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Outcomes/Predictions

## A risk scoring model based on vital signs and laboratory data predicting transfer to the intensive care unit of patients admitted to gastroenterology wards



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#### ABSTRACT

*Purpose:* To compare the ability of a score based on vital signs and laboratory data with that of the modified early warning score (MEWS) to predict ICU transfer of patients with gastrointestinal disorders. *Materials and methods:* Consecutive events triggering medical emergency team activation in adult patients ad-

mitted to the gastroenterology wards of the Asan Medical Center were reviewed. Binary logistic regression was used to identify factors predicting transfer to the ICU. Gastrointestinal early warning score (EWS-GI) was calculated as the sum of simplified regression weights (SRW).

*Results:* Of the 1219 included patients, 468 (38%) were transferred to the ICU. Multivariate analysis identified heart rate  $\geq$  105 bpm (SRW 1), respiratory rate  $\geq$  26 bpm (SRW 2), ACDU (Alert, Confused, Drowsy, Unresponsive) score  $\geq$  1 (SRW 2), SpO<sub>2</sub>/FiO<sub>2</sub> ratio < 240 (SRW 2), creatinine  $\geq$ 2.0 mg/dL (SRW 2), total bilirubin  $\geq$ 9.0 mg/dL (SRW 2), prothrombin time/international normalized ratio (INR)  $\geq$ 1.5 (SRW 2), and lactate  $\geq$ 3.0 mmol/L (SRW 2) for inclusion in EWS-GI. The area under the receiver operating characteristic curve of the EWS-GI was larger than that of MEWS (0.76 vs. 0.64; P < 0.001).

*Conclusions:* EWS-GI may predict ICU transfer among patients admitted to gastroenterology wards. The EWS-GI should be prospectively validated.

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#### 1. Introduction

Medical emergencies, including upper (variceal and non-variceal) and lower gastrointestinal bleeding, acute liver failure, severe sepsis/ septic shock, and respiratory insufficiency, occur frequently in patients admitted to gastroenterology wards [1-5]. Several risk assessment and scoring systems have been developed for upper gastrointestinal bleed-ing [1,2,6] and advanced liver disease [7]. However, to our knowledge, overall assessment scores for the risk of transfer to intensive care units (ICUs) have not been developed for patients with gastrointestinal disorders.

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Early warning scores (EWS) are bedside evaluation tools based on physiologic measurements (e.g., blood pressure, heart rate, respiratory rate) obtained at admission or by monitoring during hospitalization. EWS provide a simple method for categorizing a patient's condition and indicating when a patient may require additional attention [8-10]. Despite the potential ability of EWS to identify physical deterioration in acute care settings, improvements are needed [11]. In addition, most EWS are disease-nonspecific that do not consider the characteristics of patients with certain diseases [10]. For example, physiologic characteristics may differ in critically ill patients with and without gastrointestinal disorders [12], and traditional EWS may perform differently in these groups of patients. These factors indicate a need for EWS tailored to different patient groups.

The number of patients who can be monitored and treated in ICUs is restricted owing to resource limitations. Early identification of patients at risk of deterioration and the selection of those who might benefit from ICU care can be crucial. This study was therefore designed (i) to develop a risk screening tool, the gastrointestinal EWS (EWS-GI), using

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vital signs and laboratory data for patients admitted to gastroenterology wards; and (ii) to compare the ability of the EWS-GI and the previously described modified EWS (MEWS) [8] to predict ICU transfer of these patients.

