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***Retrospective Cohort Study***

**Comprehensively evaluate the short outcome of small bowel obstruction: A novel medical-economic score system**

Xu WX *et al.* Comprehensive medical-economic score system for SBO

Wei-Xuan Xu, Qi-Hong Zhong, Yong Cai, Can-Hong Zhan, Shuai Chen, Hui Wang, Peng-Sheng Tu, Wen-Xuan Chen, Xian-Qiang Chen, Jun-Rong Zhang

**Wei-Xuan Xu, Qi-Hong Zhong, Yong Cai, Can-Hong Zhan,** Shengli Clinical Medical College, Fujian Medical University, Fuzhou 350122, Fujian Province, China

**Shuai Chen, Hui Wang, Peng-Sheng Tu, Wen-Xuan Chen, Xian-Qiang Chen, Jun-Rong Zhang,** Department of General Surgery (Emergency Surgery), Fujian Medical University Union Hospital, Fuzhou 350001, Fujian Province, China

**Author contributions:** Xu WX designed and performed the research and drafted the manuscript; Zhang JR designed the research and supervised and reviewed the report; Chen XQ supervised the report and provided funding acquisition; Zhong QH, Cai Y, Zhan CH, designed the research and contributed to the analysis; Chen WX, Chen S, Wang H, Tu PS collected data and provided methodology

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**Corresponding author: Jun-Rong Zhang, MD, Attending Doctor, Chief Doctor, Surgeon, Surgical Oncologist,** Department of General Surgery (Emergency Surgery), Fujian Medical University Union Hospital, No. 29 Xinquan Road, Fuzhou 350001, Fujian Province, China. junrongzhang@fjmu.edu.cn

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**Abstract**

BACKGROUND

Small bowel obstruction (SBO) still imposes a substantial burden on the health care system. Traditional evaluation systems for SBO outcomes only focus on a single element. The comprehensive evaluation of outcomes for patients with SBO remains poorly studied. Early intensive clinical care would effectively improve the short-term outcomes for SBO, however, the full spectrum of the potential risk status regarding the high complication-cost burden is undetermined.

AIM

We aim to construct a novel system for the evaluation of SBO outcomes and the identification of potential risk status.

METHODS

Patients who were diagnosed with SBO were enrolled and stratified into the simple SBO (SiBO) group and the strangulated SBO (StBO) group. A principal component (PC) analysis was applied for data simplification and the extraction of patient characteristics, followed by separation of the high PC score group and the low PC score group. We identified independent risk status on admission *via* a binary logistic regression and then constructed predictive models for worsened management outcomes. Receiver operating characteristic curves were drawn, and the areas under the curve (AUCs) were calculated to assess the effectiveness of the predictive models.

RESULTS

Of the 281 patients, 45 patients (16.0%) were found to have StBO, whereas 236 patients (84.0%) had SiBO. Regarding standardized length of stay (LOS), total hospital cost and the presence of severe adverse events (SAEs), a novel principal component was extracted (PC score = 0.429 × LOS + 0.444 × total hospital cost + 0.291 × SAE). In the multivariate analysis, risk statuses related to poor results for SiBO patients, including a low lymphocyte to monocyte ratio (OR = 0.656), radiological features of a lack of small bowel feces signs (OR = 0.316) and mural thickening (OR = 1.338), were identified as risk factors. For the StBO group, higher BUN levels (OR = 1.478) and lower lymphocytes levels (OR = 0.071) were observed. The AUCs of the predictive models for poor outcomes were 0.715 (95%CI: 0.635-0.795) and 0.874 (95%CI: 0.762-0.986) for SiBO and StBO stratification, respectively.

CONCLUSION

The novel PC indicator provided a comprehensive scoring system for evaluating SBO outcomes on the foundation of complication-cost burden. According to the relative risk factors, early tailored intervention would improve the short-term outcomes.

**Key Words:** Principal component analysis; Small bowel obstruction; Outcome evaluation system; Risk factors; Intensive clinical care; Radiomics

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**Core Tip:** A novel outcome indicator based on the standardized length of stay, total hospital cost and the presence of severe adverse events provided a comprehensive system for evaluating small bowel obstruction (SBO) outcomes. Furthermore, risk statuses associated with poor results were identified; specifically, for simple SBO patients, a low lymphocyte to monocyte ratio, as well as radiological features of a lack of small bowel feces signs and mural thickening, should be noticeable. For the strangulated SBO group, higher blood urea nitrogen levels and lower lymphocytes levels were recognized. Accordingly, early clinical intensive care was applicable for outcome improvement.

**INTRODUCTION**

Small bowel obstructions (SBO) result in over 300000 hospitalizations *per* year in the United States[1]. With the increasing public health burden, the average cost for SBOs ranges from $30000-$38000 individually, and the total cost for SBOs is estimated to be approximately 9-11.4 billion dollars[2,3]. Recently, the short outcomes of SBO were evaluated by using in-hospital mortality, major complications and the length of hospital stay[3-6]. There is still lack of an integrative medical-economic system to evaluate the overall outcomes for SBO, even though previous studies have confirmed the relationship between worse outcomes and higher hospital costs[7,8]. Furthermore, the question of how to comprehensively evaluate outcomes for patients with SBO remains uncharted.

Principal component analysis (PCA) is commonly used for dimension reduction[9,10], linear correlation resolution and data simplification. By summarizing and maximizing the information encoding a set of outcome variables, a novel principal component for evaluating the clinical and economic effects on SBO is available. For SBO, patients’ statuses on admission, including longer pain duration, acute kidney injury and malnutrition, were found to be closely correlated with severe adverse events (SAEs), based on previous studies[3,5,7,11]. However, the risk factors for the integrative scoring system, including clinical and economic adverse events, have not been extensively evaluated. The method of how to fully evaluate the potential risk status regarding the high complication-cost burden is urgently needed.

As an urgent life-threatening problem, the physical status of strangulated SBO is considerably deteriorating[12-14]. To control this confounding factor[15,16] and to further identify the risk admission status, we divided patients into a simple bowel obstruction group and a strangulated bowel obstruction group for the stratification analysis. We also constructed a novel indicator combining standardized SAEs, length of stay (LOS) and total hospital cost for defining outcomes of SBO. Furthermore, we established a representative model to distinguish high-risk statuses for both the simple small bowel obstruction (SiBO) and strangulated small bowel obstruction (StBO) groups to guide clinical intensive care for SBO.

**MATERIALS AND METHODS**

***Patient population***

From October 2016 to February 2021, 479 patients diagnosed with intestinal obstructions at Fujian Medical University Union Hospital were included in the study. After excluding 180 cases with large bowel obstructions, 4 cases with missing computed tomography (CT) images and 13 cases with incomplete clinical data, 281 patients were recruited for the final study (shown in Figure 1). The following stratification was made according to the pathological confirmation of intestinal ischemia: A simple bowel obstruction (SiBO, *n* = 236) group and a strangulated bowel obstruction (StBO, *n* = 45) group. For patients without acute peritonitis, conservative treatment was applied. Once patients with highly suspect of bowel ischemia or failure to conservative treatment, laparoscopy as well as laparotomy was adopted for SBO patients according to different intrabdominal pressures (shown in Table 1). The study protocol was approved by the Institutional Review Board of Fujian Medical University Union Hospital (Approval No. 2021YF005-02), and all of the patients provided written informed consent for the procedure.

