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Special Report—Neonatal Resuscitation: 2010 International Consensus on Cardiopulmonary Resuscitation and Emergency Cardiovascular Care Science With Treatment Recommendations

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Note From the Writing Group: Throughout this article, the reader will notice combinations of superscripted letters and numbers (eg, “Peripartum Suctioning^{NRP-011A, NRP-012A}”). These callouts are hyperlinked to evidence-based worksheets, which were used in the development of this article. An appendix of worksheets, applicable to this article, is located at the end of the text. The worksheets are available in PDF format and are open access.

Approximately 10% of newborns require some assistance to begin breathing at birth, and <1% require extensive resuscitation (LOE 4^{1,2}). Although the vast majority of newborn infants do not require intervention to make the transition from intrauterine to extrauterine life, the large number of births worldwide means that many infants require some assistance to achieve cardiorespiratory stability. Newborn infants who are born at term and are breathing or crying and have good tone must be dried and kept warm. These actions can be provided with the baby lying on the mother’s chest and should not require separation of mother and baby.

All others need to be assessed to determine their need for one or more of the following actions in sequence:

- A. Initial steps in stabilization (dry and provide warmth, position, assess the airway, stimulate to breathe)
- B. Ventilation
- C. Chest compressions
- D. Medications or volume expansion

Progression to the next step is initially based on simultaneous assessment of 2 vital characteristics: heart rate and respirations. Progression occurs only after successful completion of the preceding step. Approximately 30 seconds is allotted to complete each of the first 2 steps successfully, reevaluate, and decide whether to progress to the next (see Figure: Newborn Resuscitation Algorithm).

Since publication of the *2005 International Consensus on CPR and ECC Science With Treatment Recommendations*,^{3,4} several controversial neonatal resuscitation issues have been identified. The literature was researched and a consensus was reached on the assessment of oxygenation and role of supplementary oxygen, peripartum management of meconium, ventilation strategies, devices to confirm placement of an advanced airway (eg, tracheal tube or laryngeal mask airway), med-

