



Quality monitoring of resuscitative endovascular balloon occlusion of the aorta using cumulative sum analysis in Korea: a case series

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Purpose: Resuscitative endovascular balloon occlusion of the aorta (REBOA) is a state-of-the-art lifesaving procedure. However, due to its high mortality and morbidity, including ischemia and reperfusion injury, well-trained medical staff and effective systems are needed. This study was conducted to investigate the learning curve for REBOA in Korea.

Methods: To monitor this learning curve, we used cumulative sum (CUSUM) analysis and graphs of mortality and aortic occlusion time within 60, 90, and 120 minutes for consecutive patients. The procedures performed between July 2017 and June 2021 were divided into pre-trauma center (pre-TC; July 2017–February 2020) and TC (February 2020–June 2021) periods.

Results: REBOA was performed for 31 consecutive patients with trauma. The pre-TC (n=12) and TC (n=19) periods did not differ significantly with regard to Injury Severity Score, age, injury mechanism, initial systolic blood pressure, prehospital cardiopulmonary resuscitation (CPR), or CPR in the emergency department. At the 17th consecutive patient during the TC period, CUSUM failure graphs for mortality and aortic occlusion time exhibited a downward inflection, indicating an improvement in performance.

Conclusions: The mortality and aortic occlusion time of REBOA improved, and these parameters can be monitored using CUSUM analysis at the hospital level.

Keywords: Aorta; Balloon occlusion; Endovascular procedures

INTRODUCTION

Severe hemorrhage due to torso injury is a leading cause of death [1,2]. To reduce mortality from severe torso hemorrhage, damage control surgery and resuscitation have been introduced [1]. The

core concept of damage control is prompt hemostasis, such as via emergency surgery or interventional radiology [1]. However, some patients are vulnerable to severe bleeding before effective hemostasis can be achieved and die before surgery or interventional radiology is performed. To rescue such patients, resuscita-

tive endovascular balloon occlusion of the aorta (REBOA) was introduced for the temporary cessation or limitation of aortic blood flow [3]. REBOA is used as a bridge until definitive hemorrhagic control is achieved [4]. Resuscitative thoracotomy is substantially invasive and has high morbidity [5]; thus, REBOA has received considerable attention [6]. Moreover, the introduction of a 7F or 8F sheath minimizes iatrogenic limb ischemia.

Well-trained staff and well-organized medical resources are required for an effective REBOA procedure. Medical staff should demonstrate proficiency in hemorrhage detection, diagnosis, vascular approach, identification of balloon position, and subsequent prompt hemostatic procedures such as damage control surgery and interventional radiology. Facilities and equipment such as trauma bays, interventional radiology rooms, operating rooms, point-of-care ultrasonography, and portable X-ray equipment should be well-organized. However, training and experience with medical resources are time-consuming. Moreover, experience with REBOA is insufficient in most trauma centers because the procedure is usually performed only in rare situations, such as in patients in severe shock [6,7]. Recently, centers with high REBOA utilization were found to be associated with lower mortality than low-utilization centers [8]. This implies that the learning curve at the hospital level is crucial. However, the learning curve associated with REBOA has not been examined in previous studies. Recently, the learning curve for damage control laparotomy in a Korean regional trauma center was evaluated using cumulative sum (CUSUM) analysis, which is a useful method for monitoring the performance of procedures [9–11]. Here, we investigated the improvement in the quality of the REBOA procedure and associated mortality using CUSUM at the hospital level.

METHODS

Ethics statements

This study was approved by the Institutional Review Board of Cheju Halla General Hospital (No. 2022-L03-01). The requirement for informed consent was waived due to the retrospective nature of the study.

Study design

We reviewed the Korean Trauma Database for records from our trauma center from July 2017 to June 2021 to identify patients with trauma who underwent REBOA. Patients who died before REBOA, who did not undergo balloon inflation, or who underwent balloon insertion into the inferior vena cava due to a retro-

hepatic inferior vena cava injury were excluded from the study. Patient demographic and clinical data including mechanism of injury, age, sex, laboratory findings, vital signs, Injury Severity Score, Abbreviated Injury Scale score, postoperative outcomes, and REBOA-related time parameters were collected and analyzed.