***CT findings***

All of the patients with suspected SBO underwent CT scans before receiving treatment. The features of the CT scans that were recorded in this study were separated into mesenteric fluid, ascites, spiral signs, concentric circle signs, small bowel feces signs and edema of the bowel wall categories[17-20]. All of the CT scan images were cross-reviewed and evaluated by two senior general surgeons (Chen XQ and Zhang JR, and both surgeons had abundant experience in abdominal emergency surgery. The definitions of CT characteristics are shown in Supplementary Figure 1 and supplied in Supplementary Table 1[21-25].

***Clinical characteristics and laboratory tests***

Baseline demographics consisted of sex, age, body mass index (BMI), comorbidity, temperature, pain duration and history of abdominal pain. Biochemical parameters, including white blood cell count, neutrophil percentage, lymphocyte concentration, monocyte concentration, hemoglobin concentration, platelet concentration, albumin, alanine aminotransferase, aspartate aminotransferase (AST), calcium concentration, chloride concentration, potassium concentration, sodium concentration, blood urea nitrogen (BUN), serum creatinine, glucose, prothrombin time (PT), activated partial thromboplastin time (APTT), D-dimer (DDI) and fibrinogen, were collected within 24 h of admission. Combinations of inflammatory parameters, such as the neutrophil to lymphocyte ratio and lymphocyte to monocyte ratio (LMR), were calculated and recorded accordingly.

***Outcome definition***

Posttreatment outcomes were both clinically and economically evaluated.

Postoperative complications were defined as any deviation from the normal postoperative course during the index admission for SBO treatment, which was guided by the European Perioperative Clinical Outcome definitions[7,26]. The severity of complications was graded according to the Clavien-Dindo (CD) system[27], which is a validated classification system that categorizes complication severity based on the level of required treatment. Grade I was defined as complications without the need for pharmacological treatment or surgical, endoscopic and radiological interventions, as well as only minor interventions such as vomiting; grade II was defined as complications requiring pharmacological or other treatments, such as blood transfusions and total parenteral nutrition; grade III was defined as complications requiring surgical interventions or other interventional treatments; grade IV was defined as life-threatening complications, including central nervous system, cardiac and pulmonary complications, as well as renal failure and those interventions requiring intensive care unit (ICU) management; and grade V was defined as death. CD grade I to grade III were classified as non-SAE, and CD grade IV to grade V were classified as SAE.

The LOS was defined as the number of days from admission to discharge. Total hospital cost was defined as the total expenditure for medical resource utilization during hospitalizations, which included fees for operations (materials and occupancy of the operating room), medications, radiology, laboratory tests, microbiology tests, ward stay, ICU days, feeding and blood products[28].

***PCA***

PCA was used to achieve data simplification by expressing multivariate outcome indicators with fewer dimensions. With standardized LOS, total hospital cost and the presence of SAEs, a novel principal component was extracted: PC score = 0.429 × LOS + 0.444 × total hospital cost + 0.291 × SAE. Furthermore, the patient population was classified in the following manner according to the quartile PC score: The low PC score group (below the 75% quartile) and the high PC score group (in the upper 75% quartile). This analysis was performed in R V.4.1.3 (R Foundation for Statistical Programming, Vienna, Austria) by using the psych packages.

***Statistical analysis***

Categorical variables were compared by using the chi-square test or Fisher’s exact test between the two groups. Data are presented as the mean ± standard deviation or median for continuous variables. Independent t tests or Kruskal-Wallis tests were applied according to the characteristics of the variables. The association of admission status with higher PC scores was evaluated by using univariate logistic regression and summarized with an odds ratio (OR) and 95% confidence interval (CI). After setting the variables with a significance level of *P* < 0.05 and variance inflation factors < 5, a multivariate logistic regression with “binomial” method was performed, and independent risk factors were determined. We extracted the following risk score formulas based on these independent risk factors: Risk score 1 (RS1) = [0.291 × (bowel wall thickness) - 1.150 × (small bowel feces sign) - 0.421 × (LMR)] and RS2 = [-2.632 × (lymphocyte concentration) + 0.391 × (BUN concentration)] for the SiBO group and StBO group, respectively. Receiver operating characteristic curves and the area under the curve were calculated to assess the accuracy of the models. All of the statistical analyses were performed in R Version.4.1.3. The statistical methods of this study were reviewed by Yin YR.

**RESULTS**

***Outcome analysis***

For 281 patients with SBO who were included in this study, posttreatment outcomes were evaluated by LOS, total hospital cost and the presence of SAEs. *Via* the univariate analysis, admission risk status, including lower LMR (*P* = 0.005), higher BUN concentration (*P* = 0.022), higher glucose concentration (*P* = 0.007) and higher DDI concentration (*P* = 0.001), was significantly associated with higher hospital costs. Patients with SAE had lower levels of lymphocyte concentration (*P* = 0.003), higher levels of AST (*P* = 0.027), higher levels of potassium (*P* < 0.000), higher levels of BUN (*P* < 0.000), higher levels of serum creatinine (*P* < 0.000) and coagulation and fibrinolysis disturbances, including longer PT (*P* = 0.001), APTT (*P* = 0.012) and higher levels of DDI (*P* < 0.000). Furthermore, at admission, lower LMR (*P* = 0.003), higher monocyte concentration (*P* = 0.003), lower hemoglobin concentration (*P* = 0.038), higher level of glucose (*P* = 0.049), higher level of DDI (*P* = 0.004) and abnormal electrolyte and metabolic changes, such as lower calcium concentration (*P* = 0.042), lower chloride concentration (*P* = 0.003) and lower sodium concentration (*P* = 0.043), were closely related to a longer LOS (Figure 2 and Supplementary Table 2).

***PCA***

After maximizing the possible information and variation of the above-mentioned outcome indicators, including total hospital cost, LOS and SAEs, data simplification was performed. *Via* PCA, one principal component was extracted (Supplementary Figure 2). The PC score was calculated according to weights given to each outcome indicator: PC score = 0.429 × LOS + 0.444 × total hospital cost + 0.291 × SAE (Figure 1).