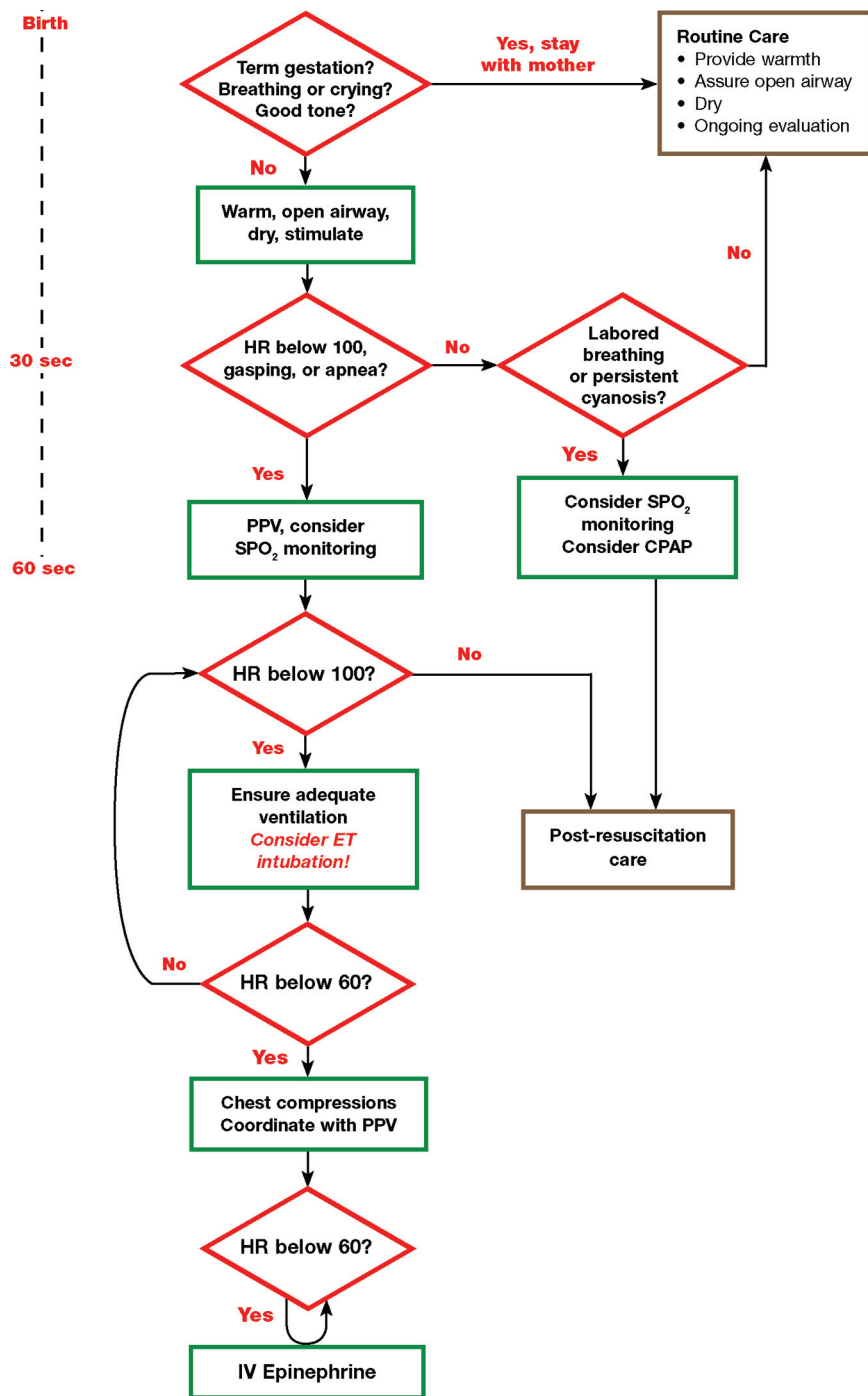


FIGURE.
Newborn Resuscitation Algorithm.

ications, maintenance of body temperature, postresuscitation management, and considerations for withholding and discontinuing resuscitation. Educational techniques for teaching, assessing, and maintaining resuscitation knowledge

and skills and issues regarding the personnel needed at cesarean sections were also debated. The following are the major new recommendations:

- Progression to the next step following the initial evaluation is now

defined by the simultaneous assessment of 2 vital characteristics: heart rate and respirations. Oximetry should be used for evaluation of oxygenation because assessment of color is unreliable.

- For babies born at term it is best to begin resuscitation with air rather than 100% oxygen.
- Administration of supplementary oxygen should be regulated by blending oxygen and air, and the concentration delivered should be guided by oximetry.
- The available evidence does not support or refute the routine endotracheal suctioning of infants born through meconium-stained amniotic fluid, even when the newborn is depressed.
- The chest compression-ventilation ratio should remain at 3:1 for neonates unless the arrest is known to be of cardiac etiology, in which case a higher ratio should be considered.
- Therapeutic hypothermia should be considered for infants born at term or near-term with evolving moderate to severe hypoxic-ischemic encephalopathy, with protocol and follow-up coordinated through a regional perinatal system.
- It is appropriate to consider discontinuing resuscitation if there has been no detectable heart rate for 10 minutes. Many factors contribute to the decision to continue beyond 10 minutes.
- Cord clamping should be delayed for at least 1 minute in babies who do not require resuscitation. Evidence is insufficient to recommend a time for clamping in those who require resuscitation.

INITIAL ASSESSMENT AND INTERVENTION

Assessment of Cardiorespiratory Transition and Need for Resuscitation^{NRP-001A, NRP-001B, NRP-014A, NRP-014B}

Consensus on Science

A prompt increase in heart rate remains the most sensitive indicator of resuscitation efficacy (LOE 5⁵). Of the

clinical assessments, auscultation of the heart is the most accurate, with palpation of the umbilical cord less so. However, both are relatively insensitive (LOE 2⁶ and LOE 4⁷). Several studies have addressed the accuracy of pulse oximetry in measuring heart rate in the delivery room and have shown the feasibility of pulse oximetry during newborn resuscitation. However, none of these studies examined the impact of these measurements on resuscitation outcomes (LOE 4^{7,8}). Pulse oximetry (SpO₂) and heart rate can be measured reliably after 90 seconds from birth with a pulse oximeter designed to reduce movement artifact and a neonatal probe (LOE 4^{9,10}). Preductal values, obtained from the right wrist or hand, are higher than postductal values.^{8,11} Applying the oximeter probe to the subject before connecting it to the instrument will produce reliable results more quickly (LOE 4¹⁰).

