Pre-hospital advanced airway management for adults with out-of-hospital cardiac arrest: nationwide cohort study

Junichi Izawa, assistant professor, research fellow,1,2 Sho Komukai, assistant professor,3 Koichiro Gibo, staff physician/biostatistician,4 Masashi Okubo, assistant professor,5 Kosuke Kiyohara, assistant professor,6 Chika Nishiyama, senior lecturer,7 Takeyuki Kiguchi, assistant professor,8 Tasuku Matsuyama, assistant professor,9 Takashi Kawamura, professor,9 Taku Iwami, professor,9 Clifton W Callaway, professor,5 and Tetsuhisa Kitamura, assistant professor10

1 Department of Anesthesiology, The Jikei University School of Medicine, Tokyo 105-8491, Japan
2 Center for Critical Care Nephrology, Department of Critical Care Medicine, University of Pittsburgh, Pittsburgh, PA, USA
3 Division of Biomedical Statistics, Department of Integrated Medicine, Graduate School of Medicine, Osaka University, Osaka, Japan
4 Department of Emergency Medicine, Okinawa Chubu Hospital, Okinawa, Japan
5 Department of Emergency Medicine, University of Pittsburgh, Pittsburgh, PA, USA
6 Department of Food Science, Otsuma Women’s University, Tokyo, Japan
7 Department of Critical Care Nursing, Kyoto University Graduate School of Human Health Science, Kyoto, Japan
8 Kyoto University Health Service, Kyoto, Japan
9 Department of Emergency Medicine, Kyoto Prefectural University of Medicine, Kyoto, Japan
10 Division of Environmental Medicine and Population Sciences, Department of Social and Environmental Medicine, Graduate School of Medicine, Osaka University, Osaka, Japan

Correspondence to: J Izawa izawa.junichi.87r@st.kyoto-u.ac.jp

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Abstract

Objective
To determine survival associated with advanced airway management (AAM) compared with no AAM for adults with out-of-hospital cardiac arrest.

**Design**

Cohort study between January 2014 and December 2016.

**Setting**

Nationwide, population based registry in Japan (All-Japan Utstein Registry).

**Participants**

Consecutive adult patients with out-of-hospital cardiac arrest, separated into two sub-cohorts by their first documented electrocardiographic rhythm: shockable (ventricular fibrillation or pulseless ventricular tachycardia) and non-shockable (pulseless electrical activity or asystole). Patients who received AAM during cardiopulmonary resuscitation were sequentially matched with patients at risk of AAM within the same minute on the basis of time dependent propensity scores.

**Main outcome measures**

Survival at one month or at hospital discharge within one month.

**Results**

Of the 310,620 patients eligible, 8459 (41.2%) of 20,516 in the shockable cohort and 121,890 (42.0%) of 290,104 in the non-shockable cohort received AAM during cardiopulmonary resuscitation. After time dependent propensity score sequential matching, 16,114 patients in the shockable cohort and 236,042 in the non-shockable cohort were matched at the same minute. In the shockable cohort, survival did not differ between patients with AAM and those with no AAM: 1546/8057 (19.2%) versus 1500/8057 (18.6%) (adjusted risk ratio 1.00, 95% confidence interval 0.93 to 1.07). In the non-shockable cohort, patients with AAM had better survival than those with no AAM: 2696/118,021 (2.3%) versus 2127/118,021 (1.8%) (adjusted risk ratio 1.27, 1.20 to 1.35).

**Conclusions**

In the time dependent propensity score sequential matching for out-of-hospital cardiac arrest in adults, AAM was not associated with survival among patients with shockable rhythm, whereas AAM was associated with better survival among patients with non-shockable rhythm.

**Introduction**

Basic life support, which consists of early cardiopulmonary resuscitation and defibrillation with automated external defibrillators, improves outcomes in patient with out-of-hospital cardiac arrest. Provision of oxygen and ventilation of the lungs via bag-valve-mask ventilation, supraglottic airway placement, or endotracheal intubation are important resuscitation skills for emergency medical services (EMS) personnel. Supraglottic airway placement and endotracheal intubation constitute advanced airway
management (AAM), which secures better protection for the airway than bag-valve-mask ventilation does. Although AAM is a common practice, the risk-benefit ratio of AAM during pre-hospital resuscitation is unclear.\textsuperscript{5,6,7}