Of the 281 patients with SBO who were included in this study, 45 patients (16.0%) were found to have StBO, whereas 236 patients (84.0%) were found to have SiBO. The low PC score group (< 75% quartile) and high PC score group (> 75% quartile) were identified according to the quartile PC score. For both the SiBO and StBO groups, no significant difference was observed between the two PC score groups for sex, age, BMI, comorbidity status, temperature or history of abdominal operation (all *P* values > 0.05, Table 1). For patients with SiBO, a higher PC score was significantly related to longer pain duration (*P* < 0.000), higher monocyte concentration (*P* = 0.011), lower LMR (*P* = 0.002), lower hemoglobin concentration (*P* = 0.016), lower platelet count (*P* = 0.002) and low level of chloride (*P* = 0.015). Through the univariate analysis of radiological characteristics, we determined that a lack of small bowel feces signs and mural thickening were risk factors for a high PC score. In contrast, in the StBO group, low levels of lymphocytes (*P* = 0.020), high levels of AST (*P* = 0.022), high levels of BUN (*P* = 0.002) and coagulation and fibrinolysis disturbances, including abnormal DDI concentrations (*P* = 0.024), PTs (*P* = 0.004) and APTTs (*P* = 0.012), were significantly associated with higher PC scores. None of the risk radiological characteristics were observed in this stratification.

***Univariate and multivariate analyses of risk statuses***

*Via* the univariate analysis of the admission clinical-laboratory features, we determined potential risk status, including longer pain duration (*P* = 0.048), higher monocyte concentration (*P* = 0.003), lower LMR (*P* = 0.006), lower hemoglobin concentration (*P* = 0.033), lower platelet count (*P* = 0.036) and low level of chloride (*P* = 0.031), as well as radiological characteristics of mural thickening (*P* = 0.033) and lack of small bowel feces sign (*P* = 0.006), for high PC scores in the SiBO stratification. *Via* the multivariate analysis, independent risk factors consisting of radiological findings of small bowel feces sign (OR = 0.316), mural thickening (OR = 1.338) and LMR (OR = 0.656) were identified (all *P* values < 0.05, Table 2 and Figure 3). For StBO stratification, low levels of lymphocytes (*P* = 0.038), high levels of AST (*P* = 0.027), longer PTs (*P* = 0.015), high levels of BUN (*P* = 0.004) and creatinine (*P* = 0.022) seemed to be related to high PC scores. Finally, we found that only lymphocytes (OR = 0.071) and BUN (OR=1.478) were independent risk factors for high PC scores (all *P* values < 0.05, Table 2 and Figure 3).

Based on the regression coefficient for each factor, we calculated risk scores and built prediction models for worse outcomes: RS1 = [0.291 × (bowel wall thickness) - 1.150 × (small bowel feces sign) - 0.421 × (LMR)] for the SiBO group and RS2 = [-2.632 × (lymphocyte concentration) + 0.391 × (BUN concentration)] for the StBO group. Furthermore, receiver operating characteristic curves were drawn with areas under the curve of 0.715 (95%CI: 0.635-0.795) and 0.874 (95%CI: 0.762-0.986) for the SiBO and StBO stratifications, respectively (Figure 4).

**DISCUSSION**

Given that approximately 9-11.4 billion dollars are the costs *per* year in the United States, SBO still imposes a substantial burden on the health care system[2]. In contrast to the traditional evaluation systems that only focus on a single element[3-6], in this study, the standardized LOS, total hospital cost and the presence of SAEs were considered as integrative systems to evaluate the clinical-economic outcomes of SBO *via* PCA[9]. Previous studies have confirmed the close relationship between patients’ statuses on admission (including longer pain duration, acute kidney injury and malnutrition) and adverse outcomes, which provides a potential target for improving outcomes[3,5,7,11]. Commonly, severe statuses, including severe inflammatory reactions, electrolyte disturbances and hemostatic abnormalities, tend to occur in strangulated bowel obstruction[22]. Following the formula that assigned weights to each component, we determined PC score = 0.429 × LOS + 0.444 × total hospital cost + 0.291 × SAE; thus, the posttreatment outcome of SBO could be calculated and precisely evaluated (Figure 1).

For people with SiBO, only low LMR is observed, as radiological features (such as a lack of small bowel feces signs and mural thickening) were independent risk factors for high PC scores *via* the multivariate analysis. The area under the curve (AUC) of the predictive model based on the comprehensive scores for SiBO was 0.715 (95%CI: 0.635-0.795). As acute intestinal failure accompanies the obstructive bowel[26], when mechanical obstruction develops, the bowel lumen dilates along with the accumulation of air and intestinal fluid; thus, enteric stasis initiates bacterial proliferation with the intestinal gas produced by the fermentation of ingested food[22]. Conversely, when obstruction is incomplete or mild, the lasting bowel absorptive function can allow for fluid reabsorption across the bowel wall, thus leading to the small bowel feces sign as an independent protective factor for SBO[18,29]. Furthermore, progressive bowel dilation accompanied by compromised venous reflux increases intramural tension, which causes mural edema, secondary intestinal absorptive dysfunction and the loss of mucosal integrity (both functionally and physically)[22,30]. Similarly, as a potential effect on decreasing mural edema, the use of gastrografin challenge has been identified as the standardized management for SBO[31,32]. Moreover, in this study, the LMR was much lower in the high PC group, which may be due to the immune system becoming weakened as a result of the underlying malnutrition, as well as an excessive compensatory anti-inflammatory response[33-37].

Once SiBO deteriorated into StBO, the risk factors were dynamically changed. None of the radiological characteristics were found to be related to the outcomes. In particular, coagulation and fibrinolysis disturbances (including abnormal DDI, PT and APTT), kidney injury (such as increasing BUN and creatinine levels) and relevant lymphocytes were confirmed as being risk factors. Finally, only BUN and lower lymphocyte counts were identified as being independent risk factors for high PC. Partially due to the impaired mucosal barriers[22,38], lactic acid from intestinal anaerobic glycolysis gradually accumulates, which adversely deteriorates renal function with increasing levels of BUN in the peripheral blood[39]. Similarly, it is difficult to correct conventional enteral interventions and intestinal mucosal malnutrition due to the weakened immune status[33,40], which may explain why a lower level of lymphocytes is a risk factor for poorer outcomes. The predictive model for StBO yielded an AUC of 0.874 (95%CI: 0.762-0.986), which provided an excellent differentiating ability.

There were a few limitations to the present study. Primarily, this was a retrospective study conducted in a single center. In addition, the sample size of the initial models was relatively small. However, in both group (SiBO or StBO) the patients evaluated were consecutively enrolled and this could reproduce a real-world situation. Adequately powered and well-designed studies are required to confirm these findings and to establish causality.

**CONCLUSION**

The novel PC indicator provided a comprehensive scoring system for evaluating SBO outcomes on the foundation of complication-cost burden. According to the relative risk factors, early tailored intervention would improve the short-term outcomes.

**ARTICLE HIGHLIGHTS**

***Research background***

Small bowel obstruction (SBO) still imposes a substantial burden on the health care system. Traditional evaluation systems for SBO outcomes only focus on a single element. There is still lack of an integrative medical-economic system to evaluate the overall outcomes for SBO. Moreover, patients’ statuses on admission, including longer pain duration, acute kidney injury and malnutrition, were found to be closely correlated with severe adverse events (SAEs). However, the risk factors for the integrative scoring system, including clinical and economic adverse events, have not been extensively evaluated.

***Research motivation***

SBO still imposes a substantial burden on the health care system. Traditional evaluation systems for SBO outcomes only focus on a single element. The comprehensive evaluation of outcomes for patients with SBO remains poorly studied. Early intensive clinical care would effectively improve the short-term outcomes for SBO, however, the full spectrum of the potential risk status regarding the high complication-cost burden is undetermined.

***Research objectives***

In this study, we aim to construct a novel indicator combining standardized SAEs, length of stay (LOS) and total hospital cost for defining outcomes of SBO. Furthermore, we established a representative model for distinguishing high-risk statuses on admission for the simple SBO (SiBO) or strangulated SBO (StBO) groups. Given that SBO still imposes a substantial burden on the health care system, we believe our findings will provide a new insight for comprehensively evaluation outcomes of SBO as well as a guideline for early intervention.

***Research methods***

In this study, we evaluated posttreatment outcomes of SBO both clinically and economically. Principal component analysis (PCA) was used to achieve data simplification by expressing multivariate outcome indicators with fewer dimensions. By summarizing and maximizing the information encoding in standardized LOS, total hospital cost and the presence of SAEs, a novel principal component was extracted: PC score = 0.429 × LOS + 0.444 × total hospital cost + 0.291 × SAE. Furthermore, the patient population was classified in the following manner according to the quartile PC score: The low PC score group (below the 75% quartile) and the high PC score group (in the upper 75% quartile).

***Research results***

In this study, a novel outcome indicator based on the standardized LOS, total hospital cost and the presence of SAEs provided a comprehensive system for evaluating SBO outcomes (PC score = 0.429 × LOS + 0.444 × total hospital cost + 0.291 × SAE). Furthermore, risk statuses associated with poor results were identified; specifically, for SiBO patients, a low LMR, as well as radiological features of a lack of small bowel feces signs and mural thickening, should be noticeable. For the StBO group, higher blood urea nitrogen levels and lower lymphocytes levels were recognized. Accordingly, early clinical intensive care was applicable for outcome improvement. In the future, adequately powered and well-designed studies are required to confirm these findings and to establish causality.

***Research conclusions***

In this study, PCA was innovatively used for dimension reduction, linear correlation resolution and data simplification. Furthermore, a novel comprehensive system for the evaluation of SBO outcomes was constructed and the potential risk status associated with poor results were identified.

***Research perspectives***

Large-scale and prospective studies are going to be designed to confirm these findings and to establish causality.

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**Footnotes**

**Institutional review board statement:** The study protocol was approved by the Institutional Review Board of Fujian Medical University Union Hospital (Approval No. 2021YF005-02).

**Informed consent statement:** Patients were not required to give informed consent to the study because the analysis used anonymous clinical data that were obtained after each patient agreed to treatment by written consent.

**Conflict-of-interest statement:** All authors read and approved the final manuscript and declared no conflicts of interest.

**Data sharing statement:** The original anonymous dataset is available on request from the corresponding author at junrongzhang@fjmu.edu.cn.

**STROBE statement:** The authors have read the STROBE Statement—checklist of items, and the manuscript was prepared and revised according to the STROBE Statement—checklist of items.

