Derivation and Validation of the ED-SAS Score for Very Early Prediction of Mortality and Morbidity with Acute Pancreatitis: A Retrospective Observational Study

Joseph Miller, MD (✉ jmiller6@hfhs.org)

Henry Ford Hospital, Detroit, MI, USA; Wayne State University, Detroit, MI, USA

Yiyang Wu MD, PhD

Nashville Biosciences, Nashville, TN, USA; Vanderbilt University Medical Center, Nashville, TN, USA

Rawan Safa, MD

Washington University School of Medicine, St. Louis, MO, USA

Georgiana Marusca, MD

Henry Ford Hospital, Detroit, MI, USA

Sandeep Bhatti, BS

Lake Erie College of Osteopathic Medicine, Erie, PA, USA

Guneet Ahluwalia, MD

Henry Ford Hospital, Detroit, MI, USA

Jad Dandashi, MD

Henry Ford Hospital, Detroit, MI, USA

Harold Gomez Acevedo, MD

Henry Ford Hospital, Detroit, MI, USA

Naureen Farook, MD

Henry Ford Hospital, Detroit, MI, USA

Ashley Scott, MD

Henry Ford Hospital, Detroit, MI, USA

Vidhya Nair, MD

Henry Ford Hospital, Detroit, MI, USA

Angie Adhami, MD

Henry Ford Hospital, Detroit, MI, USA

Jeffrey Dueweke, MD

Henry Ford Hospital, Detroit, MI, USA

Sudarshan Hebbar, MD

Calcimedica, La Jolla, CA, USA

Leeland Ekstrom, PhD

Nashville Biosciences, Nashville, TN, USA

Research Article

Keywords: Acute pancreatitis, predictive score, validation study, emergency department

Posted Date: September 18th, 2020

DOI: <https://doi.org/10.21203/rs.3.rs-75848/v1>

License: This work is licensed under a Creative Commons Attribution 4.0 International License.

[Read Full License](#)

Abstract

Background: Existing scoring systems to predict mortality in acute pancreatitis may not be directly applicable to the emergency department (ED). The objective of this study was to derive and validate the ED-SAS, a simple scoring score using variables readily available in the ED to predict mortality in patients with acute pancreatitis.

Methods: This retrospective observational study was performed based on patient level data collected from electronic health records across 2 independent health systems, one used for the derivation cohort and one for the validation cohort. Adult patients who were eligible presented to the ED, required hospital admission, and had a confirmed diagnosis of acute pancreatitis. Patients with chronic or recurrent episodes of pancreatitis were excluded. The primary outcome was 30-day mortality. Analyses tested and derived candidate variables to establish a prediction score and that was subsequently applied to the validation cohort to assess odds ratio for the primary and secondary outcomes.

Results: The derivation cohort included 599 patients, and the validation cohort 2011 patients. Thirty-day mortality was 4.2% and 3.9% respectively. From the derivation cohort, 3 variables were established for use in the predictive scoring score: ≥ 2 systemic inflammatory response syndrome (SIRS) criteria, age >60 years, and SpO₂ $<96\%$. Summing the presence or absence of each variable yielded an ED-SAS score ranging from 0 to 3. In the validation cohort, the odds of 30-day mortality increased with each subsequent ED-SAS point: 4.4 (95% CI 1.8 – 10.8) for 1 point, 12.0 (95% CI 4.9 – 29.4) for 2 points, and 41.7 (95% CI 15.8 – 110.1) for 3 points (c-statistic = 0.77).

Conclusion: An ED-SAS score that incorporates SpO₂, age, and SIRS measurements provides a rapid method for predicting 30-day mortality in acute pancreatitis.

Background

Recently, the incidence of acute pancreatitis has risen worldwide, with results from a retrospective analysis in the US showing a 13% increase in hospital admissions over 10 years.^{1,2} The annual incidence of acute pancreatitis in the US ranged from 13 to 45 per 100,000 people.³ While most patients who present with acute pancreatitis in the emergency department (ED) achieve positive outcomes, significant morbidity and mortality still exists.⁴ On average, up to 25% will progress to severe acute pancreatitis, and pancreatic necrosis will develop in 20% to 30% of patients, both of which are associated with significant mortality.^{5,6}

Multiple tools exist to aid clinicians in predicting morbidity and mortality risk associated with acute pancreatitis. These include well-known scoring systems such as Ranson's criteria, the Acute Physiology and Chronic Health Evaluation (APACHE)-II score, and the Bedside Index for Severity in Acute Pancreatitis (BISAP).⁷⁻⁹ However, these scoring systems largely incorporate variables collected over 24 hours or more of hospital admission and it may not be possible to obtain a final score until 48 hours post-admission.

Very early prognostic scores that apply to the ED triage and management of patients with acute pancreatitis are lacking.

The primary objective of this study was to establish a simple prognostic assessment of acute pancreatitis that clinicians can apply in the ED. In developing this objective, two key factors were observed that were repeatedly present early in severe acute pancreatitis. Systemic inflammatory response syndrome (SIRS) was a prominent indicator of disease severity and¹⁰⁻¹² mild to moderate hypoxia was common and likely associated with early acute lung injury (ALI).¹³ Hence, these findings were incorporated with other variables often available in the ED to derive and validate the ED-SAS, a simple scoring system that provides a prognostic assessment of mortality in patients admitted to the ED with acute pancreatitis.

Methods

Study Design and Setting

A retrospective analysis was performed at Vanderbilt University Medical Centers (VUMC) in Nashville, TN, USA and the Henry Ford Hospital System (HFHS) in Detroit, MI, USA. VUMC is a level 1 trauma center with approximately 125,000 annual ED visits and HFHS includes 9 Eds, 4 of which are free-standing, and 5 hospitals that experience approximately 460,000 annual ED encounters.

Selection of Participants

Data from VUMC were used for the derivation cohort (January 2013 to September 2017) and data from HFHS was used for the validation cohort (January 2014 to January 2018). Eligible patients were selected in similar manner from VUMC's Synthetic Derivative, a de-identified mirror-image of the system's electronic health records, by Nashville Biosciences, a subsidiary of VUMC established to support translational research, and from HFHS' electronic health records (EPIC, Verona, WI). Eligible patients were 18 years of age or older, had a diagnosis of acute pancreatitis (International Classification of Diseases [ICD], 9th Revision, code 577.0 or ICD, 10th Revision, code K85), and required hospital admission within 24 hours of presentation in the ED. Only the first encounter with a patient over the study assessment period was included to ensure no patient was represented more than once in the data set. We excluded patients with chronic pancreatitis, hepatic failure, or with a serum lipase <3x upper limit of the normal reference range. While prognostic assessment of patients discharged from the ED may be useful, we did not include these patients given high rates of concomitant chronic pancreatitis and reduced opportunity to obtain outcomes data given the retrospective nature of the study.

Data Collection and Processing

Data collected for the derivation and validation cohorts included subject-level information on demographics; relevant comorbidities based on ICD codes; diagnoses of respiratory failure, sepsis, acute kidney injury (AKI), and other relevant sequelae as per ICD codes; mechanical ventilation determined by

procedure codes for invasive mechanical ventilation; hospital procedure and visit records; mortality; and the first ED-recorded lab values and vital signs. Mechanical ventilation did not include non-Invasive modes of ventilation.

Data analysts pulled electronic health data using standardized methods within each health system. For the validation cohort only, additional chart abstraction was performed to assess the accuracy of electronic data collection for the diagnosis of acute pancreatitis. Physician chart abstractors were blinded to the study hypothesis. All abstractors were trained before initiating chart abstraction and were provided with a data collection manual that including variable definitions and details. Within the validation cohort, physicians abstracted 10% of study charts to authenticate electronic data capture and to test agreement with electronic chart abstraction for the diagnosis of acute pancreatitis. The *K* coefficient for the diagnosis of acute pancreatitis was 0.94. For missing data, it was assumed that a diagnosis or death was not present if not documented.