In observational studies assessing intra-cardiac arrest interventions, “resuscitation time bias” (that is, the fact that patients undergoing longer resuscitation tend to receive more interventions) is a crucial bias.\textsuperscript{8} Because of this bias effect, two observational studies that evaluated the impact of adrenaline (epinephrine) on patients with out-of-hospital cardiac arrest, using the same national database in Japan but different statistical approaches, reported conflicting findings.\textsuperscript{9,10} One study used traditional propensity score matching,\textsuperscript{9} whereas the other used time dependent propensity score sequential matching, accounting for resuscitation time bias and timing of administration of adrenaline.\textsuperscript{10} A recent, large randomised controlled trial showed better survival in the adrenaline group than the placebo group,\textsuperscript{11} and these findings were closer to the results of the second observational study from Japan.\textsuperscript{10,12} This confirms the importance of overcoming resuscitation time bias in observational studies.\textsuperscript{12} Previous studies of in-hospital cardiac arrest have evaluated the effect of intubation while accounting for resuscitation time bias and timing of intubation,\textsuperscript{13,14} but no studies among patients with out-of-hospital cardiac arrest have considered the timing of AAM.

In Japan, a nationwide registry includes approximately 368,000 adult patients with out-of-hospital cardiac arrest with available data on the timing of pre-hospital AAM between 2014 and 2016. This study investigated whether AAM by EMS personnel increased survival for adult patients with out-of-hospital cardiac arrest, adjusting for time dependent interventions and covariates.

**Methods**

**Study design and setting**

We used the All-Japan Utstein Registry database to do a cohort study with time dependent propensity score sequential matching. The All-Japan Utstein Registry of the Fire and Disaster Management Agency is a prospective, population based, nationwide registry of out-of-hospital cardiac arrest launched in January 2005, which records data according to the international Utstein-style reporting system.\textsuperscript{15} Details of the registry have been described previously.\textsuperscript{16} The registry started collecting time stamp data for successful AAM by out-of-hospital cardiac arrest personnel in January 2014. Cardiac arrest was defined as the cessation of cardiac mechanical activity, as confirmed by the absence of signs of circulation.\textsuperscript{15} Each EMS authority submitted anonymised data.

**Emergency medical services system in Japan**

The EMS system in Japan was previously described elsewhere.\textsuperscript{5,16} Briefly, pre-hospital life support is provided 24 hours each day by a fire station based EMS system. Among EMS personnel, specially trained emergency care providers known as emergency life saving technicians are authorised to use automated external defibrillators. With support and direction from online medical control, emergency life saving technicians are also authorised to perform advanced life support such as administering adrenaline and placing advanced airway for patients with out-of-hospital cardiac arrest. In Japan, EMS providers are generally not permitted to terminate resuscitation in the field, and all patients on whom resuscitation is
attempted are transported to hospitals. Resuscitation is not attempted only under particular conditions (for example, decapitation, incineration, decomposition, rigor mortis, or dependent cyanosis). Although all emergency life saving technicians can use supraglottic airway devices such as laryngeal tubes or laryngeal masks, only specially trained, certified emergency life saving technicians are permitted to perform endotracheal intubation under direction from online medical control. These technicians are allowed to perform endotracheal intubation but only during cardiac arrest (that is, they are not allowed to intubate after return of spontaneous circulation). Treatments for cardiac arrest were based on the Japanese cardiopulmonary resuscitation guidelines,\textsuperscript{17} which are based on the guidelines of the consensus of the International Liaison Committee on Resuscitation.\textsuperscript{18}

**Study participants**

We screened the data of all consecutive cases of out-of-hospital cardiac arrest between January 2014 and December 2016. We included patients according to the following inclusion criteria: age 18 years or over; cardiac arrest before arrival of EMS personnel; cardiac arrest for which EMS personnel attempted cardiopulmonary resuscitation; and cardiac arrest attended by an emergency life saving technician. The exclusion criteria were outliers with regard to age (that is, older than 120 years); unknown first documented electrocardiographic rhythm; missing or contradictory data (negative values) in time dependent variables (including time from cardiopulmonary resuscitation by EMS personnel to AAM at the scene, defibrillation at the scene, administration of adrenaline at the scene, and return of spontaneous circulation at the scene; time from call to cardiopulmonary resuscitation by EMS personnel; and time from cardiopulmonary resuscitation by EMS personnel to arrival at hospital); or time from call to cardiopulmonary resuscitation by EMS personnel 30 minutes or more.

**Data collection and quality control**

Data were collected prospectively on resuscitation related factors including age and sex of the patient, type of witness status (by family members, friends, colleagues, passers-by, or others), type of bystander initiated cardiopulmonary resuscitation (chest compressions only without rescue breathing or conventional cardiopulmonary resuscitation with rescue breathing), instruction on cardiopulmonary resuscitation by dispatcher, shock delivery by public access automated external defibrillators, presence of emergency life saving technicians, pre-hospital involvement of physician, and first documented electrocardiographic rhythm (ventricular fibrillation, pulseless ventricular tachycardia, pulseless electrical activity, or asystole), as well as return of spontaneous circulation before arrival at hospital, survival, and functional status at one month after the event. On the basis of the consensus guidelines,\textsuperscript{15,19} return of spontaneous circulation was defined according to a clinical assessment that shows signs of life comprising a palpable pulse or generation of a blood pressure. Origin of cardiac arrest (medical or non-medical) was also documented; arrest was presumed to be of medical origin unless it was caused by trauma, drug overdose, drowning, electrocution, or asphyxia, according to the current Utstein-style template.\textsuperscript{19} Prefecture and year, month, day, and time when EMS received the emergency call were documented. The times of the receipt of the emergency call, the initiation of cardiopulmonary resuscitation by EMS personnel, the first defibrillation by EMS personnel, the first administration of adrenaline by EMS personnel, return of spontaneous
circulation before arrival at hospital, the successful achievement of AAM by EMS personnel, and arrival at hospital were recorded according to the times on the clock used by each emergency medical services system.

All survivors were followed for up to one month after out-of-hospital cardiac arrest by the EMS providers who had provided their emergency care. Functional outcomes were determined by the physician responsible for the care of the patient by a follow-up interview at one month after successful resuscitation, with the use of the Cerebral Performance Category scale as follows: 1, good cerebral performance; 2, moderate cerebral disability; 3, severe cerebral disability; 4, coma or vegetative state; and 5, death or brain death. When the patient was discharged from the hospital before one month, the score at discharge was recorded but the database did not include the information about when the patients were discharged from the hospital.

EMS personnel filled out the study data form in cooperation with the physician in charge of the patient, and the data were stored in the registry system on the Fire and Disaster Management Agency database server. The data were checked for consistency by the computer system and were confirmed by the Fire and Disaster Management Agency. If the data form was incomplete, the Fire and Disaster Management Agency returned it to the respective fire station where it was then completed.

**Outcomes**

The primary outcome was survival at one month or at discharge from hospital within one month. The secondary outcome was favourable functional survival at one month or at hospital discharge within one month. We defined favourable functional survival as a Cerebral Performance Category scale of 1 or 2.

**Time dependent propensity scores and sequential matching**

The main exposure was the successful achievement of AAM by EMS personnel. We separated the patients into two sub-cohorts according to their first documented electrocardiographic rhythm—shockable (ventricular fibrillation or pulseless ventricular tachycardia) or non-shockable (pulseless electrical activity or asystole)—because the current international guidelines propose two different algorithms according to the rhythm, with different suggestions regarding the timing of AAM. To adjust for all measured potential confounders, we did time dependent propensity score sequential matching analysis in each original cohort. To estimate the time dependent propensity scores for receipt of AAM during cardiopulmonary resuscitation, we applied the Fine-Gray model in the presence of competing risk with time dependent covariates. In the regression model, we dealt with pre-hospital return of spontaneous circulation before AAM as the competing risk. We also dealt with arrival at hospital as censoring in the model. The Cox regression model with or without time dependent covariates was used in previous cardiac arrest studies with time dependent propensity score sequential matching. However, we regarded pre-hospital return of spontaneous circulation before AAM as informative censoring in the time dependent propensity score model, because AAM during cardiopulmonary resuscitation never occurs after return of spontaneous circulation except in rare cases involving re-arrest. If we regarded pre-hospital return of spontaneous circulation as non-informative censoring, the propensity scores in patients with return of spontaneous circulation before AAM would be overestimated. The time dependent covariates were defibrillation by EMS personnel and administration of adrenaline by EMS personnel. The time
independent covariates in the propensity score predicting model are shown in table 1 and described in the supplementary methods. We included prefecture categories according to the proportions of patients who received supraglottic airway placement (<15%, 15% to <30%, 30% to <45%, ≥45%) and endotracheal intubation (<5%, 5% to <10%, 10% to <15%, ≥15%) by EMS personnel as prefecture elements as applied in a previous study. Although we adopted the same variables in the time dependent propensity score model for both the shockable and non-shockable original cohorts, we calculated the time dependent propensity scores separately in each rhythm cohort.

On the basis of the predicted time dependent propensity scores, we separately and sequentially propensity score matched each patient receiving AAM at any given minute (from minute 0 to minute 59) with a patient who was at risk of receiving AAM within the same minute. Sequential matching attenuates resuscitation time bias. In sequential matching, at risk patients included those who were still undergoing cardiopulmonary resuscitation at the scene and had not yet received AAM before or within the same minute. In addition, replacement of unexposed patients was possible—that is, matched controls with no AAM in a risk set at each timing were allowed to match again later until they received AAM. At risk patients therefore also included patients who received AAM later, as the matching was not dependent on future events.

We sequentially matched patients with AAM to those without AAM who had the nearest propensity score in a one to one fashion. We set the calliper width for nearest neighbour matching at 0.2 standard deviations of the propensity score in the logit scale, as recommended. We calculated standardised differences to evaluate the balance of variables in each predicted propensity score matched cohort. We first regarded standardised differences less than 0.1 as having well matched balance, but we could not achieve the value for the variable of “defibrillation before matching” in the shockable cohort even with a very narrow calliper width (0.001). When we attempted to achieve better balancing of standardised differences (<0.1) by setting the calliper width much narrower (<0.001), we lost a large number of patients. In the end, we decided to avoid losing these patients by using a tight range of target and chose a value of 0.25 rather than 0.1 of standardised differences, as some statisticians have suggested, before doing our final analyses.

After sequential matching, we calculated the time from cardiopulmonary resuscitation by EMS personnel to sequential matching.

**Statistical analysis**

We did all the analyses separately for each of the two cohorts defined by the first documented electrocardiographic rhythm. We presented data as medians with an interquartile range for continuous variables and as proportions for categorical variables. We used standardised differences for the description of the data in both the original cohorts and the sequentially matched cohorts. We estimated unadjusted and adjusted risk ratios on the primary and secondary outcomes by multivariable regression to evaluate overall outcomes in the original cohorts (see the supplementary methods). Matched controls within a risk set stratum in each minute were independent from each other, whereas some controls in the matched cohort after jointing of all strata could be duplicated and were no longer independent. In the outcome analysis stage after matching, we handled this problem by adjusting for frequency weights that indicate the number of duplicated controls. Among all risk set strata in each minute, treated patients were paired up with their respective controls, creating within pair correlations. At the outcome analysis stage, therefore, we used
generalised estimating equations to estimate the unadjusted and adjusted risk ratios for the primary and secondary outcomes, considering within pair correlation of sequential matching at the same timing.27 We adjusted risk ratios for defibrillation before matching, adrenaline administration before matching, and all the time independent covariates in the propensity score predicting model.

To confirm the consistency of the effects between the types of AAM, we stratified the patients into two groups according to each patient’s type of AAM (supraglottic airway placement or endotracheal intubation) and calculated adjusted risk ratios with the same multivariable model used in the primary analysis. In this stratified analysis, we picked the pairs of supraglottic airway placement and the pairs of endotracheal intubation out of the sequentially matched cohorts. Therefore, patients were still nested within pairs at the same timing, as we selected the control from within each pair of patients—that is, the sensitively matched pairs in each risk set stratum were not collapsed even after the stratification. As a sensitivity analysis, we also did traditional, non-time dependent propensity score matching analysis as described in the supplementary methods.

We used R (R Foundation for Statistical Computing, version 3.4.3) for all statistical analyses. We did not impute missing data because no information was missing in any key variables after we excluded patients according to the exclusion criteria described above. We calculated all risk ratios with their 95% confidence intervals by log-binomial regression in generalised estimating equations with the robust variance estimators. All tests were two tailed; we regarded P values of less than 0.05 as statistically significant. The entire author group outlined and agreed on the statistical analysis plan before we did any analyses, unless stated otherwise. The other details are described in the supplementary methods.

Patient and public involvement

No patients were involved in setting the research question or the outcome measures, nor were they involved in developing plans for recruitment, design, or implementation of the study. No patients were asked to advise on interpretation or writing up of results. There are no plans to disseminate the results of the research to study participants or the relevant patient community.

Results

Original cohort and outcomes

We screened all 372,926 cases of out-of-hospital cardiac arrest from the All-Japan Utstein Registry during 2014–16. We divided the 310,620 adult patients who met our eligibility criteria into two sub-cohorts according to each patient’s first documented electrocardiographic rhythm; 20,516 had shockable rhythm and 290,104 had non-shockable rhythm (fig 1). Table 1 shows the characteristics of the patients in the original cohort. Among the patients with shockable rhythm, 8459 (41.2%) received AAM; among patients with non-shockable rhythm, 121,890 (42.0%) received AAM. Among patients who received AAM, the median time from cardiopulmonary resuscitation by EMS personnel to AAM was 8 (interquartile range 5–12) minutes in the shockable cohort and 8 (5–12) minutes in the non-shockable cohort. AAM was associated with worse survival in both rhythm cohorts, with multivariable regression adjusting for
covariates: 1625/8459 (19.2%) versus 4277/12 057 (35.5%), adjusted risk ratio 0.61 (95% confidence interval 0.58 to 0.65) in the shockable cohort; and 2783/121 890 (2.3%) versus 3939/168 214 (2.3%), adjusted risk ratio 0.82 (0.78 to 0.86) in the non-shockable cohort (supplementary table A).

**Sequentially matched cohort**

After sequential matching, 16 114 patients in the shockable cohort and 236 042 in the non-shockable cohort were matched (table 2). All of the variables incorporated into the propensity score model were well balanced in both cohorts except for defibrillation by EMS personnel before matching in the shockable cohort (standardised difference 0.19). In both the shockable and non-shockable cohorts, the largest numbers of patients were matched during five to less than 10 minutes from the time EMS personnel started cardiopulmonary resuscitation: 7040 (43.7%) in the shockable cohort and 98 410 (41.7%) in the non-shockable cohort. Among patients who had not yet received AAM (that is, patients in the “no AAM” group), 2605/8057 (32.3%) in the shockable cohort and 35 920/118 021 (30.4%) in the non-shockable cohort received AAM later.

**Outcomes in sequentially matched cohort of shockable rhythm**

In the shockable cohort, survival did not differ between patients who received AAM and those who did not receive AAM (1546/8057 (19.2%) versus 1500/8057 (18.6%); adjusted risk ratio 1.00, 0.93 to 1.07) (table 3). Patients who received AAM had lower risks of favourable functional survival (776/8057 (9.6%) versus 865/8057 (10.7%); adjusted risk ratio 0.87, 0.79 to 0.96). When we stratified the matched patients by the type of AAM (supplementary table B), neither supraglottic airway placement nor endotracheal intubation was associated with survival: adjusted risk ratios 1.00 (0.93 to 1.07) for supraglottic airway placement and 0.97 (0.79 to 1.18) for endotracheal intubation (fig 2).

**Outcomes in sequentially matched cohort of non-shockable rhythm**

In the non-shockable cohort, patients with AAM had greater risks of survival (2696/118 021 (2.3%) versus 2127/118 021 (1.8%); adjusted risk ratio 1.27, 1.20 to 1.35) (table 3). Favourable functional survival did not differ between patients who received AAM and those who did not receive AAM (499/118 021 (0.4%) versus 489/118 021 (0.4%); adjusted risk ratio 1.11, 0.97 to 1.26). When we stratified the matched patients by the type of AAM (supplementary table C), both supraglottic airway placement and endotracheal intubation were associated with survival: adjusted risk ratios were 1.11 (1.03 to 1.19) for supraglottic airway placement and 2.00 (1.74 to 2.30) for endotracheal intubation (fig 2).

**Sensitivity analysis**

In the cohorts matched using traditional, non-time dependent propensity score matching, AAM was associated with worse outcomes in both rhythm cohorts (supplementary table A). The adjusted risk ratios were similar to the results of multivariable regression analysis in the original cohorts.

**Discussion**
In this large cohort study with time dependent propensity score sequential matching of adults with out-of-hospital cardiac arrest, the association of AAM with favourable outcomes depended on the first documented electrocardiographic rhythm (shockable or non-shockable). AAM did not show any advantage for shockable rhythm, whereas for non-shockable rhythm AAM was associated with higher one month survival.

**Relation to previous studies**

In previous observational studies, pre-hospital AAM by EMS personnel was associated with worse outcomes. A previous study using traditional propensity score matching in the Japanese nationwide registry reported that AAM was associated with worse outcomes than bag-valve-mask ventilation. However, some of these previous observational studies may have had biased results favouring no AAM because patients who received AAM were likely to have needed AAM because they were more severely ill before arrest, a status that is potentially associated with less favourable outcomes. In addition, patients who received AAM during cardiopulmonary resuscitation would never have achieved return of spontaneous circulation before receiving AAM, a phenomenon now known as “resuscitation time bias” that would favour better outcomes in the no AAM group. Other recent cardiac arrest studies have used sequential matching with time dependent propensity scores to mitigate resuscitation time bias.

**Strengths and weaknesses of study**

Our study has several strengths. Firstly, its population based design using the nationwide registry enabled us to do the final analysis with complete data and no missing information. Secondly, the large sample size allowed us to estimate the associations between AAM and one month survival even when we divided our cohort by the first documented electrocardiographic rhythm. Thirdly, to overcome resuscitation time bias, we used time dependent propensity score sequential matching analysis. With this method, the groups were closely matched with respect to the timing of AAM. If we had not considered the timing of AAM, we would have had different results, as we showed in the multivariable regression analysis and in the sensitivity analysis using traditional propensity score matching (see supplementary material). We believe that this difference between the results is evidence of the presence of resuscitation time bias. Our approach is believed to have a lower risk of bias in its results compared with previous studies.

Our study also has some limitations, however. Firstly, we did not have data about whether the first documented electrocardiographic rhythm changed to another rhythm during cardiopulmonary resuscitation. Patients with non-shockable rhythm are thought to be likely to receive AAM more frequently and earlier than those with shockable rhythm, according to the algorithms of the international guidelines. Nevertheless, patients in both of our cohorts had similar proportions and timing of AAM. This might indicate that many patients whose first documented electrocardiographic rhythm is shockable develop a non-shockable rhythm during cardiopulmonary resuscitation and subsequently receive AAM according to the non-shockable algorithm. As post-defibrillation changes to non-shockable rhythms are common, and such patients have poorer outcomes than patients without rhythm changes, this would bias the results towards favouring no AAM among patients with a shockable first documented electrocardiographic rhythm.
Secondly, the number of AAM attempts per patient, especially for endotracheal intubation, and information on unsuccessful intubation were not available in our registry. Failed intubation is known to be associated with poor outcomes.\textsuperscript{32} If AAM was delayed, therefore, the patient could be matched in the no AAM group, and this could bias the results towards favouring the AAM group. The lack of information about proportion of successful intubation also decreases the generalisability of our study outside Japan. Thirdly, our database did not contain information on whether each patient was treated by a specially trained emergency life saving technician. As endotracheal intubation is performed only by these technicians and they are generally more experienced and trained than ordinary technicians, this could bias the results towards favouring AAM. Fourthly, we could not address potential confounding by indication in this study. For example, our database did not have any information about patients’ comorbidities that could affect the decisions of EMS personnel. If patients with more significant comorbidities were less likely to receive interventions including AAM, this could bias the results towards favouring AAM. In contrast, if patients were more severely ill before arrest and more likely to have received AAM, this confounding could favour better outcomes in the no AAM group.

**Meaning of study: possible explanations and implications for clinicians and policy makers**

In this study, we separated our original cohort by the first documented electrocardiographic rhythm because the international guidelines suggest different approaches in the algorithms for shockable and non-shockable rhythms.\textsuperscript{20,21} Although the association in our sequential matching analysis was much weaker than the associations in the original cohort analysis and the traditional propensity score matching analysis, AAM was not associated with better one month survival and was still associated with a lower chance of favourable functional survival among patients with shockable rhythm. Among patients with non-shockable rhythm, in contrast, AAM was associated with better survival. Similarly, in a recent observational study of in-hospital cardiac arrest, a significant interaction affecting the primary outcome, survival to hospital discharge, was seen between patients with initial shockable and non-shockable rhythms.\textsuperscript{14} On the basis of our results, different airway management strategies for shockable and non-shockable rhythms should be more strongly emphasised; immediate defibrillation and continuous chest compression rather than ventilatory support are essential for patients with shockable rhythm, as recommended in the guidelines,\textsuperscript{20,21} whereas those with non-shockable rhythm could benefit from delivery of oxygen with protected airway ventilation.

**Unanswered questions and future research**

A recently published randomised controlled trial, in which all pre-hospital endotracheal intubation was done by experienced emergency physicians with a very low intubation failure proportion of 2%, failed to show non-inferiority of bag-valve-mask ventilation relative to endotracheal intubation, but bag-valve-mask ventilation was associated with complications such as regurgitation.\textsuperscript{2} Two recent randomised controlled trials compared pre-hospital supraglottic airway placement and endotracheal intubation during cardiopulmonary resuscitation.\textsuperscript{33,34} One showed that supraglottic airway placement was not significantly superior to endotracheal intubation,\textsuperscript{33} whereas the other reported that supraglottic airway placement was more effective in terms of patient outcomes than endotracheal intubation.\textsuperscript{34} In the second trial, however, the proportion of successful initial endotracheal intubation was low at 51%.\textsuperscript{34,35} These results imply that the findings of future trials conducted under circumstances that allow for higher proportions of successful
intubation might be different. In addition, no studies have investigated the effectiveness of supraglottic airway placement compared with bag-valve-mask ventilation. Given the findings of these recent trials, future research needs to compare supraglottic airway placement and bag-valve-mask ventilation.

**Conclusions**

In this study of 310,620 cases of out-of-hospital cardiac arrest in adults in a Japanese nationwide, population based out-of-hospital cardiac arrest registry, results differed according to the patient’s first documented electrocardiographic rhythm. After time dependent propensity score sequential matching, AAM was not associated with survival in the shockable cohort but was associated with better survival in the non-shockable cohort. When we did not account for the timing of AAM and used multivariable regression or traditional propensity score matching, the associations suggested an unfavourable effect of AAM and were similar to the results of previous observational studies. Our findings suggest that different airway management strategies should be emphasised on the basis of initial electrocardiographic rhythm. Our results also imply that we should account for the timing of interventions in observational research.

**What is already known on this topic**

- Many previous observational studies reported unfavourable associations between pre-hospital advanced airway management and patients’ outcomes after out-of-hospital cardiac arrest
- A recent randomised controlled non-inferiority trial failed to show non-inferiority of bag-valve-mask ventilation versus endotracheal intubation, with more complications in the former group

**What this study adds**

- In this study of 310,620 adults with out-of-hospital cardiac arrest, results differed according to each patient’s first documented electrocardiographic rhythm
- Advanced airway management was not associated with survival in the shockable cohort but was associated with better survival in the non-shockable cohort, although the absolute effect was very small
- These findings suggest that different airway management strategies should be emphasised on the basis of the initial electrocardiographic rhythm: shockable or non-shockable

**Acknowledgments**

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**Web extra.**

Extra material supplied by authors
Web appendix: Supplementary material

Notes

Contributors: JI designed the study, analysed the data, and wrote the first draft of the manuscript. SK, KG, and TKitamura reviewed all statistical analyses and critically revised the manuscript. MO, KK, CN, TKiguchi, TM, TKawamura, TI, and CWC supervised the analysis of the data and critically revised the manuscript. All the authors read and approved the final manuscript. The corresponding author attests that all listed authors meet authorship criteria and that no others meeting the criteria have been omitted. JI is the guarantor.

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Competing interests: All authors have completed the ICMJE uniform disclosure form at www.icmje.org/coi_disclosure.pdf (available on request from the corresponding author) and declare: no support from any organisation for the submitted work other than that described above; no financial relationships with any organisations that might have an interest in the submitted work in the previous three years; and no other relationships or activities that could appear to have influenced the submitted work.

Ethical approval: The medical institutional review board of Osaka University Graduate School of Medicine approved this study and waived the need for informed consent because only de-identified data were used.

Data sharing: No additional data available.

Transparency: The lead author (JI) affirms that the manuscript is an honest, accurate, and transparent account of the study being reported; that no important aspects of the study have been omitted and that any discrepancies from the study as planned (and, if relevant, registered) have been explained.

References


Figures and Tables
Table 1
Characteristics of adult patients with out-of-hospital cardiac arrest with and without advanced airway management during resuscitation in original cohort. Values are numbers (percentages) unless stated otherwise.

<table>
<thead>
<tr>
<th>Characteristics</th>
<th>Shockable rhythm</th>
<th>Non-shockable rhythm</th>
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<td></td>
<td>No AAM (n=12057)</td>
<td>AAM (n=8459)</td>
<td>Standardised difference</td>
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<td>2010 (23.8)</td>
<td></td>
</tr>
<tr>
<td>Summer</td>
<td>2713 (22.5)</td>
<td>1903 (22.5)</td>
<td>0.03</td>
</tr>
<tr>
<td>Autumn</td>
<td>2915 (24.2)</td>
<td>1952 (23.1)</td>
<td></td>
</tr>
<tr>
<td>Winter</td>
<td>3530 (29.3)</td>
<td>2594 (30.7)</td>
<td></td>
</tr>
<tr>
<td>Day:</td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

AAM = advanced airway management; CPR = cardiopulmonary resuscitation; EMS = emergency medical services; EMS CPR = cardiopulmonary resuscitation by emergency medical services personnel; IQR = interquartile range.
Flow of out-of-hospital cardiac arrest cases from All-Japan Utstein Project Database between January 2014 and December 2016. AAM=advanced airway management; CPR=cardiopulmonary resuscitation; EMS=emergency medical services; EMS CPR=cardiopulmonary resuscitation by emergency medical services personnel
## Table 2

Characteristics of adult patients with out-of-hospital cardiac arrest with and without advanced airway management during resuscitation in time dependent propensity score sequentially matched cohort. Values are numbers (percentages) unless stated otherwise.

<table>
<thead>
<tr>
<th>Characteristics</th>
<th>Shockable rhythm</th>
<th>Non-shockable rhythm</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>No AAM (n=8057)</td>
<td>AAM (n=8057)</td>
</tr>
<tr>
<td>Year:</td>
<td></td>
<td></td>
</tr>
<tr>
<td>2014</td>
<td>2731 (33.9)</td>
<td>2776 (34.5)</td>
</tr>
<tr>
<td>2015</td>
<td>2627 (32.6)</td>
<td>2604 (32.3)</td>
</tr>
<tr>
<td>2016</td>
<td>2699 (33.5)</td>
<td>2677 (33.2)</td>
</tr>
<tr>
<td>Season:</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Spring</td>
<td>1886 (23.4)</td>
<td>1914 (23.8)</td>
</tr>
<tr>
<td>Summer</td>
<td>1678 (20.8)</td>
<td>1812 (22.5)</td>
</tr>
<tr>
<td>Autumn</td>
<td>1867 (23.2)</td>
<td>1850 (23.0)</td>
</tr>
<tr>
<td>Winter</td>
<td>2626 (32.6)</td>
<td>2481 (30.8)</td>
</tr>
</tbody>
</table>

AAM=advanced airway management; CPR=cardiopulmonary resuscitation; EMS=emergency medical services; EMS CPR=cardiopulmonary resuscitation by emergency medical services personnel; IQR=interquartile range.
Table 3
Outcomes in time dependent propensity score sequentially matched cohort

<table>
<thead>
<tr>
<th>Outcomes*</th>
<th>No (%) patients with outcome/total patients</th>
<th>Unadjusted risk ratio (95% CI)</th>
<th>Adjusted risk ratio† (95% CI)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>No AAM</td>
<td>AAM</td>
<td></td>
</tr>
<tr>
<td><strong>Shockable rhythm</strong></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Survival</td>
<td>1500/8057 (18.6)</td>
<td>1546/8057 (19.2)</td>
<td>1.09 (1.02 to 1.17)</td>
</tr>
<tr>
<td>Favourable functional survival</td>
<td>865/8057 (10.7)</td>
<td>776/8057 (9.6)</td>
<td>0.97 (0.88 to 1.07)</td>
</tr>
<tr>
<td><strong>Non-shockable rhythm</strong></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Survival</td>
<td>2127/118 021 (1.8)</td>
<td>2696/118 021 (2.3)</td>
<td>1.26 (1.19 to 1.34)</td>
</tr>
<tr>
<td>Favourable functional survival</td>
<td>489/118 021 (0.4)</td>
<td>499/118 021 (0.4)</td>
<td>1.10 (0.96 to 1.26)</td>
</tr>
</tbody>
</table>

AAM=advanced airway management.

*At one month or at hospital discharge within one month.

†Each risk ratio was adjusted for prefecture preference categories for performing supraglottic airway placement and endotracheal intubation, year, season, day, time, age, sex, cause of cardiac arrest, witness category, basic life support by bystander with or without dispatcher instruction, use of public access defibrillator, pre-hospital involvement of physician, time from call to cardiopulmonary resuscitation by emergency medical services personnel, defibrillation before matching, and administration of adrenaline before matching.
### Fig 2

<table>
<thead>
<tr>
<th>Study</th>
<th>No of patients with outcome/total patients (%)</th>
<th>Adjusted risk ratio† (95% CI)</th>
<th>Adjusted risk ratio† (95% CI)</th>
</tr>
</thead>
<tbody>
<tr>
<td>No AAM*</td>
<td>AAM</td>
<td></td>
<td></td>
</tr>
<tr>
<td><strong>Shockable rhythm</strong></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td><strong>Survival</strong></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Supraglottic airway placement</td>
<td>1295/6921 (18.7)</td>
<td>1365/6921 (19.7)</td>
<td>1.00 (0.93 to 1.07)</td>
</tr>
<tr>
<td>Endotracheal intubation</td>
<td>205/1136 (18.0)</td>
<td>181/1136 (15.9)</td>
<td>0.97 (0.79 to 1.18)</td>
</tr>
<tr>
<td><strong>Favourable functional survival</strong></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Supraglottic airway placement</td>
<td>749/6921 (10.8)</td>
<td>699/6921 (10.1)</td>
<td>0.89 (0.81 to 0.99)</td>
</tr>
<tr>
<td>Endotracheal intubation</td>
<td>116/1136 (10.2)</td>
<td>77/1136 (6.8)</td>
<td>0.75 (0.56 to 1.00)</td>
</tr>
<tr>
<td><strong>Non-shockable rhythm</strong></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td><strong>Survival</strong></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Supraglottic airway placement</td>
<td>1753/94 470 (1.9)</td>
<td>1918/94 470 (2.0)</td>
<td>1.11 (1.03 to 1.19)</td>
</tr>
<tr>
<td>Endotracheal intubation</td>
<td>374/23 551 (1.6)</td>
<td>778/23 551 (3.3)</td>
<td>2.00 (1.74 to 2.30)</td>
</tr>
<tr>
<td><strong>Favourable functional survival</strong></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Supraglottic airway placement</td>
<td>403/94 470 (0.4)</td>
<td>379/94 470 (0.4)</td>
<td>1.04 (0.90 to 1.21)</td>
</tr>
<tr>
<td>Endotracheal intubation</td>
<td>86/23 551 (0.4)</td>
<td>120/23 551 (0.5)</td>
<td>1.46 (1.09 to 1.96)</td>
</tr>
</tbody>
</table>

Analysis stratified by type of advanced airway management (AAM) in matched cohort. *Matched control at same timing of supraglottic airway placement or endotracheal intubation. †Adjusted for prefecture preference categories for performing supraglottic airway placement and endotracheal intubation, year, season, day, time, age, sex, cause of cardiac arrest, witness category, basic life support by bystander with or without dispatcher instruction, use of public access defibrillator, pre-hospital involvement of physician, time from call to cardiopulmonary resuscitation by emergency medical services personnel, defibrillation before matching, and administration of adrenaline before matching.