At our trauma center, two dedicated trauma bays, two operating rooms, and one interventional radiology room near the trauma bay were equipped for use by dedicated trauma staff. We divided our study into pre-trauma center (pre-TC; July 2017–February 2020) and TC (February 2020–June 2021) periods. Before February 2020 (the pre-TC period), trauma procedures were performed in the emergency department (ED), where nontrauma patients were also managed. From February 2020 onward (the TC period), all trauma procedures and patients were managed by attending trauma surgeons in a trauma bay. In the pre-TC period, the ultrasonographic image quality was poor due to aging equipment. Surgical instruments for laparotomy or thoracotomy were not prepared in the ED, and surgeons had to bring them from the operating room. In addition, the nursing staff members were not proficient in preparing REBOA kits, and most of them did not even know what REBOA was. The angiography room was located on a different floor from the ED, so patients had to be transferred by elevator. In contrast, in the TC period, point-of-care ultrasonography, a REBOA kit, portable X-ray equipment, surgical equipment for ED laparotomy and ED thoracotomy, and a trauma angiography room next door to the trauma bay were available after the establishment of dedicated trauma facilities. The dedicated nursing staff members were educated regarding the REBOA kit and surgical instruments and became more proficient at preparing them. In the pre-TC period, only three to five dedicated trauma surgeons worked at the trauma center. During the TC period, six to ten trauma surgeons and four emergency medicine faculty members were assigned to the trauma center.

The indications for REBOA were patients with unstable vital signs (systolic blood pressure [SBP] < 90 mmHg) and patients with severe intra-abdominal or pelvic hemorrhage. Femoral arterial puncture for REBOA was performed by a trauma surgeon using point-of-care ultrasonography or a blind method. The surgeon inflated the REBOA balloon by infusing 5 to 20 mL of saline. The balloon position was identified using portable X-ray equipment. When prompt hemostasis was required, ED laparotomy was performed appropriately. For patients with impending cardiac arrest before or after the REBOA procedure, ED thoracotomy was performed. After the return of spontaneous circula-

tion, the aortic clamp used during the thoracotomy was converted to a REBOA setup. A hemostatic procedure was defined as the control of bleeding by laparotomy or angioembolization. The total REBOA occlusion time was measured from the time of initial balloon inflation to that of full deflation. A procedure was considered successful if the REBOA occlusion time was within 60, 90, or 120 minutes; this is because prolonged REBOA occlusion time can induce ischemic injury, which is associated with bowel ischemia, acute kidney injury, and limb ischemia. The primary outcome of our study was mortality, and the secondary outcome was aortic occlusion time. We hypothesized that aortic occlusion time is a surrogate marker of success in REBOA procedures.

Statistical analysis

Continuous data are presented as medians and interquartile ranges, and data were analyzed using the Mann-Whitney U-test. Categorical data are presented as proportions. Proportions were compared using the chi-square test or the Fisher exact test, as appropriate. P-values less than 0.05 were considered to indicate statistical significance. All statistical analyses were performed using Microsoft Excel (Microsoft Corp) and R ver. 4.1.2 (R Foundation for Statistical Computing).

CUSUM analysis

The CUSUM procedure is a graphical method that is widely used for quality monitoring [9–11]. In this study, the CUSUM was calculated as follows: $S_n = (X_i - p_{oi})$, where S_n is the summation of the score, $X_i = 0$ for success (for example, patient survival) and $X_i = 1$ for failure (for example, patient death), and p_{oi} denotes the predicted probability of failure of the procedure. The graph starts at 0 and is plotted from left to right on a horizontal axis. The curve moves up by $1 - p_{oi}$ for every case of failure (penalty) and down by p_{oi} for every case of success (reward) on the cumulative failure graph. The improvement or deterioration of surgical outcomes can be identified intuitively based on the inflection of the CUSUM curve.

RESULTS

This is a case series. From July 2017 to June 2021, the REBOA procedure was performed for 31 trauma patients who were admitted to our hospital. A total of five trauma surgeons performed REBOA, with 20 procedures performed by one surgeon, eight procedures by another surgeon, and one procedure each by the remaining three surgeons. The baseline characteristics and a comparison between the pre-TC and TC periods are summa-