Outcome Measures

The primary outcome was 30-day mortality. Secondary outcomes included 180-day mortality; intensive care unit (ICU) admission; length of hospital stay; presence of sepsis, respiratory failure, or AKI; and need for dialysis or mechanical ventilation. To determine these latter outcomes, we used ICD codes documented during the patient's index hospitalization for acute pancreatitis.

Primary Data Analysis

We selected a priori predictor variables known to be associated with mortality from acute pancreatitis based on existing disease models.^{8,10,14,15} The variable selection was refined based on their availability during routine ED evaluation and consistent documentation within electronic health records. The a priori predictor variables included age >60 years, gender, self-reported race/ethnicity, the Charlson comorbidity index,¹⁶ the presence of ≥ 2 SIRS criteria (heart rate >90/min, respiratory rate >20/min, temperature >38.0°C or <36.0°C, white blood cells $>12 \times 10^9/L$, or white blood cells $<4 \times 10^9/L$), the presence of SpO2 saturation <96%, and hematocrit >44%. Of importance, the SpO2 cut-off of <96% was used rather than <95% as it includes the descending portion of the oxygen saturation curve.

For the derivation cohort, these variables were analyzed for their association with 30-day mortality with univariate analysis and retained variables for the final model with $p < 0.10$. A complete case analysis was also performed. To reduce bias in parameter estimates given the low event rate, multivariable logistic regression with Firth's penalized likelihood estimate was utilized.¹⁷ To determine the most parsimonious predictor model, the Akaike Information Criterion was assessed. In addition, model calibration using graphical assessment by loess smoothers¹⁸ and model discrimination with area under the receiver operating characteristic curve (*c* statistic) were evaluated.

The final three variables included in the model were SpO2 <96%, age >60 years, and ≥ 2 SIRS criteria (Table 1). Each variable was equally weighed and contributed 0 or 1 points to an overall score based on

the presence or absence of these three variables resulting in an ED-SAS range of 0 to 3 points. We named the score ED-SAS for SpO₂, age, and SIRS. Logistic regression for these score categories with 0 as the reference to assess the odds of death and all secondary outcomes within the derivation data set was performed and applied this derived predictor score to the validation sample to assess its association with primary and secondary outcomes. No model recalibration was performed within the validation data. All analyses were performed with SAS 9.4 (Cary, NC).

Results

The derivation cohort included 599 patients, and the validation cohort included 2011 patients. In general, mean age, SpO₂, and respiratory rate were similar between the two cohorts (Table 2). The 30-day mortality rate was similar across the two cohorts with 25 (4.2%) deaths in the derivation cohort and 78 (3.9%) in the validation cohort (Table 3). Respiratory failure rates were also similar between the two cohorts (4.2% vs 5.4%, respectively), but rates of sepsis were higher in the derivation cohort than in the validation cohort (13.4% vs. 6.6%, respectively; Table 3). In addition, length of hospital stay, and ICU admissions were higher in the derivation cohort (Table 3).

While four a priori variables were associated with 30-day mortality in univariate analysis in the derivation cohort, only the presence of ≥ 2 SIRS criteria, an ED SpO₂ saturation $< 96\%$, and age > 60 years were significant upon the multivariate analysis and were retained in the final ED-SAS Score (Table 4). After application of the ED-SAS score, the probability of 30-day mortality increased incrementally with higher ED-SAS scores for both cohorts (Figure 1). In addition, increases in the odds ratio (OR) for 30-day mortality, 180-day mortality, AKI, respiratory failure, sepsis, and ICU admission were directly associated with increased ED-SAS score in the validation cohort (Table 5).

Discussion

The incidence of acute pancreatitis has been increasing worldwide, and while multiple risk stratifying tools have previously been developed, they come with limitations.^{1,2,7-9} The 3 most commonly used risk stratification tools in the ED include the BISAP score, the APACHE II Scoring System, and the Ranson's criteria.¹ Imaging-based scores also exist, which grade computed tomography findings of pancreatitis, but may not be performed routinely and may underestimate severity in patients who present early in the disease course.^{14,19-21} While these scoring systems are regularly used to predict mortality, they were not designed specifically for utility in the ED.^{1,7-9}

