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A new clinical severity score for the management of acute small bowel obstruction in predicting bowel ischemia: a cohort study

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Geneva, May 3rd, 2022

Object: Highlights

- Small bowel obstruction (SBO) is a common hospital admission diagnosis but remains a clinical challenge
- Identification of patients requiring emergency surgery and by extension, bowel resection is the main goal of surgical physicians
- A practical clinical severity score guiding patient's care
- Early surgical relief of the SBO increases the chance to avoid bowel resection
- Avoiding unnecessary surgery could reduce patients' morbidity

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Object: Data statement

Due to the sensitive nature of the questions asked in this study, survey respondents were assured raw data would remain confidential and would not be shared.

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Declarations of interest: none

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The results of this original article were presented at the Swiss Society for Surgery Annual meeting on the 2nd of June 2021

Abstract

Background: Small bowel obstruction (SBO) is a common hospital admission diagnosis.

Identification of patients who will require a surgical resection because of a non-viable small bowel remains a challenge. Through a prospective cohort study, we aimed to validate risk factors and scores for intestinal resection, and to develop a practical clinical score designed to guide surgical vs. conservative management.

Materials and Methods: All patients admitted for an acute SBO between 2004 and 2016 in our center were included. Patients were divided in three categories depending on the management: conservative, surgical with bowel resection and surgical without bowel resection.

The outcome variable was small bowel necrosis. Logistic regression models were used to identify the best predictors.

Results: Seven hundred and thirteen patients were included in this study, 492 in the development cohort and 221 in the validation cohort. Sixty-seven percent had surgery, of which 21% had small bowel resection. Thirty-three percent were treated conservatively. Eight variables were identified with a strong association with small bowel resection: age ≥ 70 years, first episode of SBO, no bowel movement for ≥ 3 days, abdominal guarding, C-reactive protein (CRP) ≥ 50 , and three abdominal Computer Tomography scanner (CT-scan) signs: small bowel transition point, lack of small bowel contrast enhancement, and the presence of >500 mL of intra-abdominal fluid. Sensitivity and specificity of this score were 65% and 88%, respectively, and the area under the curve was 0.84 (95% CI 0.80 – 0.89).

Conclusion: We developed and validated a practical clinical severity score designed to tailor management of patients presenting with an SBO.

Trial registration: clinicaltrials.gov Identifier: NCT01125280

1. Introduction

Small bowel obstruction (SBO) is defined as a total or partial blockage of the intestinal lumen. It can result from incarcerated hernia, volvulus, intussusception, primary or secondary tumors, peritoneal carcinomatosis, strictures (secondary to inflammatory disorders, fibrosis, radiotherapy), gallstones or foreign bodies [1, 2]. However, 60 to 70% of SBO results from adhesions, either as a single fibrous band or as multiple bands all over the abdominal cavity, and therapeutic strategy in those situations is more challenging [3, 4]. In most cases, adhesions are the consequence of previous abdominal surgery, but they can also occur spontaneously in patients without any surgical history [5]. Post-operative adhesions incidence is related to the site of surgery, with a range from 0.05% after a Caesarean-section (C-section), 1% after an appendectomy and up to 10% after colorectal surgery [4, 6, 7]. In the absence of strangulation, evident ischemia or peritonitis, a significant proportion of patients can be treated conservatively with bowel rest, gastric decompression with a nasogastric (NG) tube, and intra-venous (IV) fluid. Water-soluble contrast imaging is now routinely used during the non-operative trial. A portion of cases will present a complete obstruction with bowel strangulation, compromising blood flow and resulting in bowel ischemia, commonly followed by perforation. Those patients frequently require a bowel resection when surgery is not performed within the first 6 hours [8]. The identification of patients who are at risk or who will develop small bowel ischemia is essential, as it has been associated with a mortality rate of 30% compared to 3% without ischemia [9]. Potential complications of delayed management include aspiration pneumonia, sepsis and acute renal injury, with an increased 1-year mortality rate [10]. On the other hand, the identification of patients at low risk of bowel ischemia is equally important since it allows to avoid unnecessary surgery.

We previously designed and investigated a 6-item SBO/small bowel resection risk score which included: abdominal pain ≥ 4 days, abdominal guarding, leukocytosis (≥ 10 G/ml), C-

reactive protein (CRP) ≥ 75 , ≥ 500 mL intra-abdominal free fluid and/or lack of small bowel contrast enhancement on Computer Tomography scanner (CT-scan) [11, 12]. Our initial score was able to predict small bowel resection with an area under the curve (AUC) receiver operating characteristic (ROC) curve of 0.87. However, no validation in a larger dataset was yet available.

The primary objective of the present study was to validate our previous score developed in a cohort of 221 patients (2004-2007, referred as the historical cohort) in a cohort of 492 patients (2008-2016, referred as the current cohort).

When applied to the current cohort, the AUC was 0.78 (95% CI 0.72 – 0.84). Because it was substantially less than the originally reported value of 0.87, we thus decided to improve the existing score using the most recent dataset.