rized in Table 1. Twelve (38.7%) and 19 patients (61.3%) were admitted during the pre-TC and TC periods, respectively. Twenty-nine patients (93.55%) experienced blunt trauma. The initial SBP levels of 13 patients (41.9%) could not be determined. Seven patients (22.6%) underwent prehospital cardiopulmonary resuscitation (CPR). Nine (29.0%) and eight patients (25.8%) underwent thoracotomy and laparotomy, respectively, in the ED. The median time from admission to REBOA was 31.0 minutes (interquartile range [IQR], 18.0–67.0 minutes). The median total REBOA occlusion time was 60.0 minutes (IQR, 47.5–90.5 minutes). In 30 patients (96.8%), the balloon used during REBOA was placed in zone 1 (above the celiac axis), while it was placed in zone 3 (between the inferior mesenteric artery and the iliac bifurcation) in one patient (3.2%). Twelve patients (38.7%) survived the REBOA procedure. No significant differences were present in age, sex, SBP, Injury Severity Score, or Abbreviated Injury Scale score between the pre-TC and TC periods. The median REBOA occlusion time was shorter in the TC than in the pre-TC period, although the difference was not statistically significant (62.5 minutes [IQR, 44.0–134.5 minutes] in the pre-TC period vs. 53.0 minutes [IQR, 47.5–78.5 minutes] in the TC period, $P = 0.465$). The survival rate was higher in the TC than in the pre-TC period, although the difference was also not significant (three patients [25.0%] in the pre-TC period vs. nine patients [47.4%] in the TC period, $P = 0.386$). Table 2 shows a summarized comparison between nonsurvivors and survivors. The initial SBP was significantly lower in the nonsurvivor group. Nonsurvivors were significantly more likely to have undergone CPR at ED, ED thoracotomy, and ED laparotomy. However, no patient in the survivor group underwent CPR at ED, ED thoracotomy, or ED laparotomy.

The CUSUM failure graph for survival is shown in Fig. 1. The CUSUM failure graphs for aortic occlusion time within 60 minutes, 90 minutes, and 120 minutes are shown in Figs. 2, 3, and 4, respectively. All CUSUM graphs showed an upward slope during the pre-TC period. At the 17th patient (indicated by arrows) in the TC period, a downward inflection was observed and a downward slope was observed after 17th case.

DISCUSSION

To our knowledge, this is the first study to illustrate the learning curve associated with REBOA using CUSUM analysis. The results indicated that the quality of the REBOA procedure improved. The accumulation of experience with the REBOA procedure may enhance performance. Additionally, we believe that the

Table 1. Comparison between pre-TC and TC period

Variable	Total	Pre-TC	TC	P-value
No. of patients	31 (100)	12 (38.7)	19 (61.3)	
Age (yr)	45.0 (35.0–57.5)	49.0 (35.5–59.0)	42.0 (32.0–54.0)	0.612
Injury type				>0.999
Blunt	29 (93.5)	11 (91.7)	18 (94.7)	
Penetrating	2 (6.5)	1 (8.3)	1 (5.3)	
Male sex	24 (77.4)	9 (75.0)	15 (78.9)	>0.999
Injury Severity Score	34.0 (25.0–44.0)	33.5 (18.0–52.5)	34.0 (28.0–36.5)	0.919
Abbreviated Injury Scale				
Head and neck	3.0 (0–4.5)	2.5 (0–5.0)	3.0 (0–4.0)	0.801
Face	0 (0–0.5)	0 (0–0.5)	0 (0–1.0)	0.771
Chest	3.0 (0.5–3.0)	2.0 (0–3.0)	3.0 (2.0–3.0)	0.291
Abdomen	3.0 (2.0–4.0)	3.0 (2.5–4.0)	2.0 (1.0–4.0)	0.347
Pelvis	0 (0–3.0)	0 (0–3.5)	2.0 (0–3.0)	0.427
Extremity	0 (0–1.0)	0 (0–1.0)	1.0 (0–1.0)	0.151
Initial systolic blood pressure (mmHg)				0.206
≥100	5 (16.1)	4 (33.3)	1 (5.3)	
80–99	5 (16.1)	2 (16.7)	3 (15.8)	
50–79	8 (25.8)	2 (16.7)	6 (31.6)	
Not checked	13 (41.9)	4 (33.3)	9 (47.4)	
Prehospital CPR	7 (22.6)	3 (25.0)	4 (21.1)	>0.999
CPR at the ED	15 (48.4)	7 (58.3)	8 (42.1)	0.609
Balloon type of REBOA				0.032
12F (Reliant™)	4 (12.9)	4 (33.3)	0	
7F (Rescue™)	27 (87.1)	8 (66.7)	19 (100)	
Target zone				>0.999
Zone 1	30 (96.8)	12 (100)	18 (94.7)	
Zone 3	1 (3.2)	0	1 (5.3)	
Identification of REBOA balloon position	15 (48.4)	7 (58.3)	8 (42.1)	0.609
Time from admission to REBOA (min)	31.0 (18.0–67.0)	64.5 (31.0–93.0)	21.0 (14.5–43.5)	0.007
REBOA occlusion time (min)	60.0 (47.5–90.5)	62.5 (44.0–134.5)	53.0 (47.5–78.5)	0.465
Within 60 min	17 (54.8)	6 (50.0)	11 (57.9)	0.952
Within 90 min	23 (74.2)	7 (58.3)	16 (84.2)	0.237
Within 120 min	24 (77.4)	7 (58.3)	17 (89.5)	0.114
ED thoracotomy	9 (29.0)	2 (16.7)	7 (36.8)	0.424
ED laparotomy	8 (25.8)	3 (25.0)	5 (26.3)	>0.999
Angioembolization	4 (12.9)	0	4 (21.1)	0.249
Laparotomy	19 (61.3)	9 (75.0)	10 (52.6)	0.386
Preperitoneal pelvic packing	2 (6.5)	1 (8.3)	1 (5.3)	>0.999
Hemostasis (surgery or angioembolization)	20 (64.5)	8 (66.7)	12 (63.2)	>0.999
PRBC transfusion within 24 hr (unit)	11.0 (6.0–17.0)	13.0 (5.5–17.5)	11.0 (6.0–15.5)	0.951
FFP transfusion within 24 hr (unit)	5.0 (2.5–10.0)	6.0 (2.5–10.0)	5.0 (2.5–11.0)	>0.999
Platelet transfusion within 24 hr (unit)	0 (0–8.0)	0 (0–6.5)	0 (0–8.0)	0.751
PRBC transfusion within 4 hr (unit)	8.0 (6.0–14.5)	10.0 (5.5–15.5)	7.0 (6.0–13.5)	0.902
FFP transfusion within 4 hr (unit)	4.0 (2.0–7.0)	5.0 (2.0–7.0)	4.0 (2.0–7.0)	0.790
Platelet transfusion within 4 hr (unit)	0 (0–0)	0 (0–0)	0 (0–0)	0.739
Hemodialysis due to AKI	1 (3.2)	1 (8.3)	0	0.814
Limb ischemia	0	0	0	-
Survival (in-hospital)	12 (38.7)	3 (25.0)	9 (47.4)	0.386
Cause of mortality				0.396
Central nervous system	3 (9.7)	2 (16.7)	1 (5.3)	
Hypovolemic	15 (48.4)	7 (58.3)	8 (42.1)	
Multiple organ dysfunction syndrome	1 (3.2)	0	1 (5.3)	