There is clear evidence that an increase in oxygenation and improvement in color may take many minutes to achieve, even in uncompromised babies. Furthermore, there is increasing evidence that exposure of the newly born to hyperoxia is detrimental to many organs at a cellular and functional level. For this reason color has been removed as an indicator of oxygenation or resuscitation efficacy. The oximeter can be used to adjust the increase in oxygenation to that of the uncompromised baby born at term.

Treatment Recommendations

Heart rate should remain the primary vital sign by which to judge the need for and efficacy of resuscitation. Auscultation of the precordium should remain the primary means of assessing heart rate. There is a high likelihood of underestimating heart rate with palpation of the umbilical pulse, but this is preferable to other palpation locations.

For babies who require ongoing resuscitation or respiratory support or both, the goal should be to use pulse oximetry. The sensor should be placed on the baby's right hand or wrist before connecting the probe to the instrument. Because of concerns about the ability to consistently obtain accurate measurements, pulse oximetry should be used in conjunction with and should not replace clinical assessment of heart rate during newborn resuscitation.

Use of Supplementary

Oxygen^{NRP-013A, NRP-013B, NRP-014A, NRP-014B}

Consensus on Science

In term infants receiving resuscitation with intermittent positive-pressure ventilation, 100% oxygen conferred no advantage over air in the short term and resulted in increased time to first breath or cry or both (LOE 2^{12,13}). Meta-analyses of these studies showed a decrease in mortality with the group for whom resuscitation was initiated with air.^{14,15}

There is evidence in newborn animal models of asphyxia that exposure to high concentrations of oxygen at resuscitation does not confer any clinical advantage and is potentially harmful at the cellular level.^{16,17} Two animal models of hypoxia-ischemia and persistent bradycardia found that those resuscitated with room air rather than 100% oxygen developed untoward biochemical changes in the brain (LOE 5^{18,19}).

In preterm infants at <32 weeks' gestation, if attempting to mimic the gradual rise in oxygen saturation of healthy term babies in the first 10 minutes after birth by titrating the concentration to the baby's saturation, initial use of air or 100% oxygen is more likely to result in hypoxemia or hyperoxemia, respectively, than initiation of resuscitation with 30% or 90% oxygen and titration to oxygen saturation (LOE 2^{11,20}). There is insufficient evidence in babies

born at 32 to 37 weeks' gestation to define the appropriate oxygen administration strategy.

Treatment Recommendation

In term infants receiving resuscitation at birth with positive-pressure ventilation, it is best to begin with air rather than 100% oxygen. If despite effective ventilation there is no increase in heart rate or if oxygenation (guided by oximetry) remains unacceptable, use of a higher concentration of oxygen should be considered.

Because many preterm babies of <32 weeks' gestation will not reach target saturations in air, blended oxygen and air may be given judiciously and ideally guided by pulse oximetry. Both hyperoxemia and hypoxemia should be avoided. If a blend of oxygen and air is not available, resuscitation should be initiated with air.

Peripartum Suctioning^{NRP-011A, NRP-012A}

Peripartum suctioning was examined from 2 perspectives: (1) suctioning of the airway in depressed neonates born through clear amniotic fluid and (2) tracheal suctioning in depressed neonates born through meconium-stained amniotic fluid.

Suctioning of the Upper Airway

Consensus on Science

There is no evidence to support or refute suctioning of the mouth and nose of depressed neonates at birth when the infant is born through clear amniotic fluid. In healthy neonates suctioning of the mouth and nose is associated with cardiorespiratory complications (LOE 1^{21,22}). In infants who are intubated, sedated, or paralyzed following resuscitation, endotracheal suctioning in the absence of secretions may result in a decrease in oxygenation, an increase in cerebral blood flow and intracranial pressure, and a decrease in compliance (LOE 5²³).

Treatment Recommendation

Routine intrapartum oropharyngeal and nasopharyngeal suctioning for infants born with clear or meconium-stained amniotic fluid is no longer recommended.

Tracheal Suