The Influence of Scenario-Based Training and Real-Time Audiovisual Feedback on Out-of-Hospital Cardiopulmonary Resuscitation Quality and Survival From Out-of-Hospital Cardiac Arrest

Bentley J. Bobrow, MD; Tyler F. Vadeboncoeur, MD; Uwe Stolz, PhD, MPH; Annemarie E. Silver, PhD; John M. Tobin, CEP; Scott A. Crawford, EMT-B; Terence K. Mason, RN; Jerome Schirmer, CEP; Gary A. Smith, MD; Daniel W. Spaite, MD

**Study objective:** We assess whether an initiative to optimize out-of-hospital provider cardiopulmonary resuscitation (CPR) quality is associated with improved CPR quality and increased survival from out-of-hospital cardiac arrest.

**Methods:** This was a before-after study of consecutive adult out-of-hospital cardiac arrest. Data were obtained from out-of-hospital forms and defibrillators. Phase 1 included 18 months with real-time audiovisual feedback disabled (October 2008 to March 2010). Phase 2 included 16 months (May 2010 to September 2011) after scenario-based training of 373 professional rescuers and real-time audiovisual feedback enabled. The effect of interventions on survival to hospital discharge was assessed with multivariable logistic regression. Multiple imputation of missing data was used to analyze the effect of interventions on CPR quality.

**Results:** Analysis included 484 out-of-hospital cardiac arrest patients (phase 1: 232; phase 2: 252). Median age was 68 years (interquartile range 56-79); 66.5% were men. CPR quality measures improved significantly from phase 1 to phase 2: Mean chest compression rate decreased from 128 to 106 chest compressions per minute (difference -23 chest compressions; 95% confidence interval [CI] -26 to -19 chest compressions); mean chest compression depth increased from 1.78 to 2.15 inches (difference 0.38 inches; 95% CI 0.28 to 0.47 inches); median chest compression fraction increased from 66.2% to 83.7% (difference 17.6%; 95% CI 15.0% to 20.1%); median preshock pause decreased from 26.9 to 15.5 seconds (difference -11.4 seconds; 95% CI -15.7 to -7.2 seconds), and median ventilation rate decreased from 11.7 to 9.5/minute (difference -2.2/minute; 95% CI -3.9 to -0.5/minute).

**Conclusion:** Implementation of resuscitation training combined with real-time audiovisual feedback was independently associated with improved CPR quality, an increase in survival, and favorable functional outcomes after out-of-hospital cardiac arrest. [Ann Emerg Med. 2013;62:47-56.]

Please see page 48 for the Editor's Capsule Summary of this article.
Editor's Capsule Summary

What is already known on this topic
Despite decades of cardiac arrest research, functional survival after out-of-hospital cardiac arrest has not improved substantially.

What question this study addressed
Whether a “bundle” of a cardiopulmonary resuscitation training program emphasizing performance metrics and the use of real-time audiovisual feedback improves survival for out-of-hospital cardiac arrest victims.

What this study adds to our knowledge
In this before-after trial of 484 patients, unadjusted survival to discharge with favorable functional outcomes was 6.5% in the before group and 10.8% (difference 4.2%; 95% confidence interval [CI] –0.8% to 9.2%) after implementation of the bundle, and the adjusted odds ratio was 2.69 (95% CI 1.04 to 6.94).

How this is relevant to clinical practice
Within the limitations of a study design that is vulnerable to temporal confounding, this study suggests that this approach might be beneficial.

Outcomes by systematically improving the CPR quality delivered by out-of-hospital providers.

In addition to novel approaches to CPR training, real-time audiovisual feedback has been shown to improve CPR quality in actual arrest scenarios both inside and outside the hospital. Hostler et al23 showed improvement in CPR quality metrics but not outcomes when real-time audiovisual feedback was used in the out-of-hospital setting. Edelson et al24 demonstrated that real-time audiovisual feedback used for inhospital arrests improved CPR quality and increased rates of return of spontaneous circulation. For in-hospital training, Wayne et al25,26 and Wayne and McGaghie27 showed significant improvement in CPR performance with simulations and a team approach.

Goals of This Investigation
Our a priori hypothesis was that an out-of-hospital initiative aimed at improving CPR quality by implementing (1) scenario-based CPR training, emphasizing a team approach to resuscitation and the importance of CPR quality metrics, and (2) real-time audiovisual feedback during CPR would improve CPR quality and survival from out-of-hospital cardiac arrest.