Values are presented as number (%) or median (interquartile range).

TC, trauma center; CPR, cardiopulmonary resuscitation; ED, emergency department; REBOA, resuscitative endovascular balloon occlusion of aorta; PRBC, packed red blood cell; FFP, fresh frozen plasma; AKI, acute kidney injury.

Table 2. Comparison between nonsurvivor and survivor (n=31)

Variable	Nonsurvivor (n=19)	Survivor (n=12)	P-value
Group			0.386
Pre-TC	9 (47.4)	3 (25.0)	
TC	10 (52.6)	9 (75.0)	
Age (yr)	48.0 (32.0–60.5)	39.5 (35.0–49.5)	0.310
Injury type			>0.999
Blunt	18 (94.7)	11 (91.7)	
Penetrating	1 (5.3)	1 (8.3)	
Male sex	15 (78.9)	9 (75.0)	>0.999
Injury Severity Score	34.0 (27.0–49.0)	33.5 (21.0–34.0)	0.166
Abbreviated Injury Scale			
Head and neck	2.0 (0–5.0)	3.5 (0–4.0)	0.784
Face	0 (0–0)	0 (0–2.0)	0.076
Chest	3.0 (1.5–4.0)	2.5 (0–3.0)	0.157
Abdomen	3.0 (2.0–4.0)	3.0 (2.0–3.0)	0.530
Pelvis	0 (0–3.5)	3.0 (0–3.0)	0.321
Extremity	1.0 (0–1.0)	0 (0–1.0)	0.151
Initial systolic blood pressure (mmHg)			0.001
≥100	3 (15.8)	2 (16.7)	
80–99	1 (5.3)	7 (58.3)	
50–79	2 (10.5)	3 (25.0)	
Not checked	13 (68.4)	0	
Prehospital CPR	7 (36.8)	0	0.051
CPR at ED	15 (78.9)	0	>0.001
Balloon type of REBOA			0.958
12F (Reliant™)	3 (15.8)	1 (8.3)	
7F (Rescue™)	16 (84.2)	11 (91.7)	
Target zone			0.814
Zone 1	19 (100)	11 (91.7)	
Zone 3	0	1 (8.3)	
Identification of REBOA balloon position	8 (42.1)	7 (58.3)	0.609
Time from admission to REBOA (min)	31.0 (19.5–70.0)	34.0 (15.0–66.0)	0.626
REBOA occlusion time (min)	60.0 (43.5–105.5)	64.0 (48.0–83.5)	0.855
Within 60 min	11 (57.9)	6 (50.0)	0.952
Within 90 min	13 (68.4)	10 (83.3)	0.615
Within 120 min	14 (73.7)	10 (83.3)	0.853
ED thoracotomy	9 (47.4)	0	0.015
ED laparotomy	8 (42.1)	0	0.029
Angioembolization	1 (5.3)	3 (25.0)	0.295
Laparotomy	12 (63.2)	7 (58.3)	>0.999
Preperitoneal pelvic packing	2 (10.5)	0	0.681
Hemodialysis due to AKI	1 (5.3)	0	>0.999
Limb ischemia	0	0	-
Hemostasis (surgery or angioembolization)	12 (63.2)	8 (66.7)	>0.999
PRBC transfusion within 24 hr (unit)	11.0 (6.0–17.5)	11.5 (6.5–15.5)	0.968
FFP transfusion within 24 hr (unit)	4.0 (2.0–9.5)	7.0 (4.5–12.5)	0.186
Platelet transfusion within 24 hr (unit)	0 (0–0)	7.0 (1.5–10.0)	0.007
PRBC transfusion within 4 hr (unit)	10.0 (5.5–16.5)	6.5 (6.0–12.0)	0.595
FFP transfusion within 4 hr (unit)	4.0 (2.0–5.5)	4.5 (2.0–8.0)	0.595
Platelet transfusion within 4 hr (unit)	0 (0–0)	0 (0–0)	0.274
Cause of mortality			<0.001
Central nervous system	3 (15.8)	0	
Hypovolemic	15 (78.9)	0	
Multiple organ dysfunction syndrome	1 (5.3)	0	

Values are presented as number (%) or median (interquartile range).

TC, trauma center; CPR, cardiopulmonary resuscitation; ED, emergency department; REBOA, resuscitative endovascular balloon occlusion of aorta; AKI, acute kidney injury; PRBC, packed red blood cell; FFP, fresh frozen plasma.

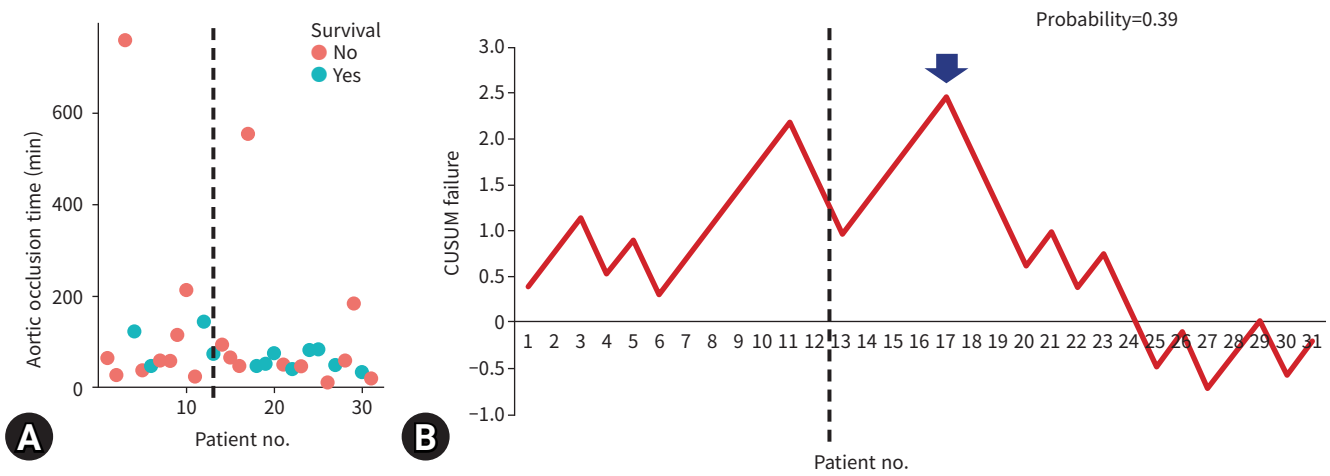


Fig. 1. Cumulative sum (CUSUM) failure graph for patient survival. (A) The scatter plot of aortic occlusion time and (B) CUSUM graph. The dotted line denotes the establishment of a regional trauma center (pre-trauma center period, patient no. 1–12; trauma center period, patient no. 13–31).

