Hands-On Defibrillation
An Analysis of Electrical Current Flow Through Rescuers in Direct Contact With Patients During Biphasic External Defibrillation

Michael S. Lloyd, MD; Brian Heeke, BS; Paul F. Walter, MD; Jonathan J. Langberg, MD

Background—Brief interruptions in chest compressions reduce the efficacy of resuscitation from cardiac arrest. Interruptions of this type are inevitable during hands-off periods for shock delivery to treat ventricular tachyarrhythmias. The safety of a rescuer remaining in contact with a patient being shocked with modern defibrillation equipment has not been investigated.

Methods and Results—This study measured the leakage voltage and current through mock rescuers while they were compressing the chests of 43 patients receiving external biphasic shocks. During the shock, the rescuer’s gloved hand was pressed onto the skin of the patient’s anterior chest. To simulate the worst case of an inadvertent return current pathway, a skin electrode on the rescuers thigh was connected to an electrode on the patient’s shoulder. In no cases were shocks perceptible to the rescuer. Peak potential differences between the rescuer’s wrist and thigh ranged from 0.28 to 14 V (mean 5.8 V). The average leakage current flowing through the rescuer’s body for each phase of the shock waveform was 283±140 μA (range 18.9 to 907 μA). This was below several recommended safety standards for leakage current.

Conclusions—Rescuers performing chest compressions during biphasic external defibrillation are exposed to low levels of leakage current. The present findings support the feasibility of uninterrupted chest compressions during shock delivery, which may enhance the efficacy of defibrillation and cardiocerebral resuscitation. (Circulation. 2008;117:2510-2514.)

Key Words: defibrillation ■ resuscitation ■ cardiopulmonary resuscitation ■ electrical stimulation

Electrocution of rescuers has been recognized as a hazard of external defibrillation therapy.1 Historically, shocks were delivered with paddles that had large, rigid, metallic electrodes.2 Arcing was common owing to inconsistencies in electrode location and contact.3,4 Variability in the distribution and performance of the conductive gel was also an important cause of stray electrical current. Guidelines have thus mandated “hands-off” periods, whereby rescuers have no contact with the patient or stretcher during shock delivery.

There have been substantial changes to external defibrillation technology since the inception of resuscitation protocols with hands-off periods. Biphasic shocks and real-time impedance monitoring have reduced peak voltages. Paddles have been replaced in many cardiac arrest settings by conformal, adhesive, pregelled electrodes, which result in better and more consistent electrode-skin coupling. Enhancements of ECG filtering permit rhythm monitoring during chest compression.

With the current state of the art, if certain types of physical contact with the patient were safe during shock delivery, continuous manual compressions might be possible during cardiac arrest. We addressed the feasibility and safety of direct contact with the patient during defibrillation by measuring the voltage and current through volunteers simulating chest compressions on patients receiving external biphasic countershocks.

Methods
The study protocol was approved by the Institutional Review Board at Emory University. Written informed consent was obtained from all patients. The patient population recruited for the present study...
included those undergoing elective cardioversion for persistent atrial fibrillation or flutter and those undergoing invasive electrophysiology study who were likely to need external cardioversion or defibrillation during the procedure. The external defibrillator used was a Lifepac 12 (Medtronic Corp, Minneapolis, Minn), which delivers a truncated exponential biphasic waveform with impedance compensation. The defibrillator power source was a grounded standard mains outlet. Shocking electrodes were self-adhesive (PadPro, Conmed Corp, Utica, NY) and were placed in the anteroposterior position for all patients. After intravenous sedation and the determination by the physician performing the procedure that it was safe to proceed, 1 of 4 healthcare workers (all coinvestigators of the study) simulated manual chest compressions on the patient. A schematic of the rescuer-patient construct is shown in Figure 1. With the coinvestigator wearing polyethylene medical gloves and standing at the right side of the patient’s bed, ~20 pounds of downward force was applied on the lower half of the patient’s sternum in the area typical for chest compression. The palm of the investigator performing the chest compressions (termed “rescuer” for the remainder of the text) was in direct contact with the bare skin of the patient’s chest adjacent to the anterior shocking electrode. Skin preparation for all electrode attachments included vigorous scrubbing with an alcohol pad and shaving when needed.

