Modeling cardiac arrest and resuscitation in the domestic pig

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Abstract

Cardiac arrest remains a leading cause of death and permanent disability worldwide. Although many victims are initially resuscitated, they often succumb to the extensive ischemia-reperfusion injury inflicted on the internal organs, especially the brain. Cardiac arrest initiates a complex cellular injury cascade encompassing reactive oxygen and nitrogen species, $\text{Ca}^{2+}$ overload, ATP depletion, pro- and anti-apoptotic proteins, mitochondrial dysfunction, and neuronal glutamate excitotoxicity, which injures and kills cells, compromises function of internal organs and ignites a destructive systemic inflammatory response. The sheer complexity and scope of this cascade challenges the development of experimental models of damage to vital internal organs following cardiac arrest and cardiopulmonary resuscitation (CPR), and to develop treatments to interrupt the lethal injury cascades. Many experimental animal preparations have been developed to decipher the mechanisms of damage to vital internal organs following cardiac arrest and cardiopulmonary resuscitation (CPR), and to develop treatments to interrupt the lethal injury cascades. Porcine models of cardiac arrest and resuscitation offer several important advantages over other species, and outcomes in this large animal are readily translated to the clinical setting. This review summarizes porcine cardiac arrest-CPR models reported in the literature, describes clinically relevant phenomena observed during cardiac arrest and resuscitation in pigs, and discusses numerous methodological considerations in modeling cardiac arrest/CPR. Collectively, published reports show the domestic pig to be a suitable large animal model of cardiac arrest which is responsive to CPR, defibrillatory countershocks and medications, and yields extensive information to foster advances in clinical treatment of cardiac arrest.

Key words: Acidemia; Asphyxia; Cardiopulmonary
Cardiac arrest remains a leading cause of death worldwide, despite tremendous improvements in emergency medical care and increased public delivery of bystander cardiopulmonary resuscitation (CPR). But progress is being achieved, thanks to the joint efforts of biomedical scientists, physicians and emergency medical personnel to translate laboratory discoveries to the ambulance and hospital. The domestic pig has proven to be a superb preclinical model of cardiac arrest, yielding a wealth of mechanistic insights and practical strategies to refine the delivery of CPR and to test promising treatments. This review examines pivotal factors in modeling cardiac arrest and CPR in the pig.

INTRODUCTION

Prior to 1960 cardiac resuscitation was administered by direct cardiac massage following thoracotomy. Based on animal experimentation a method of external cardiac massage administered by rapid, forceful compressions and passive recoil of the sternum was developed by Kouwenhoven et al[1]. Although over fifty years have passed since the inception of closed chest cardiac massage, and despite many refinements of this approach in the intervening decades, cardiac arrest remains a leading cause of death and persistent disability worldwide. All too often, victims who are initially resuscitated later succumb to extensive ischemia-reperfusion injury to their vital organs, especially the brain[2-5]. Further, many of the 10% of cardiac arrest patients who survive to hospital discharge experience persistent neurocognitive impairment which profoundly impacts their quality of life[3,6].

Although public health data and anecdotal evidence inform the refinement of cardiopulmonary resuscitation (CPR) protocols[7], knowledge of the complex mechanisms of internal organ damage, essential to foster development of effective pharmacological interventions, is incomplete. In the brain, ATP depletion, intracellular Ca²⁺ overload, excessive formation of reactive oxygen and nitrogen derivatives, inflammation and glutamate-induced excitotoxicity conspire to kill neurons and other cells and disrupt the blood-brain barrier. Currently there are no clinically effective pharmacological treatments to protect the brain during cardiac arrest and CPR[2], and therapeutic hypothermia is the only approved treatment in the United States[8]. Reliable preclinical models of cardiac arrest and resuscitation are essential to decipher the injury mechanisms and develop treatments to increase survival and improve quality of life after cardiac arrest.

Ischemia-reperfusion damage in the central nervous system is the result of a multifaceted injury cascade[9,10]. The structural complexity of the brain, which consists of integrated networks of different cell types including neurons, astrocytes, oligodendrocytes, microglia and vascular endothelium, presents fundamental challenges to developing neuroprotective treatments. The brain contains many functional regions which differ in their vulnerabilities to ischemia-reperfusion injury. Potential pharmacotherapeutic agents must first traverse the blood brain barrier, a significant permeability impediment to all but small, non-polar compounds, and act on multiple injury mechanisms, without producing untoward side effects.