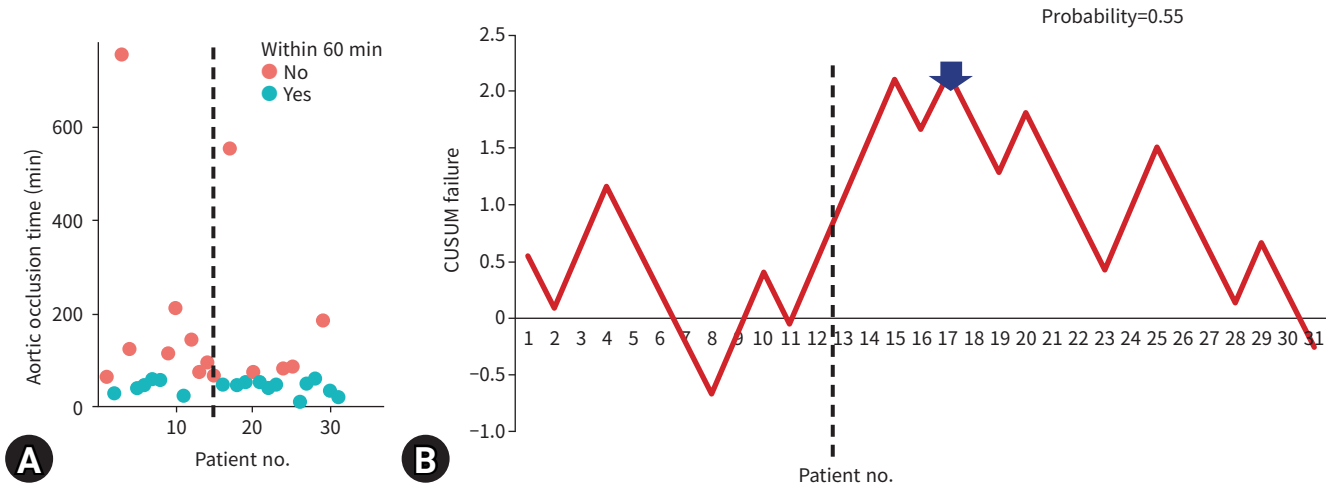


Fig. 2. Cumulative sum (CUSUM) failure graph for aortic occlusion time ≤ 60 minutes. (A) The scatter plot of aortic occlusion time and (B) CUSUM graph. The dotted line denotes the establishment of a regional trauma center (pre-trauma center period, patient no. 1–12; trauma center period, patient no. 13–31).

establishment of trauma centers plays an important role in improving the skill and knowledge of faculty and the number of dedicated facility members, which helps improve the quality of the procedure. CUSUM-based monitoring appears to be useful for the REBOA procedure. REBOA is generally performed in extremely rare situations; therefore, achieving proficiency may be difficult. We believe that the outcomes, mortality rate, and aortic occlusion time are potential indicators of the performance of REBOA. An improvement in aortic occlusion time represents prompt hemostasis and the reduction of ischemia.

Although enthusiasm has grown regarding the use of REBOA for trauma patients with severe hemorrhage, the procedure’s indications and outcomes, including mortality and morbidity, are

controversial [6,12]. In a recent systematic review and meta-analysis [6], REBOA was associated with lower mortality than ED thoracotomy, whereas no significant difference was observed between patients who underwent REBOA and those who did not. However, this meta-analysis included a limited number of studies (three and five studies comparing REBOA with ED thoracotomy and non-REBOA, respectively), and all included studies were observational. Thus, the exact effect size is unclear due to substantial selection bias. In a retrospective cohort study using the American College of Surgeons National Trauma Database with propensity score matching [7], REBOA was associated with increased mortality, even after adjustment for measured confounders. The authors reported that REBOA was not associated with