Therefore, the second objective was to provide an improved severity score for SBO with a simplified version of the score to allow its use in the absence of CRP (not routinely performed in most of the institutions) and IV contrast (usually not given to patients with an elevated creatinine).

2. Methods

2.1 Patients

The current cohort was used to validate the previous score [11]. Additionally, it has been used as the development cohort for the new clinical severity score. All adult patients (n= 824) presenting to our hospital with the International Classification of Disease-10 code K56.5 (Intestinal adhesions with obstruction) between 2008 and 2016 were considered for inclusion in the study. The SBO episode occurring during the study period was defined as the index episode. Patients with an active abdominal oncological process (primary tumors, metastases, peritoneal carcinomatosis), an SBO occurring in the immediate postoperative period (i.e. 6

weeks), an incarcerated hernia, post radiotherapy or with large bowel obstructions (n=209) were excluded from our analysis. Patients without a CT-scan, with a CT-scan without contrast or with missing data were also excluded (n=123) (similar exclusion criteria were used for the validation cohort [11, 12]). A total of 492 patients were included in the current cohort.

Patient's assessment was standardized and included: a clinical exam, a complete blood count (CBC) and a basic metabolic panel (BMP), CRP, lactate, and a CT-scan of the abdomen. The conservative versus surgical management was left at the discretion of the surgeon on call. Our previously published score was prospectively calculated for each patient, recorded as an additional information, but decision making was left at the on-call surgeon's discretion.

Patients who did not respond to conservative management within 48 hours were managed surgically. The use of laparoscopy versus open surgery, and the decision to resect/not to resect small bowel was based on the clinical judgement by the surgeon on-call. Patients were categorized in three groups depending on the management: conservative, surgical with bowel resection and surgical without bowel resection.

We used as an external validation cohort for our newly developed score, patients from the historical cohort who presented at our hospital with an SBO between 2004 and 2007 (n=221), as previously reported by our group [11, 12].

2.2 Study variables

Patient's clinical and surgical characteristics were prospectively collected in patient's electronic chart. Post-operative complications were classified according to Clavien-Dindo [13].

2.3 Score validation and development

Our initial 6-item score was calculated as previously described [11] and applied to the current cohort. ROC curve and AUC were computed. We defined 0.03 as the maximal AUC difference for validation.

For the elaboration of a new score, the 2008-2016 dataset (current cohort) was used. In addition, we used the 2004-2007 dataset (historical cohort) for external validation. We converted all variables to categorical or binary variables for score development when appropriate. Odds ratios (OR) for small bowel resection was calculated for each variable independently. Variables with an OR ≥ 2 and a p-value < 0.2 were included in the multivariate predictive model. Risk factors with a p-value > 0.05 were removed one by one, until all retained predictors were statistically significant. To assess the performance of the model, we obtained a ROC curve and the corresponding AUC. The AUC represents the probability that a randomly chosen patient who required a resection would have a higher score than a randomly selected patient who was treated conservatively. A value of 0.5 corresponds to lack of discrimination and 1 to perfect discrimination.

Risk scores were calculated by adding 1 point per risk factor present. We used multiple permutations to derive two simplified versions of the original score and retained those with AUC ≥ 0.8 and including the most clinically meaningful variables.

We arbitrarily determine 70% as the individual predicted risk of resection. For external validation, we replicated analyses (AUC, proportion at risk $\geq 70\%$, sensitivity, and specificity) for the three scores in the 2004-2007 database (historical cohort).

2.4 Statistical analysis

Continuous variables were compared between groups using a student t-test with Welch's correction or a Mann-Whitney t-test when appropriate. Categorical variables were compared by using a χ^2 test. A P-value <0.05 was considered statistically significant.

Recurrence rate was assessed and time to recurrence was defined as the duration between the date of index SBO episode admission to the recurrence date of admission. For patients who did not experience a recurrence, we used the date of last follow-up or death. A univariate analysis was performed in order to identify possible variable associated with recurrence. A multivariate regression analysis was then performed using the Cox regression model. All statistical analyses were performed using IBM SPSS Statistics version 25 (IBM corp.; Armonk, NY).

This work has been reported in line with the STROCCS criteria [14] , Supplemental Digital Content 1, <http://links.lww.com/JS9/A153>.

3. Results

We first constituted the current cohort and assessed the previous score [11]. Between 2008 and 2016, 492 patients presented a bowel obstruction and met the inclusion criteria. Three hundred and forty-five patients (70.1%) had surgery, of which 103 (20.9%) had small bowel resection. The proportion of cases managed surgically did not change over the study period (Supplementary Figure 1, Supplemental Digital Content 2, <http://links.lww.com/JS9/A154>). Hundred and forty-seven patients (29.9%) were treated conservatively. Demographics, clinical, biological and radiological characteristics of included patients are described in Table 1. There were 294 (59.8%) female and 198 (40.2%) males overall, and no gender difference was observed between groups. Patients in the surgical groups were older and had higher BMI. Patients admitted for an initial SBO episode were more likely to be in the surgical group. Pain duration, absence of bowel movement, fever, and vomiting were not significantly different

between groups. Patients treated surgically had higher leukocytosis, CRP, and CT signs of transition point, lack of small bowel enhancement, and intra-abdominal fluid.

Variables related to patients treated surgically are reported in Table 2. Among all patients operated (n=345), 204 (59.5%) were taken into the operating room in the first 24 hours and 29.9% (n=103) needed a resection because of bowel ischemia. Of the 16 patients who had a lack of small bowel contrast enhancement on CT-scan, but did not require surgical resection, the mean delay to surgery was 10.2 ± 5.8 hours compared to 12.2 ± 8.9 hours for patients with the same CT-scan findings and who needed a resection ($p=0.42$). A higher percentage of patients were operated by laparoscopy in the non-resected group in comparison with the group with bowel resection (31.4% vs 12.6%, $p<0.001$), and the conversion rate to laparotomy was significantly higher in the group with bowel resection (84.6% vs 32.9%, $p<0.001$). No association was observed between delay to surgery and complication rate and severity.

3.1 Postoperative outcomes and SBO recurrence

Length of hospital stay was longer in the surgical group compared to the conservative group. Patients treated conservatively had more subsequent SBO recurrent episodes (Table 3). One hundred twenty-two patients (35.4%) presented post-operative complications (Table 3). Complications were more frequent and more severe in the group requiring small bowel resection ($p=0.018$). SBO recurrence occurred more frequently in the conservative group (31.3%), followed by the surgical group without bowel resection (14.5%), and finally the surgical group with bowel resection (7.8%) (Table 3). Potential variables affecting SBO recurrence were assessed using a uni-/multivariate Cox proportional hazard model (Supplementary Table 1, Supplemental Digital Content 3, <http://links.lww.com/JS9/A155>). The presence of a previous SBO episode was significantly associated with an increased risk of recurrent SBO (adjusted Hazard Ratio (HR) 2.2, 95%CI: 1.4-3.4, $p<0.001$). Patient treated

surgically for their index SBO episode was protective against further recurrence (adjusted HR 0.5, 95%CI: 0.3-0.7, $p<0.001$).

3.2 Performance of our previous SBO clinical score

We applied our previously published clinical score [11] (developed in the 2004-2007 SBO cohort) to our new 2008-2016 cohort. The AUC was 0.78 (95% CI 0.72 – 0.84). Although the previous score function well in the new dataset, the AUC was substantially less than the originally reported value of 0.87. We thus decided to improve the existing score using the most recent dataset.

3.3 Elaboration of a new clinical severity score to predict the risk of small bowel resection

For the construction of the new clinical score, only the conservative group ($n=147$) and the resection group ($n=103$) were used (total $n=250$). The external validation dataset (historical cohort) included 33 patients who required resection and 82 patients who were treated conservatively, all with complete data.

We considered 13 variables measured at patient intake (Table 1). Four variables were not significantly associated with small bowel resection in univariate analysis: gender, duration of pain ($p=0.52$ for conservative versus surgery with resection, data not shown), fever, and vomiting. Of the four age groups, the two older strata were at higher risk of resection, and therefore, were grouped to define a single risk group, age ≥ 70 years. Of the three strata of CRP, only the highest (≥ 50) was associated with resection, and here too the variable was dichotomized. Nine variables remained for the multivariate logistic regression model. In this analysis, the association between leukocytosis and resection became weak and statistically non-significant, and was therefore withdrawn from the model. The final full model included 8 variables (Table 4). The AUC of this logistic regression model was 0.863 (95% CI 0.816 –

0.910). The OR ranged from 2.36 to 8.10, which suggested that some risk factors were more important than others. Nevertheless, in the interest of usability, we established a risk score as a simple sum of the risk factors present in a given patient. The score had thus a possible range of 0 to 8 (Table 5). The risk of resection increased gradually with the score, from 0% at score 0 to 100% for scores 5 to 7. No patient had a score of 8. Among 58 patients with a score of 4, 41 (70.7%) required a resection and 17 (29.3%) had a resolution of symptoms without surgery, whereas among 66 patients with a score of 3, 24 (33.4%) required a resection. We proposed that patients with scores from 4 to 8 be considered at high risk of ischemia and therefore likely to require surgical resection. By this definition, 84 (33.6%) of the 250 patients would be considered at high risk. The AUC under the ROC curve of the 8-item score was 0.84 (95% CI 0.80 – 0.89) (Figure 1).

3.4 Elaboration of two alternative scores

Because not all patients can benefit from contrast injection during CT scan (contrast allergies, renal insufficiency), we considered a 7-item score that excluded the lack of contrast enhancement on CT-scan. Finally, in order to facilitate daily use of the score, we also developed a brief 4-item score that relied only on age, guarding, transition point and presence of 500 mL of free fluid on CT scan (Table 5). Both scores were associated with the risk of resection in a gradual manner. The 7-item score can be used similarly to the 8-item score, in that a score of 4 or greater defines the high-risk group. For the brief 4-item score, a score of 2 or higher defined the high-risk category.

3.5 Comparison of the three scores

Selected items, AUC, validation AUC, high-risk thresholds and sensibility-specificity are shown in Table 6. The discriminative ability was excellent for all three scores, although decreasing along with the reduction in items, as shown by the AUC under the ROC curves (Table 6, Figure 1). The AUCs were 0.844 (8-item score), 0.826 (7-item score) and 0.802 (4-item score) (Table 6). For all three scores, the loss of discrimination was less than 0.03 in the external validation dataset (i.e. -0.023, -0.015, and -0.028, respectively).

The proportion of patients identified as being at high risk of resection (namely $\geq 0.70\%$ risk of small bowel resection) went from one third for the 8-item score to one quarter for the 4-item score (Table 6). However, when the threshold of high risk was redefined as the average of 11 values between 0.60 and 0.80, the proportions of high-risk patients were similar for the three scores. This difference between the three scores was less pronounced in the external validation dataset.

Sensitivity decreased for the 7- and 4-item risk scores compared to the full 8-item score.

However, specificity showed an opposite trend (Table 6). In the external validation dataset, these values were comparable to those of the development dataset.

The distribution of the clinical severity score was assessed among the three groups (Table 7). Sixty-five percent of patients in the surgical group with bowel resection had a score of ≥ 4 in comparison to 21.2% and 11.6% in the surgical group without resection and the conservative group, respectively. Mean and median score followed a similar trend.

4. Discussion

In this study, we address a common dilemma for the on-call surgeon in the emergency room (ER) and on the surgical floor: “should I take this patient presenting an SBO to the operating room?”. We developed and validated an SBO score with two simplified versions to aid the surgeon in the daily clinical decision making. In this study, we observed a good performance

of our previous SBO score in a new dataset. However, the reduction of the AUC in this new dataset was too important, dropping from 0.87 to 0.78, in order to validate this score.

We therefore decided to refine our initial scoring system and designed a new score, which the following 8 items: age >70 years, first SBO episode, ≥ 4 days since the last BM, small bowel transition point on CT-scan, abdominal guarding, CRP $\geq 50\text{mg/l}$, non-enhanced small bowel on CT-scan, and CT-scan free fluid $\geq 500\text{ ml}$. Four criteria were maintained from the previous score and four new criteria were added. Interestingly, we did not find significant associations between small bowel resection rate and leukocytosis, abdominal pain duration, vomiting, creatinine, lactate levels, and platelet levels. These criteria were thus abandoned.

Additionally, we present two simplified version of the score: 7-item score for patients with acute kidney insufficiency (AKI) or contrast allergies (no CT contrast enhancement evaluation required) and one with only 4-items allowing a rapid clinical and radiological assessment without the need for laboratory values. All developed scores (the 8-, 7- and 4-item) had excellent discrimination ability to predict the need for bowel resection (and thus an operation) in the current cohort. We were able to validate all three scores internally and externally. We also developed the latter simplified score keeping in mind that CRP is not routinely available/measured in most hospitals across the globe. Patients with 5 points or more in the 7- and 8-item score, and 4 points in the 4-item score had a 100% chance risk of small bowel resection. These patients should therefore be taken to the OR for a surgical exploration. For patients with an intermediate score, namely 2 to 4 for the 8- and 7-item score, and 1 to 2 in the 4-item score, a conservative approach with watchful waiting may be attempted. For patients with a low score, namely 0 to 1 for the 8- and 7-item score, and 0 in the 4-item score, a conservative approach is likely to be successful.

The management of SBO has been well described and algorithm for diagnosis and treatment have been proposed [15]. In addition to the items included in our score, the presence of the

following factors warrants a surgical exploration: a persistent obstruction after a 72h observation period, and a NG drainage volume >500 ml/24h on the 3rd day [15]. Comparing the criteria identifying peritonitis and/or ischemia from international guidelines [15] to our own data, the following can be highlighted: we abandoned leukocytosis (i.e. leukocytes > 10,000/mm³) given its weak and statistically non-significant relation with bowel resection, we lowered the “CRP >75 mg/l” criteria to 50 mg/l since this threshold was already significantly associated with resection, and kept the “free intraperitoneal fluid >500 ml” as is. Despite the profusion of guidelines regarding the management of SBO, no consensus exists when it comes to the use of a uniform score to predict the need for a surgical intervention. An “a posteriori” classification of SBO exist and was designed to predict severity and outcomes. The results have been published by the American Association for the Surgery of Trauma [16]. It includes anatomical factors (whether an operation was needed an if a resection was performed), physiological factors (the presents of systemic inflammatory response syndrome (SIRS), sepsis or septic shock), and the Charlson comorbid index [17]. This score is complex and not designed to predict the need for resection. Therefore, our proposed score and its declinations fulfill a need for standardization in the management of SBO.