### 2. Material and methods

## 2.1. Study design and study subjects

This retrospective cohort study was conducted at the Asan Medical Center, a 2680-bed university-affiliated hospital in Seoul, Korea. Consecutive events triggering medical emergency team (MET) activation in adult patients admitted to the gastroenterology wards of Asan Medical Center between March 2008 and December 2015 were reviewed. The MET at Asan Medical Center has been described [13]. The MET was automatically activated when a patient was identified by 24 h electronic medical record (EMR)-based monitoring as reaching a threshold for a relevant vital sign or laboratory measurement, based on the medical alert system criteria used at Asan Medical Center. The MET was also activated when it was telephoned or paged by a general ward (GW) nurse or resident, or when a cardiopulmonary resuscitation code blue was announced anywhere in the hospital. For patients with multiple MET activations, only the first activation was included. Events were excluded if a do not resuscitate order had been issued within 24 h of MET activation, if they involved cardiopulmonary resuscitation, or if the MET had been activated for simple procedural assistance or to educate healthcare providers. The primary study outcome was ICU transfer. The study protocol was approved by the institutional review board of Asan Medical Center (no. 2015-1101), which waived the requirement for informed consent because the study was retrospective, and the patient records were anonymized and de-identified prior to analysis.

### 2.2. Data collection and definitions

Data collected for all MET activations included demographic factors; comorbidities; causes of activation; vital signs; ACDU (Alert, Confused, Drowsy, Unresponsive) score;  $SpO_2/FiO_2$  (SF) ratio; outcome of the MET intervention; and mortality after MET activation. The vital signs and ACDU score were used to calculate the MEWS [8]. We used SF ratio instead of  $PaO_2/FiO_2$  ratio because arterial blood gas sampling was not available for every patient, but SF ratio was rapidly and easily measured by using the pulse oximetry. Moreover, previous studies found that the SF ratio, being highly correlated with  $PaO_2/FiO_2$  ratio, could be a useful tool for assessment of hypoxia in the setting of acute respiratory distress [14,15]. Laboratory data, including blood chemistry, coagulation profile, and serum lactate, were also collected on the day of MET activation.

#### 2.3. Statistical analysis

Continuous variables are reported as medians and interguartile ranges and were compared by the Mann–Whitney U test, whereas categorical variables are reported as percentages and were compared by the Chi-square or Fisher's exact test, as appropriate. The cutoff values of potential predictors of ICU transfer were selected using locally weighted scatterplot smoothing curves [16]. Multivariate regression analysis using stepwise backward selection was performed to identify factors predicting ICU transfer. The weights derived from multiple logistic regressions were simplified as natural numbers >0, and the EWS-GI was calculated as the sum of these simplified weights. The areas under the receiver operating characteristic (ROC) curve for the EWS-GI and MEWS predicting ICU transfer were compared by DeLong's test, as described [17]. The optimal cutoff values for the EWS-GI and MEWS were identified by ROC analysis. Kaplan-Meier survival estimates were stratified by EWS-GI and MEWS to compare their ability to predict mortality. All tests of significance were two tailed, and P values < 0.05 were considered significant. All analyses were performed using SPSS version 18.0 for Windows (SPSS Inc., Chicago, IL, USA) and MedCalc Statistical Software version 16.8.4 (MedCalc Software bvba, Ostend, Belgium).

#### 3. Results

During the study period, the MET was activated for 2172 patients admitted to the gastroenterology wards. Our gastroenterology department cares for approximately 19800 admissions (adult patients) per year, so there were 14.2 MET activations/1000 admissions. Of these, 953 were excluded from analysis. Of the remaining 1219 patients, 751 (62%) were treated in the GWs and 468 (38%) were transferred to the ICU (see Supplementary material for details).

The baseline characteristics and clinical outcomes of the study patients are shown in Table 1. The percentage of patients with liver cirrhosis was significantly higher in the ICU than in the GW group. MET activation was triggered automatically using only EMR-based screening criteria for 57% in the GW group and 39% in the ICU group (P < 0.001). Regarding activation cause, patients in the ICU group were more likely