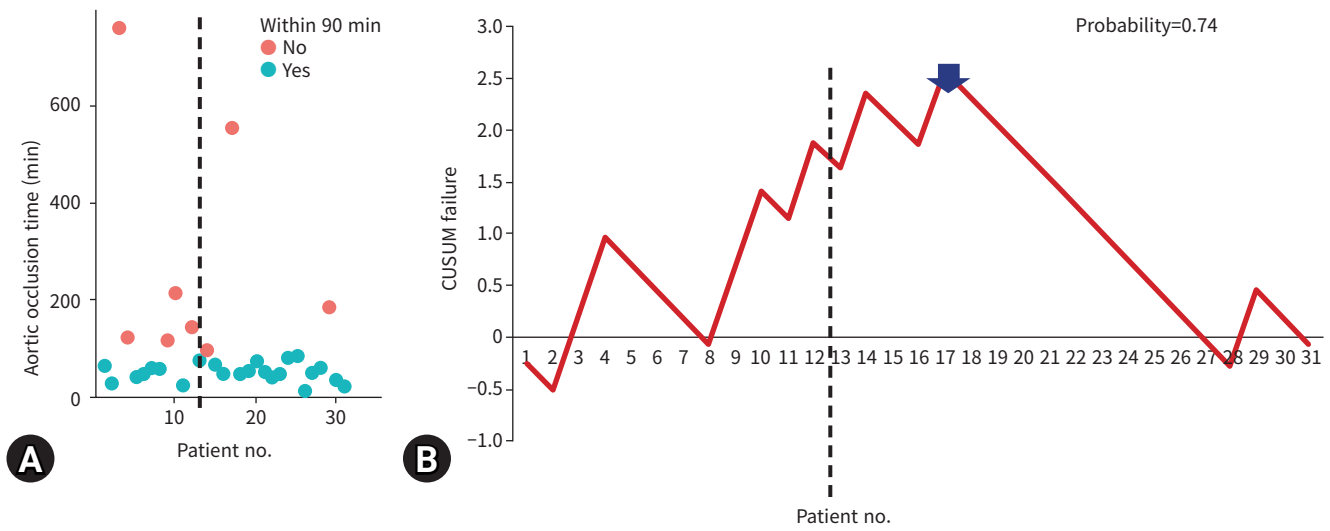


Fig. 3. Cumulative sum (CUSUM) failure graph for aortic occlusion time ≤ 90 minutes. (A) The scatter plot of aortic occlusion time and (B) CUSUM graph. The dotted line denotes the establishment of a regional trauma center (pre-trauma center period, patient no. 1–12; trauma center period, patient no. 13–31).

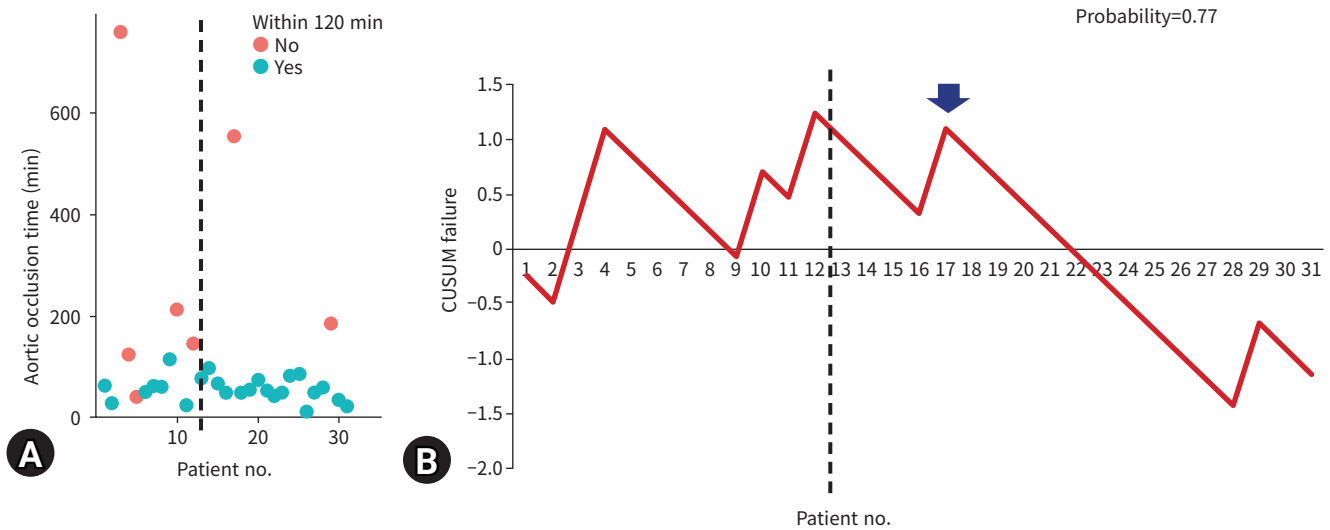


Fig. 4. Cumulative sum (CUSUM) failure graph for aortic occlusion time ≤ 120 minutes. (A) The scatter plot of aortic occlusion time and (B) CUSUM graph. The dotted line denotes the establishment of a regional trauma center (pre-trauma center period, patient no. 1–12; trauma center period, patient no. 13–31).

acute kidney injury or amputation. In our study, only one patient underwent hemodialysis, and none underwent lower extremity amputation. However, these complications are rare. In a recent review on the opinions of trauma providers regarding REBOA, interest was revealed to be widespread, but the need for training persists [13]. Thus, clinical results may need to be re-evaluated after the teaching of technical skills and dissemination of information regarding indications and complications. In our study, all patients with initial SBP levels that could not be assessed died.