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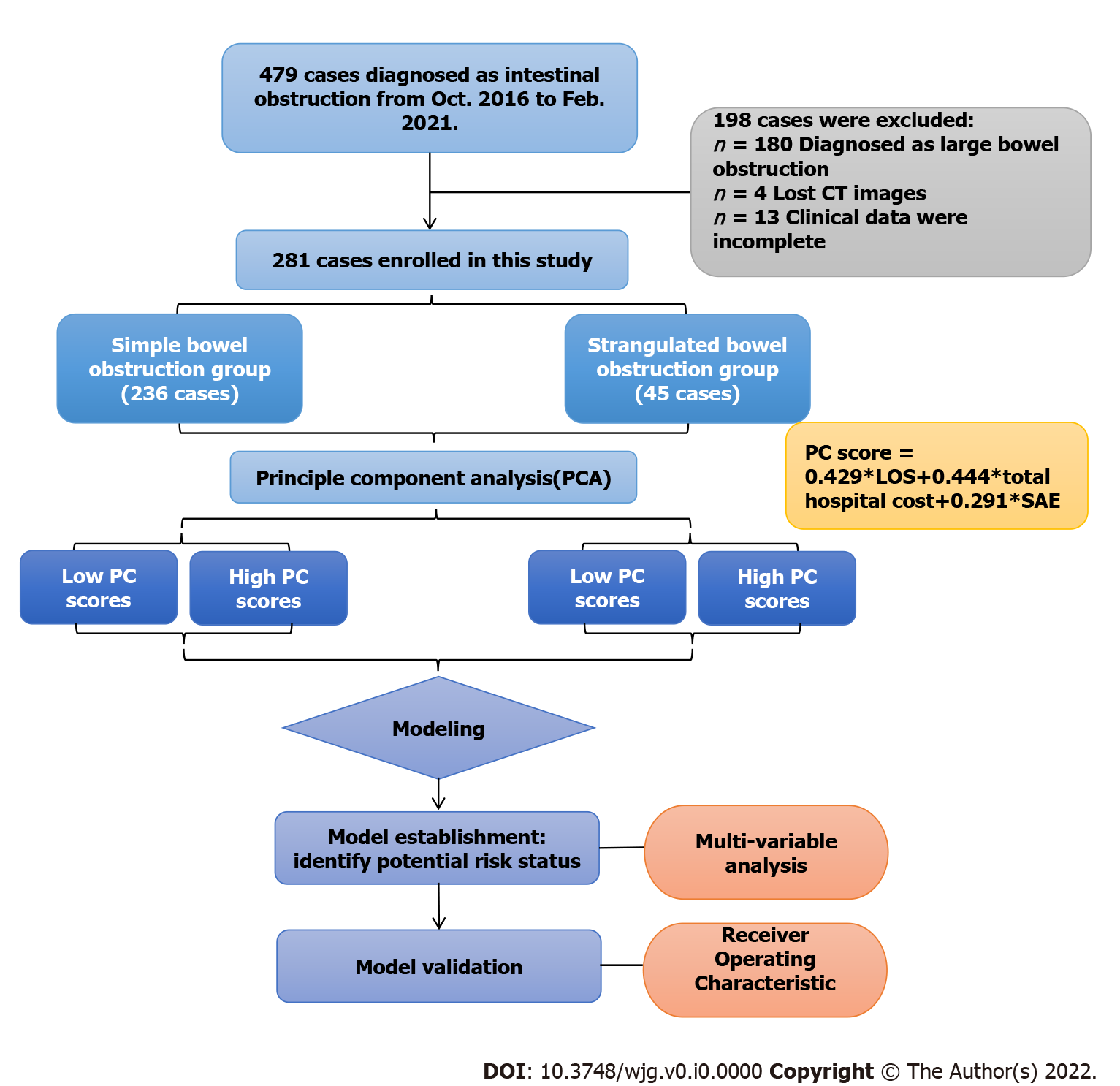
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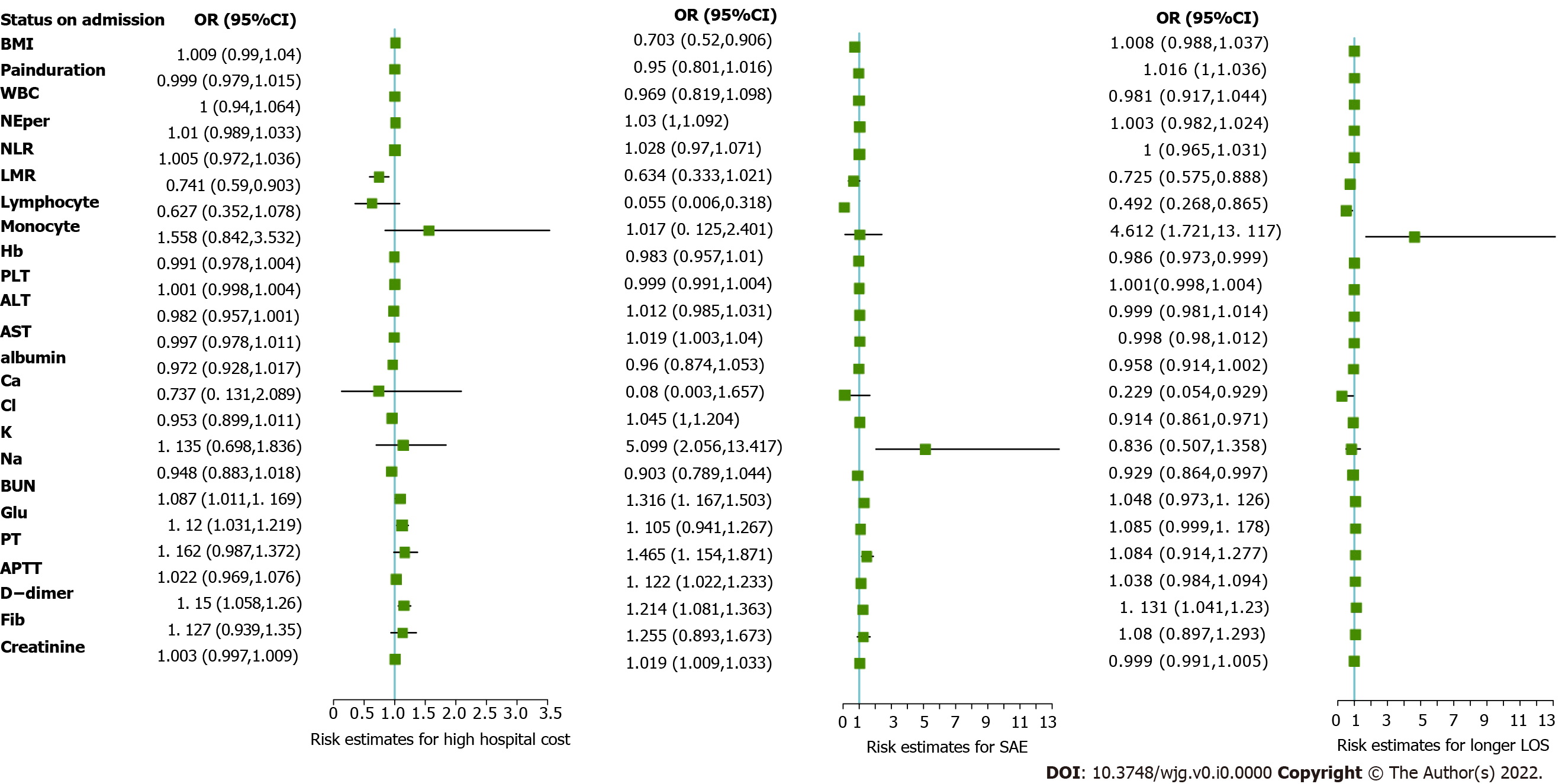
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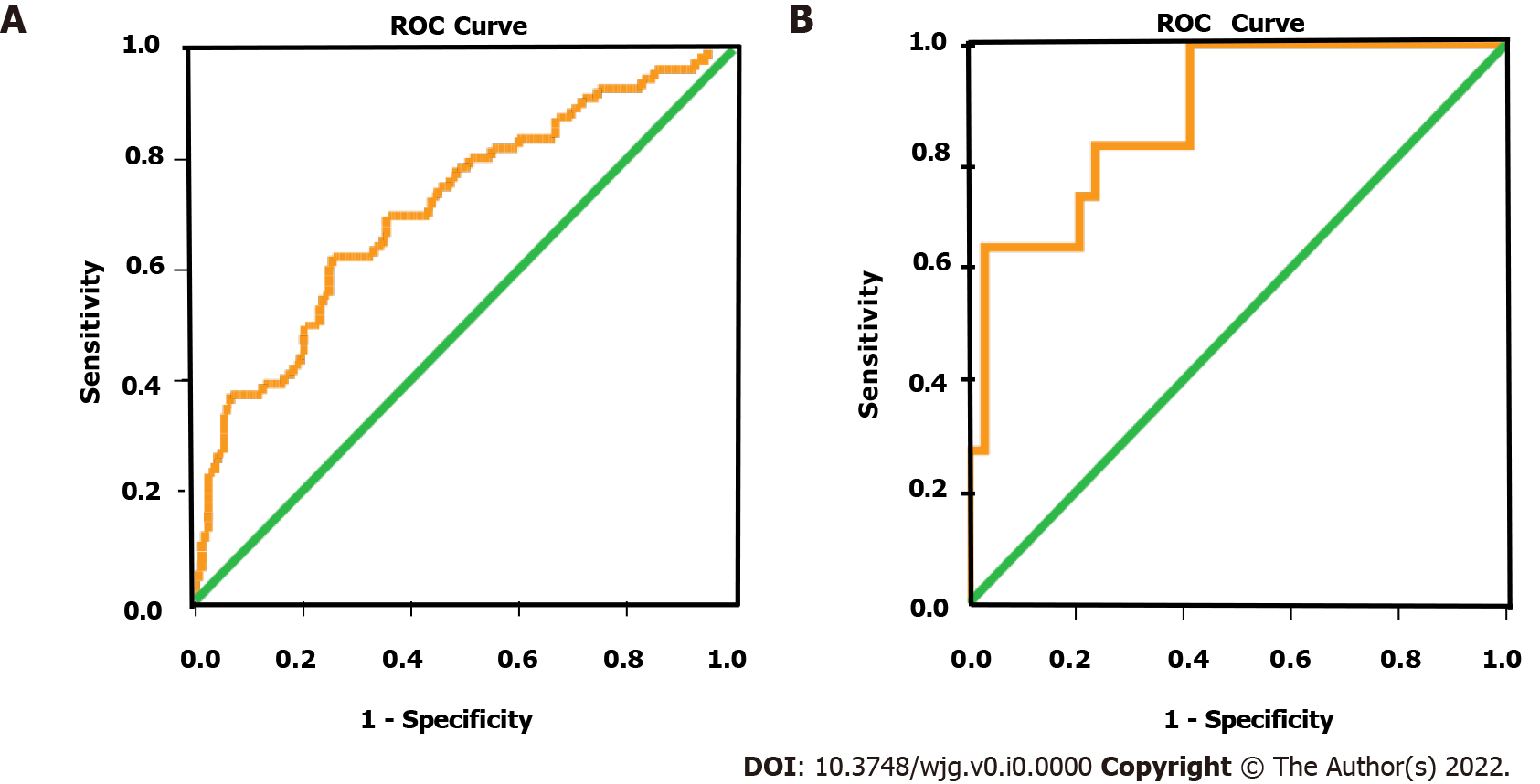
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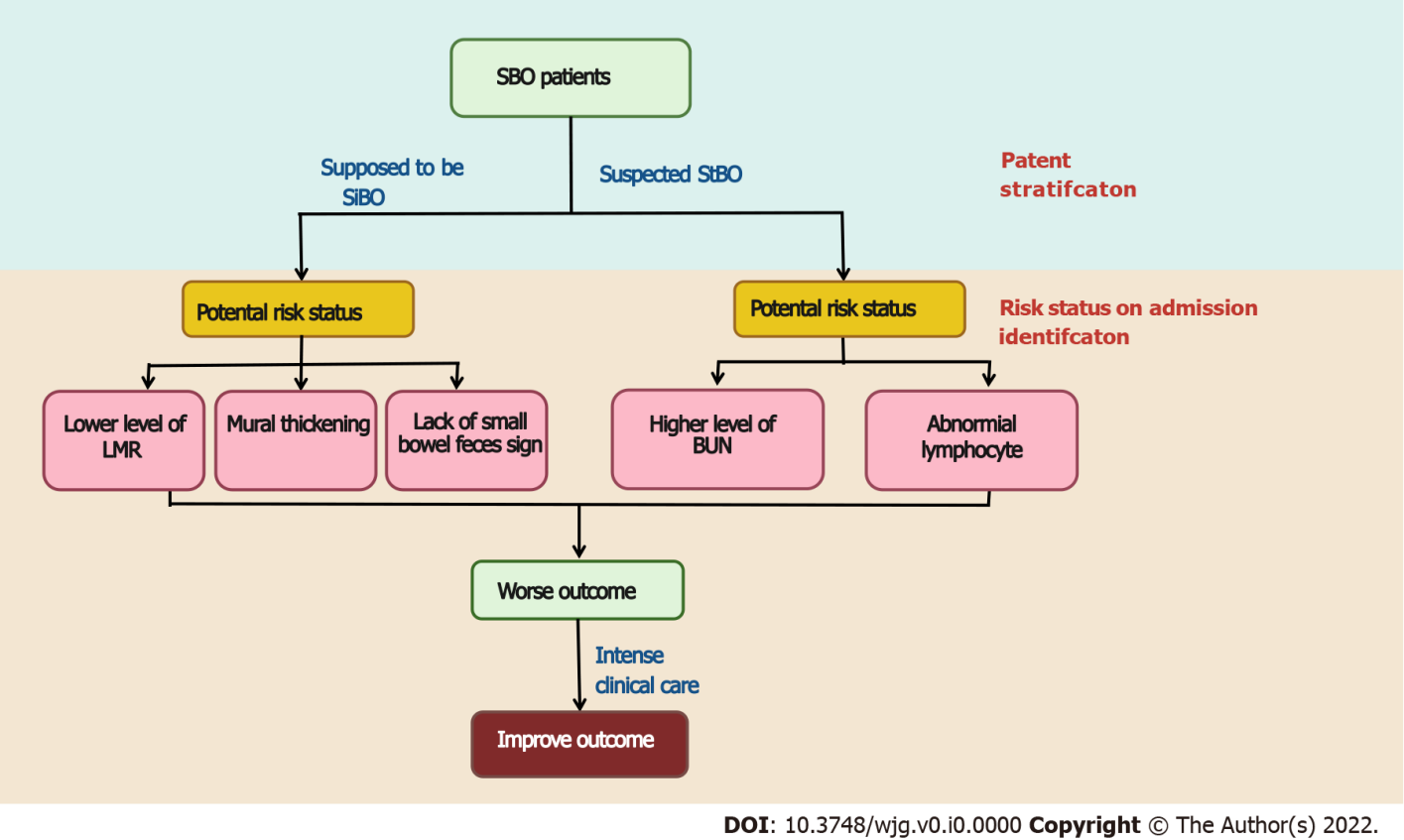
**Figure 1 Workflow of this study.** CT: Computed tomography; PC: Principal component; LOS: Length of stay; SAE: Severe adverse event.