To simulate an inadvertent conductive contact between the patient and rescuer, adhesive silver–silver chloride electrodes were placed on the prepared skin of the patient’s posterior left shoulder and the prepared skin of the rescuer’s anterior thigh (electrode surface area ~4 cm²; Conmed Corp). This electrical connection simulated an inadvertent skin-to-skin contact between the rescuer and the patient and allowed for a return pathway for leakage current across a rescuer. This connection established a “rescuer-patient circuit” in which the patient being shocked was the voltage source and the rescuer acted as the load. Measurement of the leakage current and voltage across the rescuer was performed by attaching button electrodes that connected the bare skin of the rescuer’s thigh to the rescuer’s hand and thigh. The voltage drops across the 120-Ω resistor and connected to a digital dual-channel oscilloscope (PicoScope 2202, Picotech, Cambridgeshire, United Kingdom). Voltages were subtracted from the initial 50 μs of the onset, phase change, and termination of the recording. The absolute values of the voltages recorded across the 110-kΩ resistor for each phase of the biphasic waveform were used to derive an average voltage between the rescuer’s hand and thigh. The voltage drops across the 120-Ω series resistor were used to determine current flow through the rescuer during both phases of the shock in an analogous fashion. The range, SD, and mean voltage and current were taken for 36 separate recordings and compared with International Electrotechnical Commission (IEC) 60601-1 and IEC 950 standards for electrical safety.

The authors had full access to and take full responsibility for the integrity of the data. All authors have read and agree to the manuscript as written.

**Results**

Rescuers were instrumented during 43 hands-on shocks in 39 patients. Four shocks were at 100 J, 27 were at 200 J, and 8 were at 360 J. None of the 43 shocks were perceptible to the rescuers. Seven of the 43 shocks did not produce sufficient leakage current to trigger the oscilloscope, which left 36 suitable for analysis. Representative leakage current and voltage recordings are shown in Figure 2, and summary data are in the Table. The voltage between the rescuer’s hand and the return pathway on the thigh was 5.8±5.8 V (range 0.280 to 14.1 V). Leakage current through the rescuer simulating compressions was 283±140 μA (range 18.9 to 907 μA) for each phase of the biphasic waveform. The duration of the defibrillation shocks was ~15 ms, but this number varied slightly owing to impedance compensation by the defibrillator.

With a constant pulse duration of 15 ms for calculations, the mean leakage energy was 24±12 μJ (range 0.07 to 95 μJ). Transthoracic impedances of the patient, as measured by the defibrillator, averaged 60±15 Ω (range 36 to 87 Ω). The mean rescuer circuit impedance, as calculated by dividing the recorded leakage voltage and current, was 22.7 kΩ (range 1.09 to 100 kΩ).

The present data set was compared with several benchmarks of electrical safety (Figure 3). The IEC 950 maximum allowable leakage current for non-handheld equipment is 3500 μA, which is well above the entire range of leakage current values we measured.10 For handheld equipment, the limit is 750 μA. In a single patient, 1 phase of the biphasic waveform exceeded this value (910 μA). The IEC 60601-1 guidelines for medical equipment are more rigorous owing to potential exposure to patients. Under this standard, the cutoff for enclosures is 500 μA under single-fault conditions, which is greater than the overall mean leakage current in the present study; however, 8 measurements of the 72 phases (36 biphasic shocks) exceeded this cutoff.
Discussion

The present study was meant to simulate conditions during resuscitation of cardiac arrest victims by medical personnel. We assumed that these rescuers would use gloves as part of standard precautions and would have access to adhesive defibrillation pads. Rescuers in these conditions were exposed to low levels of leakage current and voltage when performing chest compression during delivery of biphasic shocks. This was the case despite the presence of a low-resistance return pathway between the rescuer’s leg and the patient’s thigh. To the best of our knowledge, these are the first human data to directly address the safety of hands-on defibrillation.

These findings are important because the data demonstrate the safety and feasibility of continuous manual chest compressions throughout the resuscitation cycle. The adverse hemodynamic consequences of delays between chest compression and defibrillation have been clearly demonstrated.9 Most groups have concluded that defibrillation should occur during and not after chest compressions, but concerns about safety have precluded implementation of such a strategy. To circumvent this, fully automated resuscitation systems have been designed that produce continuous compressions, but the practical use of such devices has been limited in most cardiac arrest scenarios.11 We have shown that it is possible to perform continuous manual compressions during shock delivery within the constraints of our model. Elimination of delays in compression before defibrillation would also prevent substantial time lags in restoration of adequate coronary perfusion pressure when chest compressions are resumed after a pause.12 Preventing hands-off periods may improve the quality of chest compressions, which has historically been poor.13 Shock delivery during chest compressions would potentially increase the likelihood of successful defibrillation by reducing transthoracic impedance.14 Finally, uninterrupted compressions would represent a protocol simplification and possibly reduce the time to a first shock.

The present data add to prior work examining leakage current from internally delivered defibrillation shocks in animals and humans. Although leakage currents were higher than those obtained from the present models due to much lower load resistances than our directly measured values, the data likewise showed that the overall leakage energy was small.15,16

No directly comparable safety standard exists for leakage current in this specific situation. Leakage current standards vary widely depending on the type of equipment and exposure scenario. Values for patients are appropriately much smaller than for occupational exposure.17,18 Direct intracardiac exposure to very small leakage currents may cause harm.19 The amount of current needed to cause potential harm is several orders of magnitude higher for external (skin to enclosure) exposure.

We chose to compare our measurements to standards for equipment in the household and in businesses (IEC 950, Underwriters Laboratory 1950) and to those set for medical equipment (IEC 60601-1). These values are obtained by measuring current flow through several paths that incorporate the object of concern, or enclosure. These standards are usually for continuous 60-Hz alternating current and measured with a fixed load simulating that of a human body. In the present study, the patient receiving the shock can be viewed as the voltage source, or enclosure, and the direct thigh-to-shoulder electrical connection between the rescuer and patient can be viewed as a single-fault condition. Although these comparisons are reasonable, the present data

<table>
<thead>
<tr>
<th>Voltage across rescuer, V</th>
<th>5.80</th>
<th>5.77</th>
<th>0.280–14.1</th>
</tr>
</thead>
<tbody>
<tr>
<td>Current through rescuer, μA</td>
<td>283</td>
<td>140</td>
<td>18.9–907</td>
</tr>
<tr>
<td>Energy through rescuer, J</td>
<td>24</td>
<td>12</td>
<td>0.070–95</td>
</tr>
<tr>
<td>Impedance through patient, Ω</td>
<td>60</td>
<td>15</td>
<td>36–67</td>
</tr>
<tr>
<td>Impedance through rescuer-patient circuit, Ω</td>
<td>2.27×10⁴</td>
<td>1.40×10⁴</td>
<td>1.09×10⁵ to 1.00×10⁵</td>
</tr>
</tbody>
</table>
differed from these standards because leakage current was
recorded directly across rescuers’ bodies instead of a simu-
lated load.

The amount of harm that stray electrical energy could
cause a human depends mostly on the current but also on
voltage, oscillation frequency, current path, and duration of
exposure. Safety standards for the latter variables are not well
defined. The present study noted a low mean voltage (5.8 V),
similar to a single cycle of a 67-Hz alternating current, given
the biphasic nature of the waveform. The extremely short
duration of exposure (~15 ms) resulted in low total leakage
energy. These variables are far less than what would be seen
with accidental contact with an enclosure and further reduce
the likelihood of the average leakage current posing
harm to an individual.