MATERIALS AND METHODS
Setting
Data were collected from a single fire-based emergency medical services (EMS) agency located in Mesa, AZ, which responds to a suburban population of 439,000 residents, with approximately 70,000 911 calls annually.28 The agency includes 19 fire stations staffed by 202 emergency medical technician (EMT)-paramedics and 171 EMT-basics. A typical responding crew includes 2 EMT-paramedics and 2 EMT-basics. Additionally, a privately contracted ambulance company assists the fire-based rescuers with patient transport to hospitals. The Mesa Fire/Medical Department participates in the statewide cardiac resuscitation public health initiative called “SHARE—Save Hearts in Arizona Registry and Education.”29 This department has used an innovative minimally interrupted cardiac resuscitation protocol as their standard approach to adult out-of-hospital cardiac arrest from suspected cardiac cause since 2006. Minimally interrupted cardiac resuscitation has been previously described.30

Out-of-hospital cardiac arrest has been designated a major public health problem by the Arizona Department of Health Services. SHARE is the designated public health program created to measure response to out-of-hospital cardiac arrest and improve outcomes. Thus, the SHARE Program initiatives and its data collection are exempt from the Health Insurance Portability and Accountability Act. By virtue of SHARE being a health department–sponsored public health initiative, the Arizona Department of Health Services' Human Subjects Review Board and the University of Arizona institutional review board have determined that neither the interventions nor their evaluation constitutes human subjects research and have approved the publication of deidentified data.

Study Design and Selection of Participants
This is a prospective, before-after, observational cohort study of consecutive adult patients (aged ≥18 years) with out-of-hospital cardiac arrest of presumed cardiac cause who had out-of-hospital initiation of CPR. Cases were excluded from analysis if resuscitation was not initiated, the patient had a do-not-resuscitate order, arrest was witnessed by EMS, or the cause of the arrest was presumed to be noncardiac.

Interventions
Eighteen months' worth of baseline CPR quality and outcome data (October 7, 2008, to March 31, 2010) were collected during phase 1 (before). Real-time audiovisual feedback was not enabled during phase 1. The subsequent intervention included 2 hours of didactic teaching, along with 2 hours of team-centered psychomotor practice using scenario-based training, and activation of real-time audiovisual feedback. Didactic education and scenario-based training repeatedly and explicitly emphasized a team approach to resuscitation and meticulous compliance with the parameters of high-quality CPR within their minimally interrupted cardiac resuscitation protocol. Providers were educated about specific positioning and the role of each team member in a “pit crew” model of resuscitation (Appendix E1, available online at http://www.annemergmed.com), with the intent that this model would be used during actual resuscitations. The prime importance of uninterrupted, high-quality chest compressions was stressed and
the "compressor" was trained to have an unobstructed view of the defibrillator to enhance the effectiveness of real-time audiovisual feedback. In addition to the initial training, 10-minute videos were shown for both 2- and 4-provider crews to reinforce the patterned approach to resuscitation (see http://azdhs.gov/azshare/ccr_share.htm). Providers were specifically trained to avoid excessive ventilation (both rate and volume) and were educated to use the CPR interval timer (on the defibrillator) to space ventilations properly (ie, deliver 1 ventilation every 6 seconds). The training emphasized the importance of applying the combination defibrillator pads/accelerometer without interrupting compressions.

The monitor-defibrillator used in this study provides real-time audiovisual feedback through both audio and visual prompts. The visual display allows the compressor to see multiple, real-time, compression-to-compression quality parameters, including absolute compression depth, absolute compression rate, and a measure that includes a weighted summary analysis of depth, rate, and compression fraction (Appendix E1, available online at http://www.annemergmed.com). When compressions are discontinued for at least 3 seconds, an idle timer is prominently displayed, reminding the compressor to resume CPR. Rate and depth measurements are displayed numerically on the monitor. If compressions are performed outside of the target depth or rate (ie, depth <2 inches, rate <90 or >120 compressions/minute), the parameter label (rate or depth) and its numeric value are illuminated with a distinct red highlight that serves as a visual "alarm." The text "Fully Release" is automatically displayed every 30 seconds. An audio metronome, set to 100 compressions per minute, sounds any time compressions are performed. All other audio prompts related to CPR quality (eg, "push harder," "good compressions") remained disabled in both phases. The "charge during CPR" feature was enabled during phase 2 (after) to automatically charge the defibrillator before the end of each 2-minute chest compression interval, with the goal of minimizing compression interruptions while waiting for the defibrillator to charge.

On April 6, 2010, a 4-hour training session was conducted with 9 "master trainers," who later trained the remaining 364 providers between April 7, 2010, and April 29, 2010. Phase 2 began on May 27, 2010, after training was completed and the real-time audiovisual feedback and new software were enabled on the monitor-defibrillators.