The formation of adhesions is the most frequent long-term complication of virtually any type of surgery, and account for a significant number of ER visit, morbidity and mortality in the context of abdominal surgery [18]. The challenge in the management of those patients is to identify which ones will benefit from a surgical management and which one who will not. Failure to recognize early enough the need for a surgical intervention can lead to bowel necrosis, sepsis, septic shock and death, especially in frail/older patients. On the other hand, performing unnecessary surgical explorations puts patients at risk of peri- and post-operative

complications and the development of new adhesions. An SBO score helps to standardize patient's presentation to the team and can guide the surgeon make the final clinical decision. Finding a reliable and accessible predictor for bowel ischemia is crucial. It should be specific to bowel ischemia and sensitive enough, in order to rule out other pathologies. A recent systematic review analyzed all serological human markers for bowel ischemia investigated over the last few years [19]. Common laboratory values such as lactate dehydrogenase, white blood cells count, base excess and creatine kinase did not demonstrate any strong association with bowel ischemia. More interestingly, lactate level was not predictive either for ischemia, which we confirmed in our data; lactate seems limited by being an unspecific marker for tissue hypoperfusion. Furthermore, it was demonstrated that bowel necrosis can occur even without a significant increase in lactate levels [20, 21]. We found and validated a role for CRP to be one of the predictors of bowel ischemia in two different cohorts, including a total of more than 700 patients. However, CRP remains an unspecific marker of inflammation and its use is very center-dependent, and thus probably not broadly applicable. Procalcitonin (PCT) is another marker of inflammation and bacterial infection [22] that has attracted attention for its ability to predict bowel ischemia [23-26]. While its measurement is becoming more and more routinely performed, the use of PCT should be encouraged as a useful marker for initial work-up for SBO.

In their systematic review, Derikx et al. proposed other interesting markers for bowel ischemia such as D-Dimer, endothelin-1, ischemia modified albumin, α -Glutathione S-transferase (α -GST) and Intestinal fatty acid binding protein (I-FABP) [19], the latter being the most promising as it is specific to enterocytes. However, the turnover for these tests is still too long to be applicable and cut-off values are to be determined.

Over the last decades, the surgical management of acute SBO has refined with the appearance of laparoscopy [27], and progressively let more space for semi-interventional with the introduction of water-soluble contrast imaging as a dual diagnostic and therapeutic method [28]. The use of water-soluble contrast imaging potentially allows to reduce the need for surgery in selected patients [29], although we did not observe that trend in our cohort. In our population, 70.1% of patients presenting an index SBO episode were treated surgically and among them, 29.9% had a bowel resection. Our operative management rate is higher than those previously reported in the literature, namely ranging from 22% to 51% [10, 28, 30-33]. This might be explained by the fact that we specifically focused on adhesion-related SBO episodes and excluded hernia-, cancer-, and radiation-related SBOs. This could also be explained by our policy favoring surgical management which was justified by the 50% reduction in the need for readmission and 60% reduction in recurrent SBO symptoms, as shown by us and others [10-12, 33]. This is especially true when a single adhesion causing recurring abdominal pain and obstruction is present [28]. The timing of surgery is another important parameter. A delay of 24 hours or more is usually associated with higher mortality and post-operative complications rate [34]. We could however not reproduce this observation, potentially due to our more aggressive management including a smaller proportion of patients managed conservatively.

Our study has limitations. This is a mono-centric design, which can lead to selection bias, limitations in the number of possible included patients, and conclusions that rely on local practices that are not necessarily transferable to other centers. We provided a simplified 4-item score which retained excellent discrimination ability (AUC of 0.80 in the development cohort and 0.77 in the validation cohort) in order to be used more easily in other centers. This simplified score is based on objective and universally available variables to predict the need for small bowel resection; which represents an objective outcome by itself, and which should

be similar and translatable to most centers. Another inherent limitation in all the developed predictive models and clinical scores is that they are founded on skewed datasets as clinicians already balance their management decisions based on available clinical variables (that may later be included into the score). Although the new score was not used to make clinical decisions per se, the attention directed to a part of the individual variables may have contributed to some degree of selection bias. Validation in pre-score era cohort somehow mitigated the risk of confounding. Another possible bias is that we excluded patients without CT-scan contrast injection. The simplified score did function relatively well without the “IV contrast” information and patients without IV contrast should be included in future studies. The use of CRP in the full score can also represent a limitation which was addressed by the development of the simplified score as mentioned above. Finally, the objectivity of “small bowel resection” as an outcome will remain conditional to the on-call surgeon clinical decision and dependent upon the timing of the operation. In other terms, a similar situation, a strangulation might or might not result in a resection, depending upon how early is the surgical exploration. We thought about using a criterion such as “intraoperative ischemic bowel loop presence/absence” but found that this was inconsistently reported and not a fully objective criterion either. It is also likely that this “gray” zone (unclear if resection is needed) does not represent the majority of the SBO cases. Alternatively, one could also argue that an operation might be warranted and beneficial, even if the need for resection is absent, particularly in non-resolving/smoldering clinical situations. In this regard, we believe that the score should remain an aid but not a substitute for clinical judgment.