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**Figure 2 Risk factors for worse outcome of small bowel obstruction.** Risk estimates for high hospital cost; Risk estimates for severe adverse event; Risk estimates for longer length of stay. OR: Odds ratio; CI: Confidence interval; BMI: Body mass index; WBC: White blood cell; NE%: Neutrophil percentage; NLR: Neutrophil to lymphocyte ratio; LMR: Lymphocyte to monocyte ratio; Hb: Hemoglobin; PLT: Platelet, ALT: Alanine aminotransferase; AST: Aspartate aminotransferase; Ca: Calcium; Cl: Chloride; K: Potassiun; Na: Sodium; BUN: Blood urea nitrogen; Glu: Glucose; PT: Prothrombin time; APTT: Activated partial thromboplastin time; DDI: D-dimer; Fib: Fibrinogen; SAE: Severe adverse event; LOS: Length of stay.

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**Figure 3 Receiver operating characteristic curve for high principal component score prediction.** The areas under the curve were 0.715 (95%CI: 0.635-0.795), 0.874 (95%CI: 0.762-0.986), respectively. A: Receiver operating characteristic curve of simple small bowel obstruction group for high principal component score prediction. B: Receiver operating characteristic curve of strangulated small bowel obstruction group for high principal component score prediction. ROC: Receiver operating characteristic.



**Figure 4 Proposal early clinical intensive care for small bowel obstruction patients on admission.** SBO: Small bowel obstruction; SiBO: simple small bowel obstruction; StBO: Strangulated small bowel obstruction; LMR: Lymphocyte to monocyte ratio; BUN: Blood urea nitrogen.