Further study is warranted for such patients, and more rigorous indication may be needed.

In a 5-year retrospective analysis based on the American Association for the Surgery of Trauma’s Aortic Occlusion in Resuscitation for Trauma and Acute Care Surgery (AORTA) multi-institutional database [8], a center was defined as high-volume if it had more than 30 cases of REBOA over 5 years, low-volume if it had less than 10, and average-volume if it had 11 to 30 cases. The results indicated that survival was higher in high-volume

centers than in low-volume ones [8]. From this perspective, our trauma center can be regarded as high-volume because over 5 years, REBOA was performed for 31 patients. Moreover, we used CUSUM analysis to demonstrate that the performance of the REBOA procedure improved after the 17th case. We believe that this study provides new insights for trauma surgeons. REBOA is a bridge procedure used prior to achieving definitive hemostasis. REBOA requires the prompt availability of medical staff and facilities such as a proficient trauma or vascular surgeon, well-trained nurses, point-of-care ultrasonography and portable X-ray equipment, a vascular access device, a trauma bay, an operating room, an interventional radiology room, and a REBOA kit [4]. Harmoniously organizing these components may be time-consuming. At our trauma center, more than 17 cases were needed to achieve proficiency in the REBOA procedure. We believe that assigning dedicated medical staff and facilities via trauma centers may promote such improvements. Such organization may enhance the proficiency of staff and the effectiveness of the overall trauma system. Trauma surgeons may also improve their understanding of REBOA after a trauma center has been established.

Our study had several limitations. First, it was retrospective and observational. However, no randomized controlled trials have been conducted regarding REBOA. Second, the single-cohort nature and small sample size of the study are crucial limitations, and larger-scale prospective studies are needed. Third, the most important limitation is the variation in the skill and experience of the surgeons. This may have impeded consistency in proficiency in the REBOA procedure. However, investigation at the hospital level is crucial because REBOA requires a multidisciplinary approach. Fourth, conventional statistical methods showed no statistically significant difference between the pre-TC and TC periods, although mortality and occlusion time differed non-significantly between them. This lack of significant findings may have been due to the small number of patients ($n=31$) studied. However, acquiring sufficient datasets is very difficult because REBOA is rarely performed. Therefore, CUSUM monitoring is a potential alternative method. Fifth, the contribution of the establishment of trauma centers to the improvement in REBOA outcomes remains ambiguous. Further studies are required to confirm this hypothesis. However, we observed that the performance of the procedure improved after the 17th patient. This may provide meaningful insights to trauma surgeons who wish to begin performing the REBOA procedure. Sixth, we could not perform risk-adjusted CUSUM analysis because of the small sample size. In a previous study, a risk-adjust-

ed CUSUM methodology was used to adjust for individual risk and minimize selection bias based on a multivariable logistic regression model [9,11]. However, such models are statistically unstable when the sample size is small [14]. Future large-scale studies are required. Seventh, we postulated that aortic occlusion time is a surrogate marker of success in REBOA procedures. No evidence is available regarding this issue. However, guidelines recommend balloon deflation as soon as possible [4]. Prolonged occlusion time induces ischemic damage and reducing occlusion time may require a well-organized trauma system and proficient staff. For example, prompt detection of the balloon position, early preparation for surgery, or angioembolization may be needed. We hope that this can provide new insight for researchers. Finally, we initially attempted partial REBOA (5 to 10 mL of balloon inflation) and added fluids according to the patient's status. However, unfortunately, our medical records included no records of the inflating fluid in some patients. Thus, we could not distinguish partial from total REBOA. We could not identify even intermittent REBOA due to the limited medical records available.

In conclusion, the performance of the REBOA procedure in terms of mortality and aortic occlusion time improved after the 17th case. CUSUM analysis may be useful for monitoring the procedure when the sample size of the cohort is not sufficient to expect statistical stability. However, further large-scale prospective studies are warranted to confirm the true effect size.

NOTES

Conflicts of interest

The authors have no conflicts of interest to declare.

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None.

Data sharing statement

The data of this article are available from the corresponding author upon reasonable request.

Author contributions

Conceptualization: HC, WSK; Data curation: all authors; Formal analysis: WSK; Methodology: HC, WSK; Project administration: WSK; Visualization: WSK; Writing—original draft: HC, WSK; Writing—review & editing: all authors. All authors read and approved the final manuscript.

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