The ability to identify patients with acute pancreatitis who are at high-risk for mortality using a stratification tool focused on early ED variables could potentially lead to early and rapid identification of patients with poor prognoses. In this study across two health systems and multiple EDs, we derived and validated the ED-SAS, a simple prognostic tool that focuses on 3 variables readily available in the early evaluation of a patient with acute pancreatitis: age, pulse oximetry reading on room air (SpO₂ saturation $< 96\%$), and the presence of SIRS criteria. Results from this analysis suggest an increased risk of 30-day

mortality for each incrementally higher score in the validation cohort which could potential give ED physicians the ability to predict mortality based on ED-SAS scores and could guide hospital transfer decisions and prognostic discussions with patients and families.

The inclusion of advanced age, SIRS criteria, and low pulse oximetry are consistent with the existing body of literature. Age has been validated in existing inpatient prediction scores and SIRS is well known to impact the clinical course and outcome in acute pancreatitis.^{10,22,23} SIRS is also an early predictor of severe acute pancreatitis with studies demonstrating that the severity of acute pancreatitis is greater among patients with SIRS on day 1.²⁴ While less recognized than SIRS, moderate to severe arterial hypoxia is common in patients with acute pancreatitis and is related to the ALI often associated with the inflammatory response seen with acute pancreatitis.^{10,13} Among patients that die early from acute pancreatitis, ALI is often a prominent finding and is present in approximately 60% of all elderly patients that pass from acute pancreatitis within the first week.^{15,25} Although prior studies used PaO₂ arterial blood gas measurements to assess arterial hypoxia, we relied on pulse oximetry in this study given the infrequent use of arterial blood gas in the ED for these patients.¹³

This retrospective, observational, chart review study relied on commonly documented predictor variables comes with some limitations. While standardized methods for data collection were used, there remains a potential for bias, and there may be important predictors of morbidity and mortality only available through prospective data collection that were not considered. We did not perform a direct comparison between the ED-SAS score and other common scoring systems or ED predictors of mortality such as the quick Sepsis Related Organ Failure Assessment score, which may have utility in evaluating acute pancreatitis as it does in sepsis.²⁶ In addition, the design for assessing mortality may underestimate the true rates given subjects were not directly contracted and electronic health record data were used to determine 30- and 180-day mortality. While the ED-SAS model provides a simple assessment of high-risk patients with acute pancreatitis early in their ED course, it was not designed to assess outcomes or the intensity of inpatient services that patients require to adequately determine eligibility for ED discharge. However, as this information would be of great interest to emergency physicians, additional studies to assess this variable would be of value. Finally, we did not compare the ED-SAS score to clinical gestalt and are therefore unable to assess whether it provides value added to the clinical judgement of an experienced physician.

Conclusions

In this retrospective, observation analysis of patients hospitalized for acute pancreatitis, the ED-SAS was derived and validated as a simple predictive tool that could be used to estimate the 30-day risk of mortality using patient characteristics commonly available in the ED. This tool has the potential to aid clinicians in prognosticating patients with acute pancreatitis.

Abbreviations

AKI, acute kidney injury

ALI, acute lung injury

APACHE-II, Acute Physiology and Chronic Health Evaluation II score

BISAP, Bedside Index for Severity in Acute Pancreatitis

ED, emergency department

ED-SAS, emergency department SpO₂, age, and SIRS

HFHS, Henry Ford Hospital System

ICD, International Classification of Diseases

ICU, intensive care unit

SIRS, systemic inflammatory response syndrome

VUMC, Vanderbilt University Medical Centers

Declarations

Ethics approval and consent to participate: The study was approved by the institutional review boards at the Henry Ford Health System and Vanderbilt University. A waiver of informed consent was granted by both institutional review boards for conduct of the study.

Consent for publication: Not applicable.

Availability of data and materials: Data are available on reasonable request from the corresponding author.

Competing Interests: JM, YW, RS, GM, SB, GA, JD, HGA, NF, AS, VN, AA, JD, and LE report no conflicts of interest related to this work. SH serves as chief medical officer for CalciMedica (La Jolla, CA, USA).