In conclusion, we developed and validated a score to guide the clinical decision to whether or not a given patient should be taken to the operating room for an acute SBO episode. We provided and validated two additional versions of our SBO score to accommodate most

clinical situation. The decision to operate will always retain a subjective part based on the surgeon's experience and the clinical situation, however, our SBO score has proven helpful for standardized clinical decision, management, and outcomes evaluation.

Provenance and peer review

Not commissioned, externally peer-reviewed

ACCEPTED

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Figure 1. Receiver operating characteristic curves of the full 8-item score (solid line), the 7-item score (dotted line), and the brief 4-item score (dashed line).

Arrows point to thresholds for high risk (≥ 4 for the 8-item and 7-item scores, ≥ 2 for the 4-item score).

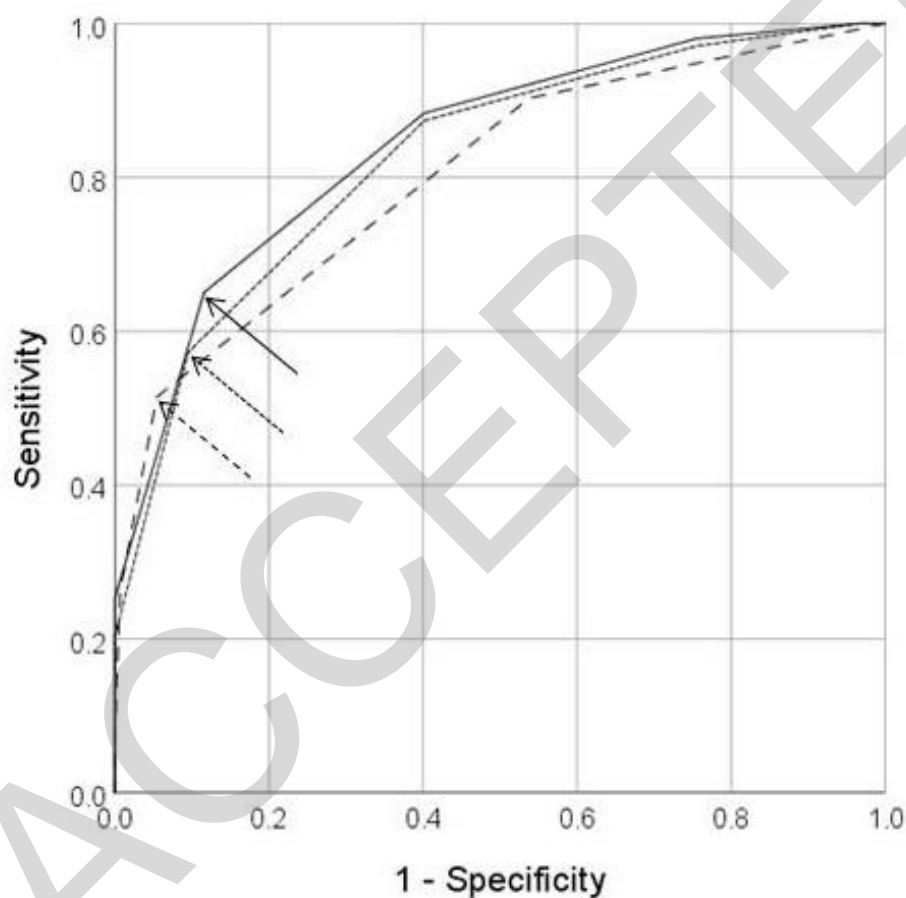


Table 1. Comparison of clinical, biological and radiological parameters of patients recorded during the first hospitalization for a SBO episode

	Surgical, no resection N=242	Surgical resection N=103	Conservative N=147	P-value [†]
Gender (%)				0.46
Female	147 (60.7)	65 (63.1)	82 (55.8)	
Male	95 (39.3)	38 (36.9)	65 (44.2)	
Age, years \pm SD	64.6 \pm 18.2	71.6 \pm 16.9	62.9 \pm 18.4	<0.001
Age, years (%)				<0.001
22-59	91 (37.6)	21 (20.4)	64 (43.4)	
60-69	44 (18.2)	13 (12.6)	24 (14.3)	
70-79	45 (18.6)	30 (29.1)	27 (18.4)	
80-96	62 (25.6)	39 (37.9)	32 (21.8)	
BMI, kg/m ² \pm SD	23.9 \pm 4.8	22.6 \pm 3.4	24.0 \pm 4.9	0.04
Previous SBO (%)				<0.001
Yes	38 (15.7)	17 (16.5)	56 (38.1)	
No	204 (84.3)	86 (83.2)	91 (61.9)	
Pain duration, days (%)				0.074
0-3	213 (88.0)	91 (88.3)	134 (91.2)	
≥ 4	29 (12.0)	12 (11.7)	13 (8.8)	
Absence of bowel movement, days	1.8 \pm 1.6	2.1 \pm 2.1	1.5 \pm 0.9	0.035
Absence of bowel movement, days (%)				0.083
0-2	208 (86.0)	84 (81.6)	134 (91.2)	
≥ 3	34 (14.0)	19 (18.4)	13 (8.8)	
Fever $>38^{\circ}\text{C}$ (%)				0.16
Yes	15 (6.2)	12 (11.7)	9 (6.1)	
No	227 (93.8)	91 (88.3)	138 (93.9)	
Vomiting (%)				0.53
Yes	185 (76.4)	78 (75.7)	105 (71.4)	
No	57 (23.6)	25 (24.3)	42 (28.6)	
Abdominal guarding (%)				<0.001
Yes	82 (33.9)	42 (40.8)	13 (8.8)	
No	160 (66.1)	61 (59.2)	134 (91.2)	
Leukocytosis, G/l \pm SD	11.2 \pm 3.9	12.3 \pm 5.3	10.8 \pm 6.6	0.056
Leukocytosis, >10 G/l (%)				0.050

Yes	145 (59.9)	67 (65.0)	74 (50.3)	
No	97 (40.1)	36 (35.0)	73 (49.7)	
CRP, mg/l \pm SD	33.0 \pm 69.1	43.5 \pm 67.4	23.5 \pm 40.1	0.041
CRP, mg/l (%)				0.035
0-9	138 (57.0)	54 (52.4)	82 (55.8)	
10-49	61 (25.2)	20 (19.4)	45 (30.6)	
50-470	43 (17.8)	29 (28.2)	20 (13.6)	
Creatinine, \pm SD	33.0 \pm 69.1	43.5 \pm 67.4	23.5 \pm 40.1	0.157
Lactates, \pm SD	1.7 \pm 0.9	2.0 \pm 1.4	1.8 \pm 2.0	0.536
Platelets, \pm SD	277.4 \pm 98.5	297.2 \pm 110.5	259.1 \pm 93.1	0.012
CT – transition point (%)				0.044
Yes	219 (90.5)	93 (90.3)	117 (79.6)	
No	23 (9.5)	10 (9.7)	30 (20.4)	
CT – contrast enhancement (%)				<0.001
Yes	226 (93.4)	79 (76.7)	144 (98.0)	
No	16 (6.6)	24 (23.3)	3 (2.0)	
CT – IAFF >500 mL (%)				<0.001
Yes	62 (25.6)	45 (43.7)	13 (8.8)	
No	180 (74.4)	58 (56.3)	134 (91.2)	

SBO: small bowel obstruction, CRP: C-reactive protein, IAFF: intra-abdominal free fluid.

† χ^2 test for binary or categorical variable (global p value), T-test for continuous variables

Table 2. Operative characteristics and data compared between the two surgical groups.

	Surgical, no resection N=242	Surgical resection N=103	P-value [†]
ASA (mean ± SD)	2.5 ± 0.7	2.6 ± 0.7	0.21
Time before surgery, hours (mean ± SD)	33.9 ± 44.6	24.4 ± 30.3	0.026
Time before surgery, hours (median min-max)	17.7 (0.8-361.8)	15.7 (2.5-236.2)	0.054
Laparoscopy (%)			<0.001
Yes	76 (31.4)	13 (12.6)	
No	166 (68.6)	90 (87.4)	
Conversion rate (%)			<0.001
Yes	25 (32.9)	11 (84.6)	
No	51 (67.1)	2 (15.4)	

SD: standard deviation, BMI: body mass index. ASA: American Society of Anesthesiologists

[†] Student t-test with Welch's correction, or Mann-Whitney t-test, or X² test for binary or categorical variable

Table 3. Comparison of outcomes after the first hospitalization for a SBO episode

	Surgical, no resection N=242	Surgical resection N=103	Conservative N=147	P-value [†]
Length of hospital stay, days ± SD	14.6 ± 15.7	17.4 ± 13.7	7.8 ± 13.7	<0.001
Length of hospital stay, days (%)				<0.001
1-6	58 (24.0)	7 (6.8)	99 (67.3)	
7-12	95 (39.2)	42 (40.8)	31 (21.1)	
13-174	89 (36.8)	54 (52.4)	17 (11.6)	
Post-surgical complications, Clavien-Dindo classification (%)			NA	NA
0	166 (68.6)	57 (55.3)		
1	27 (11.1)	8 (7.8)		
2	32 (13.2)	20 (19.4)		
3	5 (2.1)	6 (5.8)		
4	6 (2.5)	5 (4.9)		
5	6 (2.5)	7 (6.8)		
SBO recurrence (%)				<0.0001
Yes	35 (14.5)	8 (7.8)	46 (31.3)	
No	207 (85.5)	95 (92.2)	101 (68.7)	

SD: standard deviation

[†] X² test for binary or categorical variable

Table 4. Risk factors for small bowel resection versus conservative treatment in patients with a small bowel obstruction, multivariate logistic regression model.