**Table 1 Compared the clinical and laboratory characteristics of the patients with low or high principal component score**

|  |  |  |  |  |  |  |
| --- | --- | --- | --- | --- | --- | --- |
| **Characteristics** | **Simple obstruction (*n* = 236)** | | ***P* value** | **Strangulated obstruction (*n* = 45)** | | ***P* value** |
| **Low PC score** | **High PC score** | **Low PC score** | **High PC score** |
| Baseline data |  |  |  |  |  |  |
| Gender, *n* (%) |  |  | 1.0001 |  |  | 0.4212 |
| male | 117 (69.2%) | 39 (69.6%) |  | 18 (52.9%) | 8 (72.7%) |  |
| female | 52 (30.8%) | 17 (30.4%) |  | 16 (47.1%) | 3 (27.3%) |  |
| Age, (yr) | 60 (47, 69) | 65 (53, 71) | 0.081 | 63 (52.25, 70.00) | (61.0, 71.5) | 0.321 |
| BMI, (kg/m2) | 20.70 (18.83, 22.98) | 20.94 (18.21, 22.65) | 0.196 | 20.20 (18.16, 22.00) | 18.75 (17.72, 19.81) | 0.228 |
| Comorbidity, *n* (%) |  |  | 0.2451 |  |  | 1.0002 |
| none | 128 (75.7%) | 38 (67.9%) |  | 27 (79.4%) | 9 (81.8%) |  |
| yes | 41 (24.3%) | 18 (32.1%) |  | 7 (20.6%) | 2 (18.2%) |  |
| Pain duration, (d) | 2 (1, 5) | 6 (3, 12.5) | < 0.000 | 2.00 (1.00, 3.75) | 2.0 (1.0,4.0) | 0.989 |
| History of abdominal operation, *n* (%) |  |  | 0.4711 |  |  | 0.6032 |
| none | 43 (25.4%) | 17 (30.4%) |  | 11 (32.4%) | 2 (18.2%) |  |
| yes | 126 (74.6) | 39 (69.6%) |  | 23 (67.6%) | 9 (81.8%) |  |
| Temperature, (Degrees celsius) | 36.6 (36.5, 36.8) | 36.6 (36.5, 36.8) | 0.401 | 36.6 (36.5, 36.8) | 36.60 (36.50, 36.75) | 0.956 |
| CT characteristics |  |  |  |  |  |  |
| Mesenteric fluid, (%) |  |  | 0.430 |  |  | 0.9852 |
| none | 32 (18.9%) | 8 (14.3%) |  | 1 (2.9%) | 1 (9.1%) |  |
| yes | 137 (81.1%) | 48 (85.7%) |  | 33 (97.1%) | 10 (90.9%) |  |
| Ascites, (%) |  |  | 0.849 |  |  | 1.0002 |
| none | 58 (34.3%) | 20 (35.7%) |  | 4 (11.8%) | 1 (9.1%) |  |
| yes | 111 (65.7%) | 36 (64.3%) |  | 30 (88.2%) | 10 (90.9%) |  |
| Spiral signs, (%) |  |  | 0.6122 |  |  | 0.4362 |
| none | 151 (89.3%) | 52 (92.9%) |  | 22 (64.7%) | 5 (45.5%) |  |
| yes | 18 (10.7%) | 4 (7.1%) |  | 12 (35.3%) | 6 (54.5%) |  |
| Concentric circle sign, (%) |  |  | 0.1322 |  |  | 0.7452 |
| none | 164 (97.0%) | 51 (91.1%) |  | 31 (91.2%) | 11 (100%) |  |
| yes | 5 (3.0%) | 5 (8.9%) |  | 3 (8.8%) | 0 (0%) |  |
| Small bowel feces sign, (%) |  |  | 0.006 |  |  | 1.0002 |
| none | 70 (41.4%) | 35 (62.5%) |  | 17 (50.0%) | 5 (45.5%) |  |
| yes | 99 (58.6%) | 21 (37.5%) |  | 17 (50.0%) | 5 (54.5%) |  |
| Mural (thickness (median) | 3.28 (2.30, 3.75) | 3.63 (2.97, 4.53) | 0.002 | 3.51 (3.16, 4.12) | 3.42 (2.67, 4.07) | 0.634 |
| Laboratory data |  | | | | | |
| WBC, (109/L) | 6.770 (4.89, 9.52) | 7.345 (4.87, 11.18) | 0.387 | 8.70 (5.89, 12.23) | 6.83 (10.01, 18.25) | 0.384 |
| NE% | 75.50 (65.9, 83.3) | 77.45 (68.5, 84.03) | 0.422 | 83.60 (69.05, 86.90) | 77.50 (73.30, 90.25) | 0.853 |
| Lymphocyte, (109/L) | 1.01 (0.74, 1.42) | 0.94 (0.64, 1.34) | 0.240 | 0.96 (0.65, 1.34) | 0.60 (0.42, 0.76) | 0.020 |
| Monocyte, (109/L) | 0.420 (0.30, 0.58) | 0.565 (0.34, 0.73) | 0.011 | 0.570 (0.407, 0.735) | 0.540 (0.33, 0.760) | 0.721 |
| NLR, (ratio) | 4.650 (3.03, 8.07) | 6.115 (3.74, 9.30) | 0.159 | 7.750 (4.085, 12.922) | 9.030 (4.990, 15.565) | 0.491 |
| LMR, (ratio) | 2.286 (1.67, 3.42) | 1.591 (1.13, 2.84) | 0.002 | 1.681 (2.131, 1.141) | 1.482 (0.957, 1.933) | 0.459 |
| Hb, (g/L) | 128.0 (115, 142) | 120.5 (108, 133) | 0.016 | 131.0 (110.0, 137.7) | 129.0 (120.5, 145.0) | 0.587 |
| PLT, (109/L) | 205.5 (162.50, 250.75) | 250.5 (180.25, 307.25) | 0.002 | 213 (163, 260) | 180.0 (152.0, 242.5) | 0.256 |
| Albumin, (g/L) | 35.9 (32.30, 40.45) | 36.1 (31.80, 39.45) | 0.403 | 34.6 (31.7, 39.6) | 37.1 (28.6, 42.0) | 0.977 |
| ALT, (U/L) | 16 (11, 24) | 16 (11, 22) | 0.727 | 15.00 (12.00, 21.75) | 15.00 (13.25, 27.75) | 0.612 |
| AST, (U/L) | 20 (16, 26) | 21 (17, 25.5) | 0.619 | 19.50 (17.00, 23.75) | 36.50 (20.75, 45.25) | 0.022 |
| Ca, (mmol/L) | 2.19 (2.04,2.32) | 2.15 (2.02,2.26) | 0.152 | 2.19 (2.09, 2.31) | 2.05 (1.95, 2.20) | 0.062 |
| Cl, (mmol/L) | 102.30 (100.0, 104.1) | 100.15 (96.85, 104.03) | 0.015 | 100.85 (98.13, 103.85) | 102.00 (101.35, 104.15) | 0.296 |
| K, (mmol/L) | 4.035 (3.78, 4.34) | 3.985 (3.74, 4.43) | 0.957 | 4.00 (3.56, 4.31) | 4.36 (3.62, 5.07) | 0.290 |
| Na, (mmol/L) | 138.40 (136.68, 140.48) | 138.15 (135.50, 141.23) | 0.533 | 138.05 (134.13, 140.30) | 135.60 (134.75, 137.75) | 0.334 |
| BUN, (mmol/L) | 5.5 (4.3, 7.2) | 5.4 (3.68, 8.23) | 0.872 | 6.45 (4.00, 8.57) | 10.6 (7.3, 15.3) | 0.002 |
| Glu, (mmol/L) | 6.78 (5.30, 8.67) | 6.59 (5.16, 9.51) | 0.515 | 8.165 (6.963, 9.300) | 8.66 (6.78, 11.04) | 0.428 |
| PT, (s) | 13.6 (13.1, 14.3) | 13.6 (13.28, 14.43) | 0.825 | 13.45 (12.90, 13.90) | 15.40 (14.20, 16.65) | 0.004 |
| APTT, (s) | 35.6 (33.3, 38.3) | 36.1 (34.0, 40.9) | 0.184 | 35.15 (32.18, 37.10) | 41.6 (36.1, 45.0) | 0.012 |
| DDI, (mg/L) | 1.45 (0.71, 2.52) | 1.94 (0.79, 4.67) | 0.151 | 1.64 (0.88, 3.50) | 5.75 (2.39, 6.72) | 0.024 |
| Fib, (g/L) | 3.49 (2.92, 4.37) | 3.78 (3.25, 4.59) | 0.150 | 3.69 (2.71, 4.60) | 3.89 (3.17, 4.82) | 0.548 |
| Creatinine, (umol/L) | 70.0 (56, 81) | 70.5 (54.75, 88.00) | 0.512 | 67 (57, 76) | 93 (80, 147) | 0.003 |
| Management |  |  | < 0.0002 |  |  | 0.2152 |
| conservative treatment | 155 (91.7%) | 17 (30.4%) |  | 1 (2.9%) | 0 (0%) |  |
| laparoscopy | 11 (6.5%) | 8 (14.3%) |  | 8 (23.5%) | 0 (0%) |  |
| laparotomy | 3 (1.8%) | 31 (55.4%) |  | 25 (73.5%) | 11 (100%) |  |
| CD, *n* (%) |  |  | < 0.0002 |  |  | < 0.0002 |
| Grade I | 141 (83.4%) | 20 (35.7%) |  | 5 (14.7%) | 0 (0%) |  |
| Grade II | 28 (16.6%) | 32 (57.1%) |  | 28 (82.4%) | 2 (18.2%) |  |
| Grade III | 0 (0%) | 1 (1.8%) |  | 1 (2.9%) | 0 (0%) |  |
| Grade IV | 0 (0%) | 3 (5.4%) |  | 0 (0%) | 9 (81.8%) |  |
| Grade V | 0 (0%) | 0 (0%) |  | 0 (0%) | 0 (0%) |  |
| SAE, *n* (%) |  |  | 0.0182 |  |  | < 0.0002 |
| none | 169 (100%) | 53 (94.6%) |  | 34 (100%) | 2 (18.2%) |  |
| yes | 0 (0%) | 3 (5.4%) |  | 0 (0%) | 9 (81.8%) |  |
| Fee, (¥) | 12070 (8830, 19935) | 54322 (41370, 74623) | < 0.000 | 51828 (33575, 66954) | 83553.0 (74146.0, 142409.5) | < 0.000 |
| Length of stay, (d) | 5 (4, 8) | 16 (13.75, 22.50) | < 0.000 | 14.00 (10.25, 17.00) | 28.0 (18.5, 35.5) | < 0.000 |

1were compared using the chi-square test.

2were adjusted p-values or Fisher’s exact test.

SiBO: Simple small bowel obstruction; StBO: Strangulated small bowel obstruction; PC score: Principle component score; BMI: Body mass index; WBC: White blood cell; NE%: Neutrophil percentage; NLR: Neutrophil to lymphocyte ratio; LMR: Lymphocyte to monocyte ratio; Hb: Hemoglobi; PLT: Platelet; ALT: Alanine aminotransferase, AST: Aspartate aminotransferase; Ca: Calcium; Cl: Chloride; K: Potassiun; Na: Sodium; BUN: Blood urea nitrogen; Glu: Glucose; PT: Prothrombin time; APTT: Activated partial thromboplastin time; DDI: D-dimer; Fib: Fibrinogen; CD: Clavien-Dindo; SAE: Severe adverse event.

**Table 2 Univariate and multivariate analyses of risk factors for high principal component score**

|  |  |  |  |  |  |  |  |  |
| --- | --- | --- | --- | --- | --- | --- | --- | --- |
| **Characteristics** | **Simple obstruction (*n* = 236)** | | | | **Strangulated obstruction (*n* = 45)** | | | |
| **Univariate** | | **Multivariate** | | **Univariate** | | **Multivariate** | |
| **OR (95%CI)** | ***P* value** | **OR (95%CI)** | ***P* value** | **OR (95%CI)** | ***P* value** | **OR (95%CI)** | ***P* value** |
| Pain duration, (d) | 1.019 (1.002,1.041) | 0.0481 |  |  |  |  |  |  |
| Small bowel feces sign, (+)/ (-) | 0.424 (0.225,0.783) | 0.0061 | 0.316 (0.158,0.612) | < 0.0001 |  |  |  |  |
| Mural thickening, (cm) | 2.119 (1.084,4.375) | 0.0331 | 1.338 (1.098,1.664) | 0.0031 |  |  |  |  |
| Lymphocyte, (109/L) |  |  |  |  | 0.097 (0.007,0.665) | 0.0381 | 0.071 (0.003,0.539) | 0.0331 |
| Monocyte, (109/L) | 5.472 (1.809,17.780) | 0.0031 |  |  |  |  |  |  |
| LMR, (ratio) | 0.708 (0.541,0.891) | 0.0061 | 0.656 (0.496,0.836) | 0.0011 |  |  |  |  |
| Hb, (g/L) | 0.983 (0.969,0.998) | 0.0331 |  |  |  |  |  |  |
| PLT, (109/L) | 1.003 (1.001,1.007) | 0.0361 |  |  |  |  |  |  |
| AST, (U/L) |  |  |  |  | 1.075 (1.018,1.156) | 0.0271 |  |  |
| Cl, (mmol/L) | 0.931 (0.871,0.993) | 0.0311 |  |  |  |  |  |  |
| BUN, (mmol/L) |  |  |  |  | 1.383 (1.133,1.786) | 0.0041 | 1.478 (1.169,2.061) | 0.0041 |
| PT, (s) |  |  |  |  | 1.568 (1.141,2.418) | 0.0151 |  |  |
| APTT, (s) |  |  |  |  | 1.109 (0.999,1.264) | 0.076 |  |  |
| DDI, (mg/L) |  |  |  |  | 1.196 (1.006,1.513) | 0.067 |  |  |
| Creatinine, (umol/L) |  |  |  |  | 1.034 (1.011,1.071) | 0.0221 |  |  |

1indicates that the parameters have statistical difference (*P* < 0.05).

SiBO: Simple small bowel obstruction; StBO: Strangulated small bowel obstruction; OR: Odds ratio; CI: Confidence interval; LMR: Lymphocyte to monocyte ratio; Hb: Hemoglobin; PLT: Platelet; AST: Aspartate aminotransferase; Cl: Chloride; BUN: Blood urea nitrogen; PT: Prothrombin time; APTT: Activated partial thromboplastin time; DDI: D-dimer.