For further perspective on safety, it is useful to compare
the average leakage current we measured through rescuers
to other, more palpable benchmarks of electrical exposure. The
maximum value of leakage current in the present data was
below a well-accepted threshold of perception, which agrees
with the finding that none of the volunteers were able to sense
the shock. The mean leakage current we measured (283 µA)
was also below the average amount of current exposure from
a home body-fat monitoring scale (500 µA) or that used in
cardiac impedance plethysmography.

The ethical considerations involving self-experimentation
by the investigators performing manual contact have been
considered and elaborated on elsewhere. Because this trial
is the first of its kind in its use of human “rescuers” instead of
a simulated load for leakage current measurement, only
investigators with knowledge and experience of the risks of
this type of maneuver participated. This knowledge and
experience came from the routine practice of our electrophys-
iology laboratory, where manual contact comparable to that
used in the present study is frequently made with patients
being defibrillated to reduce the defibrillation threshold and
avoid dislodgement of intracardiac catheters during ablative
procedures. On these grounds, our Institutional Review Board
found that informed consent of the rescuers, because they
were named investigators of the trial, was implicit in the
design and submission of the study itself. Although measure-
ments through investigators themselves introduces potential
bias, we believed that it was unethical to use true “volunteers”
without knowledge or experience in this arena.

The present study has important limitations. Our resucer-
patient model represents only 1 type of cardiac arrest sce-
nario. Rescuers performed simulated compressions while
wearing medical polyethylene gloves. Our measurements do
not address bare skin-to-skin contact between the rescuers’
and the anterior chest of the patient. We also did not
examine leakage current using rigid handheld defibrillation
paddles. The return pathway between rescuer and patient in
the present study is likely to represent a better connection
than with inadvertent contact during actual resuscitations.
Nonetheless, it is possible that more robust conduction
pathways could occur, especially if clothing or bedding is
wet. In addition, poorly adherent defibrillating electrodes
with resultant arcing could put a hands-on rescuer at risk. The
present data, although applicable to a clinically practical
construct of resuscitation, cannot be extrapolated beyond the
constraints of our model.

Leakage measurements were made with the use of elec-
tronodes with an approximate surface area of 4 cm2. This could
be a source of underestimation. Finally, contact pressure on
the anterior chest of the patient was estimated and may have
varied between measurements, possibly affecting the amount
of leakage energy coupled to the rescuer.

**Summary**

Uninterrupted manual chest compressions during shock de-

delivery are feasible. Within the constraints of our model, direct
manual contact with a patient being defibrillated was associ-
ated with exposure to low levels of electrical current. In many cases of cardiac arrest, these observations would allow for elimination of harmful delays from “all-clear” periods during resuscitation.

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**Disclosures**
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**References**

**CLINICAL PERSPECTIVE**

During resuscitation from cardiac arrest due to ventricular fibrillation or ventricular tachycardia, “all-clear” or “hands-off” periods are mandated to avoid potential electrocution of rescuers. These hands-off periods interfere with resuscitation, especially given that current strategies emphasize the importance of avoiding any interruption of chest compressions. There have been substantial changes to external defibrillation technology since hands-off precautions were developed. We addressed the feasibility and safety of direct rescuer-patient contact during defibrillation by measuring the voltage and current through mock rescuers simulating chest compressions on patients receiving external biphasic countershocks with adhesive shock electrodes. The main finding of this work was that the leakage current, the key determinant of potential harm to a rescuer, was very low. Within the constraints of our model, uninterrupted manual chest compressions in a patient being defibrillated are associated with exposure to safe, imperceptible levels of electrical current. In many cases of cardiac arrest, these observations would allow for elimination of routine harmful delays during resuscitation and simplify the resuscitation protocol.
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