# **Table 1**Clinical characteristics of the study patients<sup>a</sup>.

$\begin{array}{c ccccccccccccccccccccccccccccccccccc$		Total	General	ICU	Р
Age   61 (50-70)   62 (52-71)   58 (47-68)   <0.001     Male gender   795 (65)   488 (65)   307 (66)   0.83     Comorbidity      0.63   0.63     Diabetes   244 (20)   147 (20)   97 (21)   0.63     Chronic heart disease   296 (24)   190 (25)   106 (23)   0.29     Liver cirrhosis   541 (44)   297 (40)   244 (52)   <0.001		(n = 1219)	ward $(n - 751)$	(n = 468)	
Age   61 (50-70)   62 (52-71)   58 (47-68)   <0.001     Male gender   795 (65)   488 (65)   307 (66)   0.83     Comorbidity      0.63   0.63     Diabetes   244 (20)   147 (20)   97 (21)   0.63     Chronic heart disease   296 (24)   190 (25)   106 (23)   0.29     Liver cirrhosis   541 (44)   297 (40)   244 (52)   <0.001			(11 = 751)		
Male gender   795 (65)   488 (65)   307 (66)   0.83     Comorbidity	Age	61 (50–70)	62 (52–71)	58 (47–68)	< 0.001
Comorbidity   Diabetes   244 (20)   147 (20)   97 (21)   0.63     Chronic heart disease   296 (24)   190 (25)   106 (23)   0.29     Liver cirrhosis   541 (44)   297 (40)   244 (52)   <0.001	Male gender	795 (65)	488 (65)	307 (66)	0.83
Diabetes   244 (20)   14/ (20)   97 (21)   0.63     Chronic heart disease   296 (24)   190 (25)   106 (23)   0.29     Liver cirrhosis   541 (44)   297 (40)   244 (52)   <0.001	Comorbidity	244 (20)	147 (20)	07 (21)	0.02
Chronic heart disease 290 (24) 190 (25) 106 (23) 0.29   Liver cirrhosis 541 (44) 297 (40) 244 (52) <0.001	Diabetes Changing have to discuss	244 (20)	147 (20)	97 (21)	0.63
Liver cirrnosis   541 (44)   297 (40)   244 (52)   <0.001     Chronic kidney disease   30 (3)   18 (2)   12 (3)   0.86     Malignancy   492 (40)   348 (46)   144 (31)   <0.001	Chronic heart disease	296 (24)	190 (25)	106 (23)	0.29
Chronic kidney disease   30 (3)   18 (2)   12 (3)   0.86     Malignancy   492 (40)   348 (46)   144 (31)   <0.001	Liver cirrnosis	541 (44)	297 (40)	244 (52)	< 0.001
Malgnancy   492 (40)   348 (46)   144 (31)   <0.001     Activation type   <0.001	Chronic kidney disease	30(3)	18(2)	12(3)	0.86
Activation type   <0.001	Malignancy	492 (40)	348 (46)	144 (31)	< 0.001
EMR triggered   610 (50)   4.29 (57)   181 (39)     Call triggered   609 (50)   322 (43)   287 (61)     Activation cause   Active liver failure   183 (15)   45 (6)   138 (30)   <0.001	Activation type	610 (50)	400 (57)	101 (00)	<0.001
Call triggered   609 (50)   322 (43)   287 (61)     Activation cause   Active liver failure   183 (15)   45 (6)   138 (30)   <0.001	EMR triggered	610 (50)	429 (57)	181 (39)	
Activation cause Acute liver failure 183 (15) 45 (6) 138 (30) <0.001	Call triggered	609 (50)	322 (43)	287 (61)	
Acute liver failure $183(15)$ $45(6)$ $138(30)$ <0.001	Activation cause	102 (15)	45 (6)	120 (20)	.0.001
	Acute liver failure	183 (15)	45 (6)	138 (30)	< 0.001
Severe sepsis/septic 414 (34) 285 (38) 129 (28) <0.001 shock	shock	414 (34)	285 (38)	129 (28)	<0.001
Hypovolemic shock 164 (14) 98 (13) 66 (14) 0.60	Hypovolemic shock	164 (14)	98 (13)	66 (14)	0.60
Respiratory insufficiency 329 (27) 171 (23) 158 (34) <0.001 Vital signs	Respiratory insufficiency Vital signs	329 (27)	171 (23)	158 (34)	<0.001
Systolic blood pressure, 108 105 115 <0.001	Systolic blood pressure,	108	105	115	< 0.001
mm Hg (85–130) (85–128) (87–136)	mm Hg	(85-130)	(85-128)	(87-136)	
Heart rate, bpm 102 99 (82–116) 107 <0.001	Heart rate, bpm	102	99 (82–116)	107	< 0.001
(85–120) (91–123)	· .	(85-120)	· · · ·	(91-123)	
Respiratory rate, bpm 22 (18–28) 20 (18–26) 24 (20–30) <0.001	Respiratory rate, bpm	22 (18–28)	20 (18-26)	24 (20-30)	< 0.001
Temperature, °C 36.8 36.8 36.8 0.05	Temperature, °C	36.8	36.8	36.8	0.05
(36.4–37.7) (36.5–37.8) (36.4–37.5)		(36.4-37.7)	(36.5-37.8)	(36.4-37.5)	
ACDU score 0 (0-2) 0 (0-0) 1 (0-3) <0.001	ACDU score	0 (0-2)	0 (0-0)	1 (0-3)	< 0.001
SpO <sub>2</sub> /FiO <sub>2</sub> ratio 338 400 314 <0.001	SpO <sub>2</sub> /FiO <sub>2</sub> ratio	338	400	314	< 0.001
(222-462) (260-467) (188-452)		(222-462)	(260-467)	(188 - 452)	
Modified early warning 4 (3–5) 3 (2–5) 5 (3–6) <0.001	Modified early warning	4 (3–5)	3 (2–5)	5 (3–6)	<0.001
Laboratory data	Laboratory data				
Creatining $mg/dI$ 10(07-19) 09(07-16) 13(08-24) <0.001	Creatinine mg/dI	10(07-19)	09(07-16)	13(08-24)	< 0.001
Total bilirubin mg/dI $37$ $28(11-75)$ $63$ <0.001	Total bilirubin mg/dI	37	28(11-75)	63	< 0.001
(15-111) (21-220)	Total Diffabili, filg/al	(15-111)	2.0 (1.1-7.5)	(2.1 - 22.0)	×0.001
Prothrombin time INR $16(13-21)$ $14(12-18)$ $19(14-26) < 0.001$	Prothrombin time INR	(1.5 - 11.1) 16(13-21)	14(12 - 18)	(2.1 - 22.0) 19(14-26)	< 0.001
Lactate mmol/l $25(16-46)$ $21(14-36)$ $35(19-61) < 0.001$	Lactate mmol/I	25(16-46)	21(14-36)	35(19-61)	< 0.001
Mortality 2.5 (1.0 4.0) 2.1 (1.4 5.0) 5.5 (1.5-0.1) <0.001	Mortality	2.5 (1.0 4.0)	2.1 (1.4 3.0)	5.5 (1.5-0.1)	-0.001
14  day = 217  (18)  58  (8)  159  (34)  < 0.001	14 day	217 (18)	58 (8)	159 (34)	< 0.001
217(10) $30(0)$ $133(34)$ <0.001 28 day $322(26)$ $118(16)$ $204(44)$ <0.001	28 day	322 (26)	118 (16)	204 (44)	< 0.001
$\begin{array}{cccccccccccccccccccccccccccccccccccc$	60 day	431 (35)	185 (25)	246 (53)	<0.001

$$\label{eq:local_local_resonance} \begin{split} ICU &= \text{intensive care unit, EMR} = \text{electronic medical record, ACDU} = \text{alert/confused/}\\ drowsy/unresponsive, SpO_2 &= \text{peripheral oxygen saturation, FiO}_2 = \text{fraction of inspired}\\ oxygen, INR &= \text{international normalized ratio.} \end{split}$$

<sup>a</sup> Data are presented as median (interquartile range) or number (percentage) of patients.

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