	Unadjusted OR (95% confidence interval)	Adjusted OR (95% confidence interval)	Adjusted OR P-value
Age ≥ 70 years (vs younger)	3.03 (1.79 – 5.12)	3.47 (1.74 – 6.92)	<0.001
First episode of obstruction (vs recurrence)	3.11 (1.68 – 5.77)	2.36 (1.07 – 5.22)	0.033
Absence of bowel movement ≥ 3 days (vs 0-2 days)	2.33 (1.09 – 4.97)	4.07 (1.58 – 10.44)	0.004
Guarding present (vs absent)	7.10 (3.55 – 14.18)	7.76 (3.17 – 19.04)	<0.001
C reactive protein ≥ 50 (vs less)	2.20 (1.13 – 4.28)	3.10 (1.39 – 6.92)	0.006
CT scan: transition point (vs none)	2.38 (1.11 – 5.13)	2.99 (1.05 – 8.54)	0.040
CT scan: lack of contrast enhancement (vs presence)	14.58 (4.26 – 49.95)	8.10 (1.86 – 35.35)	<0.001
CT scan: intra-abdominal fluid ≥ 500 mL	8.00 (4.01 – 15.94)	6.16 (2.58 – 14.60)	<0.001

Table 5. Frequency distributions of the three clinical scores and probabilities of surgical resection in 250 patients with small bowel obstruction. High-risk groups are underlined.

	Full 8-item score		7-item score (no contrast)		Brief 4-item score	
Score	N (%)	Percent with resection	N (%)	Percent with resection	N (%)	Percent with resection
0	5 (2.0)	0.0%	5 (2.0)	0%	78 (31.2)	12.8%
1	33 (13.2)	6.1%	34 (13.6)	8.8%	111 (44.4)	36.0%
2	62 (24.8)	16.1%	62 (24.8)	16.1%	33 (13.2)	78.8%
3	66 (26.4)	36.4%	76 (30.4)	40.8%	21 (8.4)	95.2%
4	58 (23.2)	70.7%	52 (20.8)	73.1%	7 (2.8)	100%
5	11 (4.4)	100%	14 (5.6)	100%	-	-
6	11(4.4)	100%	4 (1.6)	100%	-	-
7	4 (1.6)	100%	3 (1.2)	100%	-	-
8	0 (0.0)	0.0%	-	-	-	-

Table 6. Comparison of three scores for the prediction of small bowel resection in patients with small bowel obstruction: full 8-item score, 7-item score (omitting injection of contrast on CT-scan) and brief 4-item score, in development dataset and in external validation dataset.

Number of items	8	7	4
Variables in score:			
◦ Age ≥ 70 years	✓	✓	✓
◦ 1 st SBO episode	✓	✓	-
◦ ≥ 3 days since BM	✓	✓	-
◦ Abdominal guarding	✓	✓	✓
◦ CRP ≥ 50 mg/l	✓	✓	-
◦ Presence of transition point (CT)	✓	✓	-
◦ Lack of SB contrast enhancement (CT)	✓	-	✓
◦ Abdominal fluid ≥ 500 ml (CT)	✓	✓	✓
Area under ROC curve (AUC)	0.844	0.826	0.802
AUC in external validation set	0.821	0.811	0.774
Percentage at risk for threshold ≥ 0.70 *	33.6	29.2	24.4
Percentage at risk for same threshold in external validation set	32.2	32.2	28.7
Sensitivity for risk threshold ≥ 0.70	65.0	57.3	51.5
Sensitivity for same risk threshold in external validation set	72.7	72.7	60.6
Specificity for risk threshold ≥ 0.70	88.4	90.5	94.6
Specificity for same risk threshold in external validation set	84.1	84.1	84.1

* Score ≥ 4 for 8-item score and 7-item score, and ≥ 2 for 4-item score

BM: bowel movement. SB: small bowel.

Table 7. Clinical severity score for each group

	Surgical, no resection N=242	Surgical resection N=103	Conservative N=147	P-value [†] No res vs Res	P-value [†] No res vs Cons	P-value [†] Res vs Cons
Clinical severity score (%)						
0	1 (0.4)	0	5 (3.4)	<0.001	<0.001	<0.001
1	9 (3.7)	2 (1.9)	31 (21.1)			
2	64 (26.5)	10 (9.7)	52 (35.4)			
3	81 (33.5)	24 (23.3)	42 (28.6)			
4	55 (22.7)	41 (39.8)	17 (11.6)			
5	24 (9.9)	11 (10.7)	0			
6	8 (3.3)	11 (10.7)	0			
7	0	4 (3.9)	0			
8	0	0	0			
Clinical severity score (mean \pm SD)	3.2 \pm 1.2	4.0 \pm 1.3	2.2 \pm 1.0	<0.001	0.001	<0.001

One point was given for each present feature: age \geq 70 years old, 1st SBO episode, \geq 3 days since last bowel movement, abdominal guarding, CRP \geq 50mg/l, change of bowel diameter, lack of contrast enhancement and abdominal fluid \geq 500ml on CT scan. The score range from 0 to 8.

[†] χ^2 test for binary or categorical variable

SD: standard deviation

* Means comparison for continuous variables using ANOVA