Funding: No funding was provided was this study. All authors had full access to all the data in the study and provided input on the analyses before and during writing of the report and take responsibility for the integrity of the data and the accuracy of the data analysis. JM and RS primarily composed the manuscript; CalciMedica (La Jolla, CA, USA) commissioned medical writing assistance from Sarah Odeh (San Francisco, CA, USA) to support subsequent drafts under the direction of all authors. All authors reviewed and revised each draft and approved the final submitted version. The corresponding author had final responsibility for the decision to submit for publication.

Authors' Contributions: JM, YW, and SH conceived and designed the study. JM, YW, GA, HGA, NF, AS, VN, AA, and JD worked on data acquisition. RS and JM assisted with the statistical analysis and interpretation. All authors participated in synthesis of the data and manuscript review. All authors approved the final manuscript. JM and RS primarily composed the manuscript and take responsibility for it as a whole.

Acknowledgements: Not applicable.

References

1. Waller A, Long B, Koyfman A, Gottlieb M. Acute pancreatitis: updates for emergency clinicians. *J Emerg Med*. 2018;55:769-79.
2. Krishna SG, Kamboj AK, Hart PA, Hinton A, Conwell DL. The changing epidemiology of acute pancreatitis hospitalizations. a decade of trends and the impact of chronic pancreatitis. *Pancreas*. 2017;46(4):482-8.
3. Yadav D, Lowenfels AB. The epidemiology of pancreatitis and pancreatic cancer. *Gastroenterol*. 2013;144:1252-61.
4. Eachempati SR, Hydo LJ, Barie PS. Severity scoring for prognostication in patients with severe acute pancreatitis: comparative analysis of the Ranson score and the APACHE III score. *Arch Surg*. 2002;137:730-6.
5. Barauskas G, Ignatavičius P, Vitkauskienė A, Pundzius J, Dambrauskas Ž. Impact of etiology on course and outcomes of severe acute pancreatitis. *Medicina (Kaunas)*. 2015;51(3):167-72.
6. Singh VK, Moran RA, Afghani E, de-Madaria. Treating acute pancreatitis: what's new? *Exp Rev Gastroenterol Hepatol*. 2015;9(7): 901-11.
7. Bradley EL. A clinically based classification system for acute pancreatitis. Summary of the International Symposium on Acute Pancreatitis, Atlanta, Ga, September 11 through 13, 1992. *Arch Surg*. 1993;128:586-90.
8. Larvin M, McMahon MJ. APACHE-II score for assessment and monitoring of acute pancreatitis. *Lancet*. 1989;2:201-5.
9. Wu BU, Johannes RS, Sun X, Tabak Y, Conwell DL, Banks PA. The early prediction of mortality in acute pancreatitis: a large population-based study. *Gut* 2008;57:1698-703.
10. Shields CJ, Winter DC, Redmond HP. Lung injury in acute pancreatitis: mechanisms, prevention, and therapy. *Curr Opin Crit Care*. 2002;8:158-63.
11. Mofidi R, Duff MD, Wigmore SJ, Madhavan KK, Garden OJ, Parks RW. Association between early systemic inflammatory response, severity of multiorgan dysfunction and death in acute pancreatitis. *The Br J Surg*. 2006;93:738-44.
12. Mofidi R, Lee AC, Madhavan KK, Garden OJ, Parks RW. Prognostic factors in patients undergoing surgery for severe necrotizing pancreatitis. *World J Surg*. 2007;31:2002-7.

13. Imrie CW, Ferguson JC, Murphy D, Blumgart LH. Arterial hypoxia in acute pancreatitis. *The Br J Surg.* 1977;64:185-8.
14. Papachristou GI, Muddana V, Yadav D, et al. Comparison of BISAP, Ranson's, APACHE-II, and CTSI scores in predicting organ failure, complications, and mortality in acute pancreatitis. *Am J Gastroenterol.* 2010;105:435-41.
15. Andersson R, Andren-Sandberg A. Fatal acute pancreatitis. Characteristics of patients never reaching hospital. *Pancreatol.* 2003;3:64-6.
16. Charlson ME, Pompei P, Ales KL, MacKenzie CR. A new method of classifying prognostic comorbidity in longitudinal studies: development and validation. *J Chronic Dis.* 1987;40:373-83.
17. Heinze G, Puhr R. Bias-reduced and separation-proof conditional logistic regression with small or sparse data sets. *Stat Med.* 2010;29:770-7.
18. Austin PC, Steyerberg EW. Graphical assessment of internal and external calibration of logistic regression models by using loess smoothers. *Stat Med.* 2014;33:517-35.
19. Bollen TL, Singh VK, Maurer R, et al. A comparative evaluation of radiologic and clinical scoring systems in the early prediction of severity in acute pancreatitis. *Am J Gastroenterol.* 2012;107:612-9.
20. Kuo DC, Rider AC, Estrada P, Kim D, Pillow MT. Acute pancreatitis: what's the score? *J Emerg Med.* 2015;48:762-70.
21. Sahu B, Abbey P, Anand R, Kumar A, Tomer S, Malik E. Severity assessment of acute pancreatitis using CT severity index and modified CT severity index: Correlation with clinical outcomes and severity grading as per the Revised Atlanta Classification. *Indian J Radio Imaging.* 2017;27:152-60.
22. McKay CJ, Gallagher G, Brooks B, Imrie CW, Baxter JN. Increased monocyte cytokine production in association with systemic complications in acute pancreatitis. *Br J Surg.* 1996;83:919-23.
23. Lundberg AH, Granger DN, Russell J, et al. Quantitative measurement of P- and E-selectin adhesion molecules in acute pancreatitis: correlation with distant organ injury. *Ann Surg.* 2000;231:213-22.
24. Singh VK, Wu BU, Bollen TL, et al. Early systemic inflammatory response syndrome is associated with severe acute pancreatitis. *Clin Gastroenterol Hepatol.* 2009;7:1247-51.
25. Jacobs ML, Daggett WM, Civette JM, et al. Acute pancreatitis: analysis of factors influencing survival. *Ann Surg.* 1977;185:43-51.
26. Hallac A, Puri N, Applebury D, et al. The value of quick sepsis-related organ failure assessment scores in patients with acute pancreatitis who present to emergency departments: a three-year cohort study. *Gastroenterol Res.* 2019;12:67-71.

Tables

Table 1. ED-SAS Score.

Parameter	Score
SpO2 <96%	1
Age >60 years	1
≥2 SIRS criteria*	1
Total Score, range	0 – 3

*Presence of 2 or more of the following: fever >38.0°C or hypothermia <36.0°C, tachycardia >90 beats/minute, tachypnea >20 breaths/minute, Leukocytosis >12*10⁹/L, or leucopenia <4*10⁹/L.

ED-SAS, Emergency Department SpO2, Age, and SIRS; SIRS, systemic inflammatory response syndrome

Table 2: Baseline characteristics of patients with acute pancreatitis admitted to the ED.*

	Derivation Cohort N=599	Validation Cohort N=2011
Gender, n (%)		
Male	316 (52.8)	977 (48.6)
Female	283 (47.3)	1034 (51.4)
Mean Age, years (SD)	53.4 (17.4)	56.1 (17.8)
Race, n (%)		
Black	81 (13.5)	487 (24.2)
White	501 (83.6)	1359 (67.5)
Other	16 (2.7)	165 (8.2)
Ethnicity, n (%)		
Hispanic	548 (6.2)	79 (3.9)
Non-Hispanic	562 (93.8)	1886 (93.8)
Unknown	54 (9.0)	20 (1.0)
Vitals, mean (SD)		
Heart Rate, bpm	91.2 (20.1)	87.9 (20.2)
Respiratory Rate, breath/min	18.4 (2.9)	18.5 (5.1)
Temperature, °C	36.7 (0.6)	36.6 (0.6)
SpO2, %	97.5 (2.7)	97.4 (2.5)
Comorbidities, n (%)		
Cancer	128 (27.83)	171 (13.96)
Stroke	85 (18.48)	120 (9.8)
Congestive Heart Failure	110 (23.91)	250 (20.41)
Dementia	5 (1.09)	90 (7.35)
Diabetes Mellitus	151 (32.83)	443 (36.16)
Myocardial Infarction	54 (11.74)	185 (15.1)
Chronic Kidney Disease	112 (24.35)	323 (26.37)
COPD	138 (30.00)	463 (37.8)
Additional Predictive Variables		
Charlson comorbidity index, mean (SD)	2.6 (2.3)	3.3 (3.7)