Methods of Measurement

chest compression quality was measured during resuscitation with a monitor-defibrillator (E-series; ZOLL Medical, Chelmsford, MA) with Food and Drug Administration–approved accelerometer-based technology that measures chest compression fraction, depth, rate, and rate of recoil. The accelerometer is integrated into defibrillator pads that are used for patient monitoring and defibrillation. The defibrillator units are equipped with Food and Drug Administration–approved technologies that provide real-time audiovisual feedback on the quality of compressions.

chest compression fraction was measured as the percentage of time compressions were performed (when indicated) throughout the entire resuscitation event. Compressions were considered indicated any time a patient was without spontaneous pulses (as documented in the patient care report and confirmed by ECG) during out-of-hospital care (ie, excluded data after arrival at the emergency department (ED)) and when compression data were valid (ie, the pads were connected and adhered properly). Time was not allotted for the performance of interventions such as ventilation, defibrillation, or intubation (ie, the timer continued to run during all interventions). Fresh shock was calculated as the number of seconds without ongoing compressions before shock delivery for patients with a shockable rhythm (ventricular fibrillation/tachycardia). "Ongoing" compressions were defined as at least 5 back-to-back compressions. Recoil was measured as the peak chest compression release velocity (milli-inches/second) during each compression. Ventilation rates were averaged for each minute of postintubation EMS care without return of spontaneous circulation. Ventilations were captured with the end-tidal CO₂ waveform from a sidestream ETCO₂ adaptor (LoFlo Sidestream CO₂ Module; Philips/Respironics; Wallingford, CT), which was placed after intubation. ETCO₂ values were averaged for each case from all out-of-hospital minutes containing valid ETCO₂ data without return of spontaneous circulation. Minutes with ETCO₂ values greater than 40 mm Hg were no: averaged because they may have been associated with return of spontaneous circulation.

The SHARE program has been previously described in detail and includes a voluntary Utstein-style out-of-hospital cardiac arrest EMS database linked with inhospital postarrest process and outcome data from hospitals. Data collected from participating EMS systems and hospitals are entered into an ACCESS 2007 (Microsoft Corporation, Redmond WA) database maintained on a secure server at the University of Arizona. The SHARE database is mapped to the Cardiac Arrest Registry to Enhance Survival Registry, the largest national out-of-hospital cardiac arrest reporting system (http://www.mycares.net). The SHARE database has multiple logic constraints for out-of-hospital cardiac arrest data elements. For example, arrival in the ED cannot occur before collapse. When values outside the realm of physical possibility are encountered or are missing for any of the Utstein data elements, secondary and tertiary data sources (such as private ambulance transporting first care reports or hospital ED records) are referenced, which allows the backfilling of missing data elements or confirmation of suspected erroneous elements. Additionally, each record goes through a manual review before being committed to the data set. Minimally interrupted cardiac resuscitation protocol compliance was determined by all 4 components: 200 preshock chest compressions, 200 postshock chest compressions, delayed intubation attempt for 3 cycles of 200 compressions and rhythm analysis, and patients having received intravenous epinephrine in the first or second cycle of chest compressions.
Outcome Measures

The main outcome variables were survival to hospital discharge, favorable functional outcome (Cerebral Performance Category score of 1 or 2) as measured at hospital discharge by formally trained hospital personnel, and CPR quality. These outcomes were compared between study phases (phase 1 versus phase 2), the main independent variable. Additional confounders and risk factors considered were initial cardiac rhythm on EMS arrival (initial rhythm), EMS dispatch-to-on-scene arrival interval (response interval), age, sex, location of arrest, witnessed versus unwitnessed arrest, provision of bystander CPR, the use of therapeutic hypothermia, and minimally interrupted cardiac resuscitation protocol compliance.

CPR quality measures included the following: chest compression fraction, depth, rate, and release velocity; preshock and postshock pause; and ventilation rate.

Primary Data Analysis

For univariate analyses, Fisher’s exact test (proportions), t test (means), or Kruskal-Wallis test (medians) was used, with \( \alpha = .05 \). Summary statistics are reported as percentages, means with 95% confidence intervals (CIs), and medians with interquartile ranges. Absolute differences with 95% CIs are reported for comparisons of means, medians, and proportions. To calculate crude and adjusted odds ratios (ORs) for survival and favorable functional outcome, we used multivariable mixed-effects logistic regression (xtmelogit, Stata version 12.1; StataCorp, College Station, TX), with the hospital providing final care as the random effect and all patients not transported treated as a single cluster.

Covariates were included in the final model if the associated P value from the likelihood ratio \( \chi^2 \) test was less than or equal to .05 or if they were judged to be a significant confounder (inclusion of covariate changed the coefficient for main risk factor >10%) of the relationship between the outcome variable and our main independent variable, pre- versus postperiod. We calculated the Hosmer-Lemeshow goodness-of-fit statistic and the area under the receiver operating characteristics curve for each final multivariable model, using the predicted probabilities incorporating the random effects from the mixed-effects modeling. We also explored model diagnostics for the mixed-effects model by examining final model residuals (Pearson, deviance, and Anscombe) to identify overly influential covariate patterns or outliers that could represent miscoded cases. We also examined model diagnostics (leverage, deviance, etc) for all final models, assuming no random effects (ordinary logistic regression), as an additional approach to identifying potential outliers. Fractional polynomial regression was used to examine the linear relationship of continuous variables with the outcome variables in the logit scale.

Univariate multiple imputation methods and approaches were explored to handle missing values for the chest compression quality metrics (mean depth, mean rate, recoil, compression fraction, percentage of compression \( \approx 2 \) inches, mean preshock and postshock pause), using the following variables as covariates for imputation: survival to discharge, pre/postperiod, out-of-hospital return of spontaneous circulation, age, sex, witnessed arrest, shockable rhythm (ventricular fibrillation/ventricular tachycardia), bystander CPR, location of arrest, and EMS response interval. The pattern of missing data was first explored with univariate analyses to examine associations between a patient’s having missing data and study covariates. Because all CPR quality data were missing for a case with any missing CPR quality data, each CPR quality metric was imputed independently. Twenty imputed data sets were created for analysis with linear regression ("mi impute regress" with random number seed “4987”), and all chest compression quality metrics were compared between phase 1 and phase 2 with either linear or median regression, as appropriate, with associated 95% CIs for differences. Ventilation and ETCO\(_2\) data were not amenable to imputation because we could not identify the time of intubation for all patients and thus were not able to determine which patients were intubated before sustaining return of spontaneous circulation. Thus, ventilation data are compared between the phase 1 and phase 2 with nonimputed data.

We conducted a post hoc analysis to identify potential secular trends or a Hawthorne effect in all-rhythm survival and positive functional outcomes by dividing phase 1 into halves and comparing outcomes (survival and functional outcome) between them. In addition, we investigated the proportion of patients who were not transported to a hospital but were treated by EMS on scene in phase 1 versus phase 2 to assess whether this was associated with any outcome differences between periods. All statistical analyses and imputations were performed with Stata (version 12.1).
RESULTS

Characteristics of Study Subjects

A total of 232 consecutive, adult, non-EMS-witnessed, out-of-hospital cardiac arrests of presumed cardiac cause with resuscitation initiated in the field occurred in phase 1 and 252 in phase 2 (see the Figure for inclusion/exclusion flow chart). Among the 484 patients in this analysis, 1 was missing survival data and 3 were missing functional outcome scores. A total of 147 patients (30.4%) were missing CPR quality data, and 71 of 228 patients (31.1%) who received shock were missing pre/post-shock pausedata. A comparison of demographics and standard Utstein data elements is presented in Table 1. Patient characteristics of phase 1 and phase 2 were similar. Overall, 113 of 484 patients (23.4%) achieved return of spontaneous circulation in the out-of-hospital setting and 55 patients (11.4%) survived to hospital discharge. Of patients with a witnessed arrest and a shockable rhythm, 35 of 93 (37.6%) survived to hospital discharge.

Main Results

Table 2 shows survival and favorable functional outcomes across study periods, along with crude and adjusted ORs. Survival increased from 8.7% (20/231) in phase 1 to 13.9% (35/252) in phase 2 (absolute difference 5.2; 95% CI —0.4 to 10.8), with a crude OR of 1.73 (95% CI 0.93 to 3.21) and an adjusted OR of 2.72 (95% CI 1.15 to 6.41), controlling for witnessed arrest, initial rhythm, provision of therapeutic hypothermia, age, and minimally interrupted cardiac resuscitation protocol compliance. Favorable functional outcome increased from 6.5% in phase 1 to 10.8% in phase 2 (absolute difference 4.2%; 95% CI —0.8% to 9.2%), with a crude OR of 1.76 (95% CI 0.88 to 3.52) and an adjusted OR of 2.69 (95% CI 1.04 to 6.94), adjusting for witnessed arrest, provision of therapeutic hypothermia, age, and minimally interrupted cardiac resuscitation protocol compliance. Age as a continuous variable was linear in the logit scale, as determined by fractional polynomial regression. The intraclass correlation (clustereffect) between both survival and favorable functional outcome and hospital was 0.125 (95% CI 0.018 to 0.528) and 0.140 (95% CI 0.023 to 0.532), respectively, and the likelihood ratio test P value comparing mixed effects versus ordinary logistic regression was .02 and .01, respectively, indicating a significant cluster effect and justifying mixed-effects logistic regression. For the final multivariable model for survival to discharge, the Hosmer-Lemeshow goodness of fit P value was .85 and area under the receiver operating characteristics curve for survival was 0.913. For the final model for positive functional outcome, the Hosmer-Lemeshow goodness of fit P value was .30 and the area under the receiver operating characteristics curve was 0.920. In witnessed arrests with a shockable rhythm, survival and functional outcomes improved significantly from phase 1 to phase 2 (survival 26.3% [15/57] to

<table>
<thead>
<tr>
<th>Characteristic</th>
<th>Overall</th>
<th>Preperiod</th>
<th>Postperiod</th>
<th>Absolute Difference to Post—Pre (95% CI)</th>
</tr>
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<tbody>
<tr>
<td>Total, No. (%)</td>
<td>484 (100)</td>
<td>232 (47.9)</td>
<td>252 (52.1)</td>
<td>NA</td>
</tr>
<tr>
<td>Age, median (IQR), y</td>
<td>68 (56—79)</td>
<td>69 (59—79)</td>
<td>68 (55—79)</td>
<td>—1 (—5 to 3)</td>
</tr>
<tr>
<td>Male sex, No. (%)</td>
<td>322 (66.5)</td>
<td>149 (64.2)</td>
<td>173 (68.7)</td>
<td>4.4 (—4.0 to 12.8)</td>
</tr>
<tr>
<td>Witnessed arrest, No. (%)</td>
<td>192 (39.8)</td>
<td>98 (42.2)</td>
<td>94 (37.3)</td>
<td>4.9 (—13.7 to 3.8)</td>
</tr>
<tr>
<td>Shockable rhythm on EMS arrival, No. (%)</td>
<td>150 (31.0)</td>
<td>79 (34.1)</td>
<td>71 (28.2)</td>
<td>—5.9 (—14.1 to 2.4)</td>
</tr>
<tr>
<td>Provision of bystander CPR, No. (%)</td>
<td>192 (39.7)</td>
<td>102 (44.0)</td>
<td>90 (35.7)</td>
<td>—8.3 (—17.0 to 0.5)</td>
</tr>
<tr>
<td>Location of arrest, No. (%)</td>
<td>353 (72.9)</td>
<td>167 (72.0)</td>
<td>186 (73.8)</td>
<td>1.8 (—6.1 to 9.8)</td>
</tr>
<tr>
<td>Residential</td>
<td>353 (72.9)</td>
<td>167 (72.0)</td>
<td>186 (73.8)</td>
<td>1.8 (—6.1 to 9.8)</td>
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<tr>
<td>Medical facility</td>
<td>68 (14.1)</td>
<td>32 (13.8)</td>
<td>36 (14.3)</td>
<td>0.5 (—5.7 to 6.7)</td>
</tr>
<tr>
<td>Public</td>
<td>63 (13.0)</td>
<td>33 (14.2)</td>
<td>30 (11.9)</td>
<td>—2.3 (—8.3 to 3.7)</td>
</tr>
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<td>EMS response interval, median (IQR), min</td>
<td>5 (4—6)</td>
<td>5 (4—6)</td>
<td>5 (4—6)</td>
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<tr>
<td>Use of TH, No. (%)</td>
<td>52 (10.7)</td>
<td>23 (9.9)</td>
<td>29 (11.5)</td>
<td>1.6 (—3.9 to 7.1)</td>
</tr>
<tr>
<td>MICR protocol compliance (complete vs partial), No. (%)</td>
<td>375 (77.5)</td>
<td>155 (66.8)</td>
<td>220 (87.3)</td>
<td>20.5 (13.2 to 27.8)</td>
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<tr>
<td>Return of spontaneous circulation, No. (%)</td>
<td>113 (23.4)</td>
<td>58 (25.0)</td>
<td>55 (21.8)</td>
<td>—3.2 (—10.7 to 4.4)</td>
</tr>
<tr>
<td>Survival to hospital discharge for all rhythms, No./total (%)*</td>
<td>55/483 (11.4)</td>
<td>20/231 (8.7)</td>
<td>35/252 (13.9)</td>
<td>5.2 (—0.4 to 10.8)</td>
</tr>
<tr>
<td>Survival to hospital discharge for witnessed arrests, shockable rhythms, No./total (%)*</td>
<td>35/93 (37.6)</td>
<td>15/57 (26.3)</td>
<td>20/36 (55.6)</td>
<td>29.2 (9.4 to 49.1)</td>
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<tr>
<td>Favorable functional outcome (CPC score = 1 or 2) for all rhythms, No./total (%)†</td>
<td>42/481 (8.7)</td>
<td>15/230 (6.5)</td>
<td>27/251 (10.8)</td>
<td>4.2 (—0.8 to 9.2)†</td>
</tr>
<tr>
<td>Favorable functional outcome (CPC score = 1 or 2) for witnessed arrests, shockable rhythms, No./total (%)†</td>
<td>27/91 (29.7)</td>
<td>11/56 (19.6)</td>
<td>16/35 (45.7)</td>
<td>26.1 (6.6 to 45.6)†</td>
</tr>
</tbody>
</table>

NA, Not applicable; IQR, interquartile range; TH, therapeutic hypothermia; MICR, minimally interrupted cardiac resuscitation; CPC, cerebral performance category.
*Missing survival outcome.
†Missing functional outcomes.
‡Discrepancy between the absolute difference and the subtraction of pre and post values is due to rounding.

Table 1. Demographics and outcomes by study period.
Table 2. Final logistic regression models for survival and neurologic outcomes (all rhythms and witnessed/shockable).

<table>
<thead>
<tr>
<th>Characteristic</th>
<th>No./Total (%)</th>
<th>Absolute Difference (95% CI)</th>
<th>Crude OR (95% CI)</th>
<th>Adjusted OR (95% CI)*</th>
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<tr>
<td><strong>Study period</strong></td>
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<td></td>
<td></td>
<td></td>
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<tr>
<td>Pre</td>
<td>20/231 (8.7)</td>
<td>5.2 (0.4 to 10.8)</td>
<td>1 [Reference]</td>
<td>1 [Reference]</td>
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<tr>
<td>Post</td>
<td>35/252 (13.9)</td>
<td>1.73 (0.93 to 3.21)</td>
<td>2.71 (1.15 to 6.41)</td>
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<tr>
<td><strong>Witnessed arrest</strong></td>
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<td></td>
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<td></td>
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<tr>
<td>No</td>
<td>13/291 (4.5)</td>
<td>17.4 (1.1 to 23.7)</td>
<td>1 [Reference]</td>
<td>1 [Reference]</td>
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<tr>
<td>Yes</td>
<td>42/192 (21.9)</td>
<td>4.19 (2.12 to 8.30)</td>
<td>4.0 (1.72 to 9.28)</td>
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<td><strong>Initial rhythm</strong></td>
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<tr>
<td>Non-shockable</td>
<td>12/334 (3.6)</td>
<td>25.3 (17.7 to 32.8)</td>
<td>[Reference]</td>
<td>1 [Reference]</td>
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<tr>
<td>Ventricular fibrillation</td>
<td>43/149 (28.9)</td>
<td>7.33 (3.59 to 14.99)</td>
<td>5.88 (2.59 to 13.36)</td>
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<td><strong>Use of TH</strong></td>
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<tr>
<td>No</td>
<td>25/431 (5.8)</td>
<td>61.2 (36.3 to 65.5)</td>
<td>1 [Reference]</td>
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<td>Yes</td>
<td>30/52 (57.7)</td>
<td>14.6 (6.98 to 30.51)</td>
<td>11.47 (5.16 to 27.33)</td>
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<tr>
<td><strong>Age, per year</strong></td>
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<td></td>
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<td></td>
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<tr>
<td>Partial</td>
<td>10/108 (9.3)</td>
<td>2.7 (0.36 to 9.1)</td>
<td>1 [Reference]</td>
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<tr>
<td>Complete</td>
<td>45/375 (12.0)</td>
<td>2.02 (0.95 to 4.28)</td>
<td>1.16 (0.46 to 2.93)</td>
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<td><strong>MICR protocol compliance</strong></td>
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<tr>
<td>Partial</td>
<td>10/108 (9.3)</td>
<td>-0.7 (0.29 to 5.5)</td>
<td>1 [Reference]</td>
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<tr>
<td>Complete</td>
<td>32/375 (8.6)</td>
<td>1.31 (0.60 to 2.86)</td>
<td>0.62 (0.23 to 1.63)</td>
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</tr>
</tbody>
</table>

VF, Ventricular fibrillation; VT, ventricular tachycardia.

*Adjusted for all variables listed in this table in final model (likelihood ratio P value for all variables included in final model ≤ .05 or judged a significant confounder).

The absolute difference in survival was 1.4% (95% CI, 0.2% to 2.6%). Among 52% of patients who survived to hospital discharge, 22% of patients who survived to hospital discharge were discharged with permanent normal sinus rhythm, and 28% of patients who survived to hospital discharge were discharged with permanent atrial fibrillation. The absolute difference in hospital discharge was 1.0% (95% CI, 0.2% to 1.8%). The absolute difference in hospital discharge was 1.0% (95% CI, 0.2% to 1.8%). The absolute difference in hospital discharge was 1.0% (95% CI, 0.2% to 1.8%). The absolute difference in hospital discharge was 1.0% (95% CI, 0.2% to 1.8%). The absolute difference in hospital discharge was 1.0% (95% CI, 0.2% to 1.8%). The absolute difference in hospital discharge was 1.0% (95% CI, 0.2% to 1.8%). The absolute difference in hospital discharge was 1.0% (95% CI, 0.2% to 1.8%). The absolute difference in hospital discharge was 1.0% (95% CI, 0.2% to 1.8%). The absolute difference in hospital discharge was 1.0% (95% CI, 0.2% to 1.8%). The absolute difference in hospital discharge was 1.0% (95% CI, 0.2% to 1.8%). The absolute difference in hospital discharge was 1.0% (95% CI, 0.2% to 1.8%). The absolute difference in hospital discharge was 1.0% (95% CI, 0.2% to 1.8%). The absolute difference in hospital discharge was 1.0% (95% CI, 0.2% to 1.8%). The absolute difference in hospital discharge was 1.0% (95% CI, 0.2% to 1.8%). The absolute difference in hospital discharge was 1.0% (95% CI, 0.2% to 1.8%). The absolute difference in hospital discharge was 1.0% (95% CI, 0.2% to 1.8%). The absolute difference in hospital discharge was 1.0% (95% CI, 0.2% to 1.8%). The absolute difference in hospital discharge was 1.0% (95% CI, 0.2% to 1.8%). The absolute difference in hospital discharge was 1.0% (95% CI, 0.2% to 1.8%). The absolute difference in hospital discharge was 1.0% (95% CI, 0.2% to 1.8%). The absolute difference in hospital discharge was 1.0% (95% CI, 0.2% to 1.8%). The absolute difference in hospital discharge was 1.0% (95% CI, 0.2% to 1.8%). The absolute difference in hospital discharge was 1.0% (95% CI, 0.2% to 1.8%). The absolute difference in hospital discharge was 1.0% (95% CI, 0.2% to 1.8%). The absolute difference in hospital discharge was 1.0% (95% CI, 0.2% to 1.8%). The absolute difference in hospital discharge was 1.0% (95% CI, 0.2% to 1.8%). The absolute difference in hospital discharge was 1.0% (95% CI, 0.2% to 1.8%). The absolute difference in hospital discharge was 1.0% (95% CI, 0.2% to 1.8%). The absolute difference in hospital discharge was 1.0% (95% CI, 0.2% to 1.8%). The absolute difference in hospital discharge was 1.0% (95% CI, 0.2% to 1.8%). The absolute difference in hospital discharge was 1.0% (95% CI, 0.2% to 1.8%). The absolute difference in hospital discharge was 1.0% (95% CI, 0.2% to 1.8%). The absolute difference in hospital discharge was 1.0% (95% CI, 0.2% to 1.8%). The absolute difference in hospital discharge was 1.0% (95% CI, 0.2% to 1.8%). The absolute difference in hospital discharge was 1.0% (95% CI, 0.2% to 1.8%). The absolute difference in hospital discharge was 1.0% (95% CI, 0.2% to 1.8%). The absolute difference in hospital discharge was 1.0% (95% CI, 0.2% to 1.8%). The absolute difference in hospital discharge was 1.0% (95% CI, 0.2% to 1.8%). The absolute difference in hospital discharge was 1.0% (95% CI, 0.2% to 1.8%). The absolute difference in hospital discharge was 1.0% (95% CI, 0.2% to 1.8%). The absolute difference in hospital discharge was 1.0% (95% CI, 0.2% to 1.8%).
The thrust of the current literature supports the concept that CPR quality is an important factor in survival from out-of-hospital cardiac arrest. It is strongly emphasized in the 2010 AHA guidelines. This analysis demonstrates that a systematic and comprehensive approach to improving out-of-hospital CPR quality in a large EMS system was associated with achieving the 2010 AHA guideline recommendations for CPR quality, an increase in survival to hospital discharge, and favorable functional outcomes.

These results demonstrate an improvement in CPR quality performance in line with the 2010 AHA guidelines for all
metrics and, most important, increased adjusted odds of survival and favorable functional outcome in our postintervention group (Tables 2 and 3). According to current understanding of the effects of CPR during cardiac arrest and the quality of CPR in most actual resuscitations, our findings are both biologically plausible and logical. For example, one major contributor to the low survival rates in most settings is prolonged inadequate myocardial and cerebral blood flow.6-8 During resuscitation efforts, the forward blood flow generated by CPR is marginal, and as such, any pause in compressions or compressions of inadequate depth have a significant negative effect on both defibrillation success and survival.6-8 Cardiac output is the major determinant of carbon dioxide delivery to the lungs during CPR. In our analysis, the higher ETCO₂ in phase 2 provides strong evidence for improved CPR quality during phase 2 and was likely the result of increased perfusion.

The recognition of the importance of continuous blood flow and the consequences of interrupting myocardial and cerebral perfusion has led to great interest in CPR quality.8-10 Numerous animal and clinical studies have demonstrated that CPR quality (chest compression depth,8-13 fraction,8-13 pre-shock pause,14-16 recoil,17-19 chest compression rate,15,20 and ventilation rate15) has a significant effect on cardiac arrest outcomes. In 2007, Kramer-Johansen et al.30 proposed a rationale for establishing common definitions and a reporting template for CPR metrics. Despite the understanding of the importance of high-quality CPR, most out-of-hospital cardiac arrest victims still do not receive optimal CPR.13,21,33-35

Previous investigators have shown that real-time audiovisual feedback can improve CPR quality.6,7,21-23 Our findings are consistent with these previous investigations and strongly support the current emphasis that the 2010 AHA guidelines place on high-quality CPR as a means to improve survival.32 Although causality cannot be proven according to a nonrandomized trial, to our knowledge, this is the first study to demonstrate an association between a dedicated CPR quality initiative using real-time audiovisual feedback and out-of-hospital cardiac arrest outcomes. Furthermore, it is the first report to show an association between performance of the 2010 AHA CPR quality metrics and increased survival.32 The interventions, which included didactic education, scenario-based training, and real-time audiovisual feedback, were specifically aimed at particular CPR quality metrics. The training emphasized the importance of CPR quality, a team approach to resuscitation, and real-time audiovisual feedback.

Hostler et al.35 recently performed a large (1,586 subjects) cluster-randomized study within the Resuscitation Outcome Consortium to assess the effect of real-time audiovisual feedback on CPR quality and outcomes. Although they found improvements in CPR quality with feedback on compared with feedback off (compression fraction 66% versus 64%, respectively, P=.02; depth 40 mm versus 38 mm, P=.005; rate
103 versus 108, P < .001; percentage with incomplete release 10% versus 15%, P < .001), they did not find improvements in rates of return of spontaneous circulation (44% versus 45%) or survival to hospital discharge (11% versus 12%). And although the study by Hostler et al24 showed statistical improvements in CPR quality, the effect sizes were small and of questionable clinical significance and CPR quality data were missing in 26% of patients. There are several fundamental differences between the study by Hostler et al24 and ours. First, the current investigation included (1) dedicated didactic and scenario-based training emphasizing CPR quality metrics, in addition to real-time audiovisual feedback as part of a bundled approach to improving CPR quality; (2) a major focus on the optimal use of the feedback-capable defibrillator; (3) a specified "pit crew" team approach to resuscitation; and (4) multiple imputation to reduce the likelihood of bias caused by excluding cases with missing CPR process data. Additionally, we used a different feedback device and our training included a targeted and standardized approach to resuscitation, with a focus on maximizing CPR quality.

In this study of out-of-hospital cardiac arrest, a carefully targeted CPR training curriculum in conjunction with real-time audiovisual feedback was independently associated with achievement of the AHA 2010 guideline-recommended metrics for CPR quality and an increased likelihood of both survival to hospital discharge and favorable functional outcome.

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Author affiliations: From the Arizona Department of Health Services Bureau of Emergency Medical Services and Trauma System, Phoenix, AZ (Bobrow); the Maricopa Medical Center/University of Arizona College of Medicine, Phoenix, AZ (Bobrow); Arizona Emergency Medicine Research Center, the University of Arizona College of Medicine (Stoil, Spaite), Tucson, AZ; the Department of Emergency Medicine, Mayo Clinic, Jacksonville, FL (Vadeboncoeur); the ZOLL Medical Corporation, Chelmsford, MA (Silver); and the Mesa Fire/Medical Department, Mesa, AZ (Tobin, Crawford, Mason, Schirmer, Smith).

Author contributions: BJB, TFV, AES, JMT, GAS, and DWS conceived and designed the study. BJB, AES, JMT, SAC, TKM, JS, and GAS supervised the conduct of the study and data collection. BJB, US, and AES managed the data, including quality control. BJB, TFV, US, AES, and DWS provided statistical advice on study design and analyzed the data. TFV drafted the article, and all authors contributed substantially to its revision. TFV takes responsibility for the paper as a whole.

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Address for correspondence: Bentley J. Bobrow, MD, E-mail bobrowb@azdhs.gov.

REFERENCES


Appendix E1.

**Figure E1.** A, E Series Defibrillator Display with Real-Time Audiovisual Feedback Enabled (ZOLL Corporation, Chelmsford MA). B, Coordinated multiprovider resuscitation schematic.

1. Make sure chest compressors can see monitor.
2. Do not interrupt chest compressors—they need to focus on delivery of quality chest compressions.

**4 Person Pit Crew Formation**

- Airway, Drug prep
- Focus on monitor
- Defib, IV/IO Meds
- Chest compressors (switch every 2 min)!!

Second compressor, IV setup and watch monitor