Sophisticated animal models are required to model the composite structure and integrated function of the central nervous system and to evaluate the benefits and potential side effects of prospective treatments for ischemia and other brain disorders. Extensive research has established the domestic pig as an excellent animal model to study the impact of cardiac arrest, resuscitation, and therapeutic interventions on the brain and other internal organs. An impressive variety of swine cardiac arrest models are reported in the literature. By examining the features that distinguish these models, this article aims to assist the reader in evaluating the literature and in designing porcine cardiac arrest models appropriate to address specific research objectives.

ATTRIBUTES OF SWINE FOR MODELING CARDIAC ARREST AND RESUSCITATION

Several attributes make the domestic pig an ideal model for cardiac arrest research[11,12]: (1) a large mammal, the pig accommodates extensive instrumentation for blood sampling, monitoring of intravascular and intracardiac pressures, electrocardiography and intravenous administration of medications and experimental treatments; (2) pigs tolerate invasive surgical procedures and rapidly regain consciousness post-anesthesia; (3) resting heart rate, blood pressure, and serum chemistries of pigs and humans are very similar[13-15]; (4) pigs have sufficient blood volume to permit collection of multiple arterial and venous samples for analyses of blood gases and serum chemistry; (5) neurological examinations have been developed to evaluate neurobehavioral function...
Table 1  Details of representative cardiac arrest protocols in pigs

| Ref.                                                                 | Lurie et al[14], 2002 | Mayr et al[15], 2004 | Tang et al[16], 2006 | Li et al[17], 2008 | Indik et al[18], 2009 | Hang et al[19], 2014 |
|---------------------------------------------------------------------|-----------------------|----------------------|----------------------|---------------------|-----------------------|---------------------|
| Pre-anesthetic, induction anesthetic                                | Ketamine 20 mg/kg iv   | Ketamine 20 mg/kg iv  | Ketamine 20 mg/kg iv | Ketamine 20 mg/kg iv | Ketamine 15 mg/kg iv  |
|                                                                     | Propofol 2.5 mg/kg iv  | Propofol 1-2 mg/kg iv| Pentobarbital 30 mg/kg iv | Pentobarbital 30 mg/kg iv | Propofol 0.5 mg/kg iv |
| Maintenance anesthesia                                              | Propofol 10 mg/kg per hour iv | Isoflurane (1%-2%) in 65% nitrous oxide | Pentobarbital 8 mg/kg per hour iv | Pentobarbital 8 mg/kg per hour iv | None |
| Method of arrest                                                    | Electrical: 60 Hz, 140-160 V | Pharmacological: 5 mg/kg bupivacaine | Electrical: 1-2 mA | LAD balloon occluder | Steel plug in LAD |
| Pre-CPR arrest                                                      | 6 min                 | 6 min                 | 7 min                 | 5 min               | 8 min               |
| Preodical compressions (%) (of chest diameter)                      | Mechanical: 80/100 (min %) | Manual: 100/min       | Mechanical: 100/100 (min %) | 100/min (Group 1 25%, Group 2 17.5%) | Manual: 100/100 (c. 33%) |
| CPR duration                                                        | 6 min                 | 2 min                 | 1 min                 | 3 min               | 2 min               |
| Ventilation during CPR?                                            | Fio2 = 1.0, 5:1       | Fio2 = 1.0, 1:15:2    | compression: ventilation | compression: ventilation | Fio2 = 1.0, 1:15:2 |
| Countershocks                                                       | 3.5 x 200 J           | 3.4, 6 J/kg          | 150-360 J            | 150 J               | 150 J               |
| CPR between countershocks                                          | None                  | 1 min/shock           | 3 min                | 2 min               | 4 J/kg              |
| Vasopressors to enhance CPR                                        | EPI 0.045 mg/kg       | AVP 0.4 or 0.8 U/kg  | None                 | None                | EPI 0.02 µg/kg, 1-3 doses |
| Definition of ROSC                                                  | Systolic BP > 70 mmHg | Mean aortic BP > 60 mmHg for > 5 min | Mean aortic BP > 60 mmHg for > 5 min | Systolic BP < 30 mmHg for > 15 min | Systolic BP > 50 mmHg for > 10 min |
| ROSC duration                                                       | 24 h                  | 1 h                   | 3 d                  | 72 h                | 24 h                |
| Pigs completing protocol                                            | Placebo: 0/7; AVP: 5/7; EPI: 4/7; AVP + EPI: 7/7 | Placebo: 0/7; AVP: 5/7; EPI: 4/7; AVP + EPI: 7/7 | Placebo: 0/7; AVP: 5/7; EPI: 4/7; AVP + EPI: 7/7 | Placebo: 0/7; AVP: 5/7; EPI: 4/7; AVP + EPI: 7/7 | Placebo: 0/7; AVP: 5/7; EPI: 4/7; AVP + EPI: 7/7 |

AVP: Vasopressin; BP: Systemic arterial blood pressure; CPR: Cardiopulmonary resuscitation; EPI: Epinephrine; LAD: Left anterior descending coronary artery; ROSC: Recovery of spontaneous circulation.

Factors to consider when modeling cardiac arrest and resuscitation in pigs

The pathophysiological complexities of sudden cardiac death and cardiopulmonary resuscitation challenge the development of animal models that accurately replicate the clinical situation. The primary factors in developing suitable animal models are the study end points and objectives. However, the myriad variables in model design and experimental protocol, which mirror the complexity of cardiac arrest and its treatment, challenge the direct comparison of results obtained in different studies. This section summarizes several factors that must be considered in developing and reporting cardiac arrest-resuscitation protocols in pigs, including the anesthetic regimen, method of inducing ventricular fibrillation, the depth, frequency and duration of chest compressions, whether or not to ventilate during resuscitation and the fraction of inspired O2 (Fio2), the pattern and intensity of defibrillatory countershocks, the criteria taken to indicate recovery of spontaneous circulation (ROSC), the use of inotropic and/or vasoconstrictor support during ROSC, and strategies to correct post-arrest systemic acidemia. With multiple options for each component, it is critical that cardiac arrest-resuscitation protocols be designed carefully to address the study’s specific needs.

FACTORS TO CONSIDER WHEN MODELING CARDIAC ARREST AND RESUSCITATION IN PIGS

- The pig’s large chest accommodates powerful precordial chest compressions and application of transthoracic defibrillatory countershocks of electrical energies similar to those used clinically;
- Pigs have the largest brains among the commonly studied laboratory animals, which provides ample tissue for extensive biochemical and histological analyses of specific brain subregions.
- Porcine models are especially well-suited to study cardiac arrest and CPR, because they are easily tailored to address specific research objectives.
may be released, wires or occluders into the Porcine models of ischemia-induced ventricular or occlusion is the leading cause of cardiac arrest. which persistent cardiac insufficiency might be a confounding factor. induction of ventricular fibrillation does not impart time of ventricular fibrillation onset. Electrical fibrillation. Aside from modeling electrocution-absence of an arterial pulse confirm ventricular "torsades" pattern on electrocardiogram (Figure 1). Ventricular endocardium, a rapid train of impulses to clinical settings. The number and intensity of the countershocks required to restore sinus rhythm is greater in porcine models of ischemically-induced vs electrically-induced arrest, as is the incidence of post-resuscitation ventricular premature beats and recurrence of ventricular fibrillation. Nevertheless, ischemia-induced cardiac arrest replicates the most common cause of cardiac arrest, affording ready translation of results to clinical settings. Asphyxiation is the second most common cause of cardiac arrest and the leading cause in children. A facile method of producing asphyxia in anesthetized swine is to block the endotracheal tube while monitoring the electrocardiogram and arterial blood pressure. Hypoxemia and hypercapnia progressively intensify until cardiac arrest ensues, typically within 10-15 min after blocking ventilation. The principal advantages of asphyxia are its accurate modeling of a major cause of pediatric cardiac arrest and mortality, including the changes in blood gases and pH, and the noninvasive approach which obviates the introduction of pacing wires or occluders into the vasculature. Depending on the study endpoints and objectives, disadvantages may include the changes in blood gas chemistry and the variable duration of asphyxia before ventricular fibrillation, which imposes hypoxemia on the brain and other internal organs even before onset of cardiac arrest. High dosages of certain chemicals, e.g., bupiva-
Caine\(^{[38]}\), may be injected into the right atrium to arrest the heart, modeling cardiac arrest secondary to drug overdose. In such models the potential systemic side effects of the chemicals must be taken into account.

**Duration of pre-CPR arrest**

The duration of pre-intervention arrest is crucial; as this interval is prolonged, cardioversion, survival and good neurological recovery become progressively less achievable. The three-phase model of cardiac arrest\(^{[39]}\) subdivides the pre-intervention period into three phases. The first 4-5 min constitute the electrical phase, during which countershocks are likely to achieve cardioversion even without pre-shock CPR. During the next 5-10 min, the circulatory phase, interventions to effect circulation, e.g., chest compressions, are essential to ensure countershocks produce cardioversion. After 10-15 min arrest, the victim enters the *metabolic phase*, in which increasingly intense metabolic derangements result in protracted or permanent organ damage and severe neurological impairment even if cardioversion is achieved. If the study requires a high survival rate, the period of pre-intervention cardiac arrest may be limited to assure a high rate of defibrillation and ROSC.

**Cardiopulmonary resuscitation: force, frequency and duration**

By affording modest delivery of \(O_2\) and metabolic fuels to the myocardium, precordial compressions may support enough myocardial ATP production to sustain ion transport and repolarize cardiomyocytes, enabling defibrillatory countershocks to restore spontaneous electrical rhythm. Indeed, in a canine cardiac arrest model, effective CPR afforded partial recovery of myocardial Gibbs free energy of ATP hydrolysis\(^{[40]}\), the immediate energy source for cardiac electromechanical activity. Cardiopulmonary resuscitation protocols are readily customized to address the study endpoints. The frequency and depth of precordial compressions can profoundly influence outcome\(^{[41-44]}\). In some studies, CPR is administered by a pneumatic, piston-driven device (e.g., Thumper\(^{®}\)), which can be adjusted to deliver forceful compressions at a predetermined frequency and depth, ensuring consistency of frequency and depth of compressions across experiments\(^{[45]}\). Alternatively, precordial compressions can be administered manually, modeling the CPR given by a bystander responding to an out-of-hospital cardiac arrest. Current American Heart Association guidelines\(^{[45]}\) recommend manual mid-sternal chest compressions be sufficiently forceful to compress the abdomen.
Figure 2  Aortic pressures and lead II electrocardiogram during cardiac arrest, cardiopulmonary resuscitation and recovery of spontaneous circulation. A: Phasic aortic pressure tracing during the period from pre-arrest baseline to 20 min ROSC. Lettered arrows indicate times at which the electrocardiograms shown in panels B-F were obtained. Vertical lines indicate: (1) induction of ventricular fibrillation cardiac arrest; (2) commencement of precordial compressions (CPR); (3) injection of vasopressin (AVP); (4) defibrillation (Defib) by 200 J countershock; and (5) initiation of intravenous phenylephrine (PE; c. 2 µg/kg per minute) to stabilize systemic arterial pressure during ROSC. Mechanical ventilation was suspended during cardiac arrest and CPR. Panels B-F show 5 s electrocardiographic recordings. CPR: Cardiopulmonary resuscitation; ROSC: Recovery of spontaneous circulation; AVP: Vasopressin; PE: Phenylephrine.
chest by one-fourth of its antero-posterior diameter, followed by release and recoil, at a rate of 100 cycles/min.

The duration of CPR before defibrillatory countershocks is an important factor. Longer intervals model the protracted CPR given by bystanders before arrival of the ambulance team, but also lower the likelihood of post-arrest survival with good neurological outcome. Another important consideration is whether or not the animal will be ventilated during CPR and, if so, at what compression: ventilation ratio and FIO₂. Extensive clinical evidence demonstrates that assisted ventilation during CPR following witnessed cardiac arrest offers little or no neurological or survival benefit⁴⁶,⁴⁶-⁴⁸. In accordance with current recommendations for bystander CPR⁵⁰, mechanical ventilation may be suspended for the duration of cardiac arrest and CPR, then resumed after confirming defibrillation to a productive sinus rhythm.

A systemic vasoconstrictor may be administered intravenously to increase the arterial pressures produced by the chest compressions, thereby increasing perfusion of brain and myocardium at the expense of peripheral organs and tissues. The most widely used vasoconstrictors include epinephrine, a physiological adrenergic agonist, and vasopressin, a non-adrenergic vasoconstrictor which may afford greater survival to hospital discharge than epinephrine, especially in patients with asystole⁵¹,⁵².

Although epinephrine has been used in this manner for decades, its potentially detrimental effects, including increased physiological shunt compromising pulmonary gas exchange⁵³,⁵⁴, intensified myocardial ATP consumption and oxygen demand⁵⁵, and the resultant post-resuscitation myocardial dysfunction⁵⁶ and ventricular arrhythmias⁵⁷,⁵⁸ have raised concerns regarding its clinical application for CPR. Preclinical and clinical evidence has shown the non-adrenergic vasoconstrictor vasopressin to be at least as effective as epinephrine at augmenting arterial pressure during precordial compressions, but without epinephrine’s untoward effects. In a porcine cardiac arrest model, vasopressin vs epinephrine produced greater myocardial and brain blood flows and mean arterial pressures during CPR⁵⁹. Thus, vasopressin was associated with higher incidence of conversion to productive sinus rhythm⁶⁰, increased post-arrest cardiac function and decreased morbidity and mortality vs epinephrine. We have found⁶¹ that vasopressin (c. 0.3 U/kg) injected into the right jugular vein at 60 s CPR improved markedly the quality of CPR, increasing the mean arterial pressures from 25-30 to c. 60 mmHg within 3 min (cf. Figure 1). Although epinephrine produced a more abrupt increase in arterial pressure following its injection, within 2 min vasopressin increased mean arterial pressure to a similar extent; during the first 15 min ROSC, the vasopressin-treated swine had less intense tachycardia and more moderate heart rate x arterial pressure product, a measure of myocardial energy expenditure, than their epinephrine-treated counterparts⁶².

Defibrillation and cardioversion
The defibrillation protocol presents the investigator several options for model design. One choice is the sequence of defibrillatory countershocks, i.e., whether the shocks will be administered singly, or in a sequence of multiple (often three) countershocks, before checking for cardioversion. The electrical energies of the countershocks must be considered, including that of the initial countershock, and, if the initial shock fails to achieve cardioversion, whether or at what progression the intensity will be increased for subsequent countershocks. It must be determined if and for how long CPR will be administered during the interval between an unsuccessful cardioversion and the next attempt. When pre-CPR arrest exceeds the electrical phase, bouts of CPR, including a minimum of 20-25 s of chest compressions following unsuccessful countershocks, are essential to ensure effective countershocks. A similar protocol of single shocks with intervening chest compressions increased post-arrest survival vs a conventional 3-shock protocol in a porcine model of ventricular fibrillation cardiac arrest⁶³.

The cardiocirculatory values that constitute ROSC, including the presence of an organized electrical rhythm and maintenance of arterial blood pressure above a predetermined target value for a minimum duration (cf. Table 1) must be specified. Core body temperature has a marked effect on post-arrest and neurological injury and mortality; indeed, moderate hypothermia is the only currently approved intervention consistently shown to produce significant clinical benefit⁶⁴-⁶⁶. Pigs do not thermoregulate effectively while under anesthesia, so typically the animal must be maintained on a heating pad during the cardiac arrest-resuscitation protocol to avoid the impact of hypothermia on study endpoints. Finally, the criteria for abandoning futile resuscitation efforts must be defined.

Post-resuscitation management
Because cardiac arrest imposes ischemia on the heart itself, cardiac mechanical function may be depressed for several hours of ROSC, a manifestation of reversible myocardial injury termed cardiac “stunning”⁶⁵. As the period of ROSC progresses, interventions may be necessary to maintain adequate arterial pressure. Intravenous saline solutions may be infused to expand extracellular fluid volume. Vasopressor agents, e.g., phenylephrine, may be administered, but it should be recognized that vasopressors may lose their efficacy over time due to desensitization of their membrane receptors⁶⁶ and, thus, may be unsuitable for long-
term maintenance of arterial pressure. Accordingly, the vasoconstrictor infusion can be tapered and ultimately discontinued as cardiac function recovers. It may be necessary to adjust tidal volume and frequency of ventilations or administer bicarbonate to compensate for post-arrest hypercapnia and/or acidemia. Isotonic saline (0.9% NaCl) may be infused iv to maintain extracellular fluid volume over the course of the protocol.

**Inspired oxygen concentration**

The oxygen concentration of medical gases used during resuscitation is an important consideration when designing a model of cardiac arrest-resuscitation. For decades, it has been recommended that patients be ventilated with 100% oxygen during resuscitation to increase oxygen delivery to ischemic tissues. Recently, however, hyperoxic ventilation during resuscitation has been shown to intensify formation of reactive oxygen and nitrogen intermediates within tissues and, thus, exacerbate ischemia-reperfusion injury. A recent meta-analysis of clinical trial data showed hyperoxia (PaO2 > 300 mmHg) to be associated with increased in-hospital mortality following cardiac arrest. Oxygen toxicity has been studied for years in a peroperative setting, but only recently has there been sufficient clinical evidence for the European Resuscitation Council to recommend that patients not be ventilated with 100% oxygen after cardiac arrest, but rather with room air supplemented with enough O2 to maintain an oxyhemoglobin saturation (spO2) of 94%-98%. Thus, when designing a cardiac arrest model, the oxygen concentration used during resuscitation may be adjusted depending on whether the study aims to mimic the conventional approach of ventilation with 100% oxygen, or newly recommended strategies such as titration of oxygen administration to maintain a desired spO2.

**CHALLENGES TO MODELING CARDIAC ARREST IN PIGS**

**Pulseless electrical activity**

Pulseless electrical activity (PEA) is a “non-shockable” cardiac electrical rhythm that does not produce ventricular contraction or forward movement of blood. Approximately 60% of out-of-hospital resuscitation attempts result in the development of PEA as the presenting rhythm. Only 2%-5% of patients who present with PEA as their initial rhythm survive to hospital discharge, well below the 15%-40% survival rate of those presenting with ventricular fibrillation. Even fewer patients in whom ventricular fibrillation converted to PEA following countershocks survive to hospital discharge. In our porcine cardiac arrest model, PEA is an ominous finding; typically, even heroic efforts fail to convert PEA to a productive sinus rhythm. None of the 9 pigs developing PEA during resuscitative efforts survived for 4 h ROSC. This situation replicates the clinical setting of out-of-hospital cardiac arrest, where a much lower rate of survival to hospital discharge is achieved in cardiac arrest victims in which PEA is the initial rhythm vs patients with an initial electrocardiographic substrate of ventricular fibrillation.

**Malignant hyperthermia**

A small minority of pigs harbor a genetic lesion in the skeletal muscle sarcoplasmic reticular Ca2+ release channels that predisposes them to develop malignant hyperthermia (aka porcine stress syndrome), often triggered by exposure to volatile anesthetics. Malignant hyperthermia has no overt clinical phenotype detectable by routine screening. As post-arrest survival and neurobehavioral recovery are negatively correlated with body temperature, an episode of malignant hyperthermia, during which core body temperature may rise above 42 ℃, can have disastrous consequences, including systemic hypotension, acidemia, hypercapnia and hyperkalemia that are refractory to conventional interventions. Indeed, in our studies none of the five anesthetized pigs (4% of the total) that developed acute malignant hyperthermia survived to 4 h ROSC, despite aggressive measures including intravenous infusion of ice-cold saline and the K+ chelator calcium gluconate.

**Limitations of porcine models**

An important limitation of many porcine cardiac arrest models is that juvenile, disease-free pigs are generally used. In clinical settings, patients who experience cardiac arrest typically are elderly and suffer from chronic disorders such as hypertension, atherosclerosis, congestive heart failure, diabetes, emphysema or end-stage renal disease. The Ossabaw swine, which is predisposed to develop metabolic syndrome when consuming a high fat diet, provides a unique, clinically relevant experimental model suitable for studying cardiac arrest and resuscitation superimposed on metabolic syndrome. Indeed, under anesthesia these swine develop severe arrhythmias, responsive to amiodarone, that may deteriorate into cardiac arrest (Johnathan D. Tune, personal communication). Unlike most porcine preparations, human victims of out-of-hospital cardiac arrest are not anesthetized when they are stricken. Cardiac arrest is an unanticipated event, and when it occurs outside the hospital, the delays to effective treatments are variable, poorly defined and all too often lethal. Most preclinical cardiac arrest studies employ well defined protocols, such as those reviewed herein. The fundamental differences between these protocols...
and the highly variable and exceedingly challenging clinical situation must be acknowledged.

CONCLUSION

Over the last few decades the collective efforts of many investigators have fostered the development of sophisticated porcine models of cardiac arrest, CPR and ROSC. The domestic pig provides an excellent large animal model of the human cardiovascular system and yields ample tissue for extensive analyses of mechanisms of injury and cytoprotection in the internal organs, such that each experiment generates a wealth of information. Although there is much to consider when constructing an experimental design, the swine model of cardiac arrest-resuscitation is easily tailored to accommodate the desired study end points. The swine model provides unparalleled translational value among current mammalian models of cardiac arrest and CPR, permitting an integrative approach to bridge the gap from bench to bedside.

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