≥ 2 SIRS criteria, n (%) [†]	182 (30.4)	512 (25.5)
SpO ₂ <96%, n (%)	96 (16.0)	336 (16.7)
WBC >12 × 10 ⁹ /L, n (%)	259 (43.2)	740 (36.8)
Hematocrit >44%, n (%)	108 (18.0)	601 (29.9)

COPD, chronic obstructive pulmonary disease; SIRS, systemic inflammatory response syndrome; WBC, white blood cells.

*Additional parameters were not available in the data set; [†]Presence of 2 or more of the following: fever >38.0°C or hypothermia <36.0°C, tachycardia >90 beats/minute, tachypnea >20 breaths/minute, leukocytosis >12*10⁹/L, or leucopenia <4*10⁹/L.

Table 3: Primary and secondary outcomes upon review of electronic health record data.

Outcome, n (%)	Derivation Cohort N=599	Validation Cohort N=2011
30-Day Mortality	25 (4.2)	78 (3.9)
180-Day Mortality	91 (15.19)	243 (12.1)
Sepsis	80 (13.36)	133 (6.61)
Acute Kidney Injury	155 (25.88)	445 (22.13)
Respiratory Failure	25 (4.17)	108 (5.37)
Mechanical Ventilation	7 (1.17)	75 (3.7)
Dialysis	9 (1.50)	58 (2.88)
ICU Admission	83 (13.86)	159 (7.91)
Length of stay, mean days (SD)	6.7 (7.6)	3.6 (4.8)

ICU, intensive care unit.

Table 4: Univariate analysis of acute pancreatitis predictor variables.

Predictor	30-Day Mortality Odds Ratio (95% CI)
Female gender	1.1 (0.5 – 2.7)
Age >60 years*	7.6 (2.5 – 23.2)
Black Race	0.3 (0.0 – 2.4)
Hematocrit >44%	0.5 (0.1 – 2.2)
SpO2 <96%*	5.7 (2.3 – 14.2)
≥2 SIRS criteria* [†]	4.3 (1.0 – 18.7)
Charlson Index	1.2 (1.0 – 1.5)

*Statistically significant predictors upon implementation of the multivariate analysis that were retained in final model; [†]Fever >38.0°C or hypothermia <36.0°C, tachycardia >90 beats/minute, tachypnea >20 breaths/minute, leukocytosis >12*10⁹/L, or leucopenia <4*10⁹/L.

SIRS, systemic inflammatory response syndrome.

Table 5. Odds of primary and secondary outcomes in validation cohort based on ED-SAS score.

ED-SAS score, OR (95% CI)*	ED-SAS = 1	ED-SAS = 2	ED-SAS = 3
30-day Mortality	4.4 (1.8 – 10.8)	12.0 (4.9 – 29.4)	41.7 (15.8 – 110.1)
180-day Mortality	3.1 (1.8 – 5.4)	6.1 (3.4 – 10.9)	14.8 (7.3 – 30.1)
Acute kidney injury	2.0 (1.6 – 2.6)	3.3 (2.4 – 4.5)	5.6 (3.4 – 9.3)
Respiratory failure	2.9 (1.6 – 4.9)	5.7 (3.1 – 10.3)	7.6 (3.3 – 17.2)
Sepsis	2.9 (1.7 – 5.0)	7.2 (4.1 – 12.7)	17.1 (8.5 – 34.3)
ICU Admission	2.3 (1.5 – 3.5)	3.1 (1.9 – 5.0)	4.6 (2.3 – 9.3)

*Reference is ED-SAS of 0.

Figures

Figure 1

Probability of 30-day mortality based on ED-SAS scores. After application of the ED-SAS score, the probability of 30-day mortality increased incrementally with higher ED-SAS scores for both cohorts