation II (APACHE II) system, general Sequential Organ Failure Assessment Score (gSOFA), as well as the Peritonitis Prognosis System (PPS) developed by the authors. The probability of the effect of 40 clinical and laboratory parameters on the outcome of patients with secondary peritonitis was analyzed via using parametric and nonparametric methods of statistical analysis (Fisher’s test, Mann-Whitney U test, Chi-squared test with Yates’s continuity correction). The criteria were selected that had a predictive power for the lethal outcome (p <0.05), and they were included in the PPS system. To compare the predictive value of the PPS, ROC analysis was conducted with construction of receiver operating characteristic curves for each analyzed system of predicting the peritonitis outcome. The STATISTICA 8 software was used for performing the statistical analysis.

Results — The following criteria were of greatest importance in predicting the lethal outcome: a patient’s age, a presence of a malignant neoplasm, the nature of the exudate, the development of sepsis (toxic shock), as well as multiple organ dysfunction not associated with the developed peritonitis. PPS exhibited the greatest accuracy in terms of predicting mortality in patients with secondary diffuse peritonitis (AUC=0.942) versus minimal in APACHE II (AUC=0.840).

Conclusion — APACHE II, MPI, WSES SSS and PPS can be considered reliable in terms of mortality prognosis in peritonitis patients. PPS has the greatest accuracy of predicting the mortality in patients with secondary diffuse peritonitis (94%).

Keywords: Peritonitis, prognosis systems, abdominal sepsis, multiple organ dysfunction syndrome.

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Introduction

A preliminary assessment of the lethality outcome likelihood is used for selecting the rational treatment tactics, along with comparing the treatment outcomes and scientific research results.

Currently, there are several general clinical systems, along with specific schemes, for predicting the outcome of peritonitis, which more or less combine various factors for early prediction of treatment results, but none of such approaches, according to Tolonen M. et al. (2018), work satisfactorily [1].

These algorithms are useful for comparing large groups of patients, but each system has its own drawbacks, the main of which is that they all have limited value in the daily clinical decision-making process for each individual patient [2].

Perhaps this is due to the fact that the parameters of such systems do not take into account (or take into account only superficially) pathological changes in the abdominal cavity observed by the surgeon during the operation [3]. Nonetheless, at present, the following systems for predicting the outcome of peritonitis are widely used: Acute Physiology and Chronic Health
Evaluation II (APACHE II) [4], Simplified Acute Physiology Score (SAPS II) [5], Sequential Organ Failure Assessment Score (SOFA) [6], Mannheim Peritoneal Index (MPI), and World Society for Emergency Surgery Sepsis Severity Score (WSES SSS) [7].

However, APACHE II is of little use in the choice of surgical tactics for treating a patient with peritonitis, since it does not take into account pathological changes in the abdominal cavity and the cause of peritonitis [8]. Besides, APACHE II is aimed at classifying patient groups rather than individual patients, making it difficult to apply to a particular patient [9].

Results of assessment via APACHE II are often reported in the literature as a score rather than a percentage of lethal outcomes. Given that APACHE II scores are presented to describe disease severity in a population, it could be difficult to accurately compare two incongruent patient populations because predicted mortality is based on indications at admission to the intensive care unit.

Use of any specific biochemical parameter as a prognostic indicator is not common [10], and virtually all studied biomarkers of sepsis to date have insufficient specificity due to the development of one or another degree of systemic inflammation in various categories of patients at intensive care units [11].

Comparison of the published data casts doubt on the reliability of the MPI prediction of the peritonitis outcome without taking into account its nature and cause. For example, Billing et al., (1994) considered the threshold value of the assessment at the level of 26 MPI points. For this value, the average sensitivity of the test was 86% (54-98%), its average specificity was 74% (58-97%), and the accuracy of predicting lethal outcome was 83%. Depending on MPI scores, average mortality of the patients varied: for a score of under 21 points, an average mortality was 2.3%; for 21-29 points, it was 22.5%; and for severity score of over 29 points, the mortality was 59.1% [12]. However, when comparing the results of the mentioned study with the data of nine independent studies of mortality in peritonitis, depending on the MPI score, it could be seen that with the same assessment parameter values, deaths occur with different probabilities, and the difference reaches 50% [13].

It is possible that the differences in the mortality rate, depending on the MPI score [14], were caused by different distribution of patients in terms of peritonitis. Unfortunately, in their publications, the authors, as a rule, did not indicate clearly how the patients were distributed by the main diagnosis. According to N.V. Lebedev et al (2017), MPI can be used to evaluate treatment outcomes and research results only in patients with the same cause of peritonitis [15].

Many studies confirmed the accuracy of the WSES SSS in predicting mortality in patients with peritonitis [16, 17, 18].

At the same time, some studies on the comparative characterization of systems for predicting the outcome of peritonitis raise doubts about the accuracy of WSES SSS. For example, studies by Tolonen M. et. al (2019) showed that SOFA score ≥ 2 (78.4%) had the highest identification rates, followed by WSES SSS ≥ 8 (73.1%), SOFA ≥ 3 (75.2%), and APACHE II ≥ 14 (68.8%). The most accurate prediction took place at SOFA score ≥ 2 with 78.4% identification rates, followed by SOFA score ≥ 3 (with 75.2%), WSES SSS ≥ 8 (with 73.1%), and APACHE II ≥ 14 (with 68.8% identification rates) [3]. According to Mohan R. et. al. (2019), MPI is superior to WSES in predicting mortality in patients with peritonitis [19].

The PIPAS system (Physiological Indicators for Prognosis in Abdominal Sepsis) was proposed in 2019 [20]. The creators of this system believed that the PIPAS indicator had a very good ability to determine the likelihood of a lethal outcome.

However, the validity of PIPAS should be tested in future large prospective series before it could be recommended for use in clinical practice. A potential limitation of using PIPAS could possibly be a large proportion of patients with acute appendicitis included in the study (42.1%) [21], as well as the inclusion of patients with severe pancreatitis, which has its own pathophysiological characteristics. Besides, this system does not take into account the intra-abdominal changes in peritonitis and, as a result, is not promising for choosing the surgical tactics. Probably, the PIPAS system is analogous to the general clinical system APACHE II or SAPS II.

The severity of a peritonitis patient condition is associated with a number of clinical and pathophysiological factors. The most important risk factors of a lethal outcome are: the nature of the abdominal lesion and the presence of sepsis with associated organ lesions [2, 22, 23]. At the same time, the presence or absence of abdominal sepsis is of particular importance for the outcome of peritonitis [24].

Other important factors include the physiological reserve of a patient, comorbidities, the state of immune system, and anatomical abnormalities caused by the disease.

The objective. To develop a novel system for predicting the outcome of secondary peritonitis (survival or death) focused on the criteria of abdominal sepsis and multiple organ dysfunction syndrome (MODS), associated or not associated with peritonitis, and to analyze its accuracy against the most common comparable systems.

Material and Methods

Study design

Our study was based on an analysis of the treatment results of 353 patients with secondary diffuse peritonitis. Patients with pancreatic necrosis and impaired mesenteric circulation were excluded from the study due to the peculiarities of pathogenesis and difficulties in predicting the further development of the disease. The clinical characteristics of study subjects are shown in Table 1.

<table>
<thead>
<tr>
<th>Parameter</th>
<th>Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Average age, years</td>
<td>55.8±9.1</td>
</tr>
<tr>
<td>Men/women ratio</td>
<td>1.2:1.0 (190:163)</td>
</tr>
<tr>
<td>Characteristics of peritonitis vs V.S. Savelyev</td>
<td>336 (95.5%)</td>
</tr>
<tr>
<td>Generalized</td>
<td>16 (4.5%)</td>
</tr>
<tr>
<td>The most widespread causes of peritonitis, n=353</td>
<td></td>
</tr>
<tr>
<td>Acute destructive appendicitis</td>
<td>122 (34.6%)</td>
</tr>
<tr>
<td>Perforations of gastric and duodenal ulcers</td>
<td>85 (24.1%)</td>
</tr>
<tr>
<td>Colon perforation of various origins</td>
<td>58 (16.4%)</td>
</tr>
<tr>
<td>Acute destructive cholecystitis</td>
<td>36 (10.2%)</td>
</tr>
<tr>
<td>Lethality structure, n=59</td>
<td></td>
</tr>
<tr>
<td>Purulent intoxication and/or sepsis</td>
<td>51 (87.9%)</td>
</tr>
<tr>
<td>Cancer intoxication</td>
<td>4 (6.9%)</td>
</tr>
<tr>
<td>Acute cardiac failure</td>
<td>3 (5.2%)</td>
</tr>
<tr>
<td>Total lethality</td>
<td>16.7%</td>
</tr>
</tbody>
</table>

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Table 2. Impact of the criteria included in the PPS system on the prognosis

<table>
<thead>
<tr>
<th>Criteria</th>
<th>p-values</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age &gt; 80 years</td>
<td>0.000317 (Mann-Whitney U Test)</td>
</tr>
<tr>
<td>Concomitant cancer</td>
<td>0.0585 (Chi-squared with Yates’s continuity correction)</td>
</tr>
<tr>
<td>Purulent exudate</td>
<td>0.0002 (Fisher’s test)</td>
</tr>
<tr>
<td>Fecal exudate</td>
<td>0.0450 (Chi-squared with Yates’s continuity correction)</td>
</tr>
<tr>
<td>gSOFA</td>
<td>0.0000 (Chi-squared with Yates’s continuity correction)</td>
</tr>
<tr>
<td>Sepsis</td>
<td>0.0000 (Chi-squared with Yates’s continuity correction)</td>
</tr>
<tr>
<td>Toxic shock</td>
<td>0.0000 (Chi-squared with Yates’s continuity correction)</td>
</tr>
</tbody>
</table>

Table 3. Peritonitis Prognosis System (PPS)

<table>
<thead>
<tr>
<th>Criteria</th>
<th>Score, points</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age, years</td>
<td>0</td>
</tr>
<tr>
<td>≤ 60</td>
<td>3</td>
</tr>
<tr>
<td>61-70</td>
<td>4</td>
</tr>
<tr>
<td>&gt; 70</td>
<td>5</td>
</tr>
<tr>
<td>Presence of a malignant neoplasm</td>
<td>2</td>
</tr>
<tr>
<td>Exudate</td>
<td>3</td>
</tr>
<tr>
<td>Purulent exudate</td>
<td>5</td>
</tr>
<tr>
<td>Fecal exudate</td>
<td>5</td>
</tr>
<tr>
<td>Systemic inflammatory response</td>
<td>5</td>
</tr>
<tr>
<td>Sepsis</td>
<td>5</td>
</tr>
<tr>
<td>Toxic shock</td>
<td>7</td>
</tr>
<tr>
<td>Organ failure not associated with peritonitis</td>
<td>4</td>
</tr>
</tbody>
</table>

Table 4. The probability of death vs. the score sensu Peritonitis Prognosis System

<table>
<thead>
<tr>
<th>PPS score</th>
<th>Patient number</th>
<th>Died</th>
<th>Lethality (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>0-3</td>
<td>208</td>
<td>1</td>
<td>0.48</td>
</tr>
<tr>
<td>4-6</td>
<td>59</td>
<td>9</td>
<td>15.25</td>
</tr>
<tr>
<td>7-10</td>
<td>51</td>
<td>21</td>
<td>41.20</td>
</tr>
<tr>
<td>11-14</td>
<td>28</td>
<td>21</td>
<td>75.00</td>
</tr>
<tr>
<td>15 and more</td>
<td>7</td>
<td>7</td>
<td>100.00</td>
</tr>
<tr>
<td>Total</td>
<td>353</td>
<td>59</td>
<td>16.70</td>
</tr>
</tbody>
</table>

We analyzed the sensitivity and specificity of the Mannheim Peritoneal Index (MPI) [25], the Sepsis Severity Assessment of the World Society of Emergency Surgery [WSES SSS] for patients with complicated intra-abdominal infections (CIAI: APACHE-II systems) [26], gSOFA scores, as well as the Peritonitis Prognosis System (PPS) developed by the authors of this article.

Sensu the recommendations for sepsis-III, sepsis is understood as a sharp change of two or more points in the assessment of general Sequential Organ Failure Assessment (gSOFA); whereas toxic shock is characterized as sepsis with persistent hypotension requiring vasopressors to maintain an average arterial pressure ≥ 65 mm Hg and persistent serum lactate content of > 2 mmol/L despite the adequate volume resuscitation [4].

**Data processing**

Statistical analysis was performed using the STATISTICA 8 software. A p-value less than 0.05 implied statistical significance, and the values of the probability of a predicted outcome (patient death) were given taking into account the 95% confidence interval [27].

To determine the statistical significance of the criteria effect on the outcome, we analyzed 40 clinical and laboratory parameters that could contribute to the outcome of peritonitis, using parametric and nonparametric methods of statistical analysis (Appendix 1). The principle of constructing the scale was based on the selection of parameters (indicated in Table 2) that were statistically significantly associated with death in patients with secondary peritonitis.

To compare the predictive value of such systems as MPI, APACHE II, WSES SSS and PPS, ROC (receiver operating characteristic) analysis was used with the construction of ROC curves for each system. The statistical difference in the predictive values of those systems was analyzed by comparing the area under the curve of each system. The positive predictive value (PPV) of an outcome, hereinafter referred to as predictive value, was also calculated: it is the proportion (probability) of unfavorable outcomes (death) among all patients with a high risk of death. The described indicators are relative values and are expressed as percentages.

**Results**

We revealed the prognostic factors significantly affecting (p<0.05) the outcome of secondary peritonitis (Table 1). As can be seen, the criteria of a patient age, the presence of a malignant neoplasm, a nature of the exudate, sepsis (toxic shock), as well as organ failure (gSOFA) not associated with the development of peritonitis were of the greatest importance in predicting the lethal outcome. These criteria, presented in Table 2, were statistically significantly associated with the lethal outcome of peritonitis (p<0.05).

Based on the obtained data, we came up with a system for predicting peritonitis (PPS) (Table 3). The scoring system was developed taking into account the clinically significant, in our opinion, influence on the ongoing process of secondary peritonitis, as well as considering the level of their statistical significance. Weights (points) for each criterion were selected empirically.

The likelihood of a peritonitis outcome (ranks), depending on the number of points sensu PPS criteria, is presented in Table 4.

![ROC curves](image-url)
To compare the predictive value of PPS versus the most widely used systems for predicting the outcome of peritonitis (MPI, APACHE II, WSES SSS), ROC analysis was used with the construction of ROC curves for each system. Statistical indicators and curves obtained as a result of the performed ROC analysis are shown in Table 5 and Figure 1.

The performed analysis suggested that PPS was the most sensitive of all tested systems (Se=92%). The greater sensitivity could be explained by a large number of true positive outcomes, that is, by a large number of deceased patients, for whom the mortality index exceeded the threshold level of 50%. Statistically significant differences in predictive values were obtained for PPS, WSES SSS, and APACHE II systems (p<0.05, Fisher’s F Test). The MPI system exhibited statistically similar results in terms of assessing mortality in patients with secondary peritonitis in this sample of patients. The predictive value was calculated according to the criteria of the high risk of a patient death, assuming the threshold values of 26 points for MPI; 11 points for PPS; ≥ 7 points for WSES SSS; and ≥ 14 points for APACHE II. Hence, the predictive values were 38.2% for MPI, 83.3% for PPS, 74.3%, for WSES SSS, and 50.9% for APACHE II. Thus, the likelihood of unfavorable outcomes among all patients at high risk of death was determined most accurately by PPS and WSES SSS.

The obtained results can be explained by the inclusion of the criteria related to mortality in the group of patients with secondary peritonitis in a thanatogenetic and pathophysiologic manner – specifically, the criteria directly leading to the development of abdominal infection, e.g., the purulent and fecal character of the exudate, sequential organ failure, sepsis, and, as a consequence, cerebral edema, brain herniation syndrome and clinical death of a patient. As a result of the conducted analysis, the most accurate system, in terms of predicting mortality in a patient with secondary diffuse peritonitis, was the PPS. This was also confirmed by the AUC parameter (area under the ROC curve), had maximum values for PPS (0.942) and the minimum values for APACHE II (0.840). Such high predictive values of PPS and MPI could be explained by high sensitivities of these prediction systems specifically for the peritonitis. WE established that WSES SSS was less specific, since it was designed for urgent surgical diseases, without taking into account the peculiarities of the secondary peritonitis course. The general clinical APACHE II system was originally developed to assess the severity of the condition of intensive care patients and therefore was less focused specifically on peritonitis. For this reason, it demonstrated low specificity.

**Discussion**

Diffuse peritonitis is a major contributor to non-trauma mortality in all urgent care settings and the second leading cause of sepsis in critically ill patients [28-30]. Moreover, even in developed countries, the number of patients with peritonitis does not tend to decline, and their mortality rates remain high [31]. Mortality in diffuse peritonitis varies from 26 to 60%, reaching a level of 90-93% with the development of abdominal sepsis and toxic shock syndrome [32, 33].

The prognosis of peritonitis outcome depends on a combination of several factors: the degree of severity of a patient state, resource capabilities of a patient, specific factors of the disease, and diagnostic and therapeutic interventions. The distribution of patients among different risk groups helps predicting the outcome of the illness, selecting patients for intensive care, and identifying the surgical risk, thereby contributing to the choice of surgical treatment tactics – for example, the semi-open method, programmed relaparotomy or, in exceptional cases, damage control tactics.

One of the leading risk factors for peritonitis is preoperative sepsis with associated organ dysfunctions and toxic shock [34]. Other factors include a patient’s adaptive characteristics, concomitant diseases, especially those entailing organ failure, immunosuppression, and anatomical disorders associated with the disease [22, 33].

The results of our study suggested that abdominal sepsis and multiple organ dysfunction syndrome (associated or not associated with peritonitis) were the most important risk factors of a patient’s death caused by the peritonitis.

Of course, when choosing the tactics of treating a patient with peritonitis, it is necessary to take into account the degree of pathological changes in the abdominal cavity (fibrin deposition, the state of intestinal loops, and degree of their dilatation). However, the inclusion of these parameters or systems in prediction of the peritonitis outcome is not frequent. The good example supporting this statement is represented by inclusion of the Abdominal Cavity Index [6] in the system for predicting the outcome of peritonitis: it is very problematic to use in practice, especially because the contribution of each symptom (except for the nature of exudate) is quite small.

A low score of systems for predicting the outcome of peritonitis does not guarantee a favorable outcome of the disease. Technical and tactical errors in the treatment of patients may lead to severe complications. For example, in the analyzed group of patients, a 53-year-old woman (on admission, APACHE II of 6 points, MPI of 22 points, PPS of 3 points) with a perforated duodenal ulcer has developed suture failure and progression of peritonitis on the second day after suturing the ulcer. The patient underwent repeated suturing of the ulcerative defect, drainage of the abdominal cavity, and management according to the program of sanitation relaparotomies (4 relaparotomies in total). The death occurred due to the onset of tertiary peritonitis and purulent intoxication.

We would also like to admit that our results are based on the treatment outcomes at just one clinic; hence they require further independent external verification.

**Conclusions**

1. APACHE II, MPI, WSES SSS and PPS systems can be considered reliable in terms of mortality prognosis in patients with peritonitis and could be used as alternatives.

2. The greatest accuracy (94%) and predictive value (83.3%) in assessing the likelihood of a lethal outcome in patients with secondary diffuse peritonitis was established for the Peritonitis Prognosis System. The Mannheim’s Peritoneal Index also exhibited high predictive accuracy (over 90%).

**Funding**

The study was carried out in accordance with the research plan of the Medical Institute at the Peoples’ Friendship University of Russia.

**Conflict of interest**

The authors declare that they have no conflicts of interest.
Ethical approval

The study was approved by the Ethics Committee at the Peoples’ Friendship University of Russia. Ethical endorsement applied to all procedures performed in research involving humans, to the ethical standards of the institutional and national research committee, and to the 1964 Declaration of Helsinki and its later amendments or comparable ethical standards. For this type of study, formal informed consent was not required.

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Appendix 1. Studied clinical and laboratory indicators of the lethality risk in a patient with peritonitis

Gender, age, time since the onset of the disease, the source of peritonitis, the presence of a malignancy, the presence of organ failure (associated or not associated with peritonitis), localized or diffuse or generalized peritonitis, stomach rumble, the nature of the exudate in the abdominal cavity and its volume, the presence or absence of toxic shock, leukocytes in the blood, hemoglobin, hematocrit, red blood cells, body temperature, average arterial pressure, systolic arterial pressure, heart rate, respiratory rate, oxygenation (A-aDO2 or PaO2), mm Hg, sodium, potassium, serum creatinine, Glasgow Coma Scale (eye opening, motor response, speech), urea nitrogen, serum albumin, glucose, bilirubin, assessment of the acid-base status (pH/pCO2), urine erythrocytes and protein, procalcitonin, C-reactive protein, presepsin, central venous pressure, capillaroscopy data, results of ultrasound examination of the abdominal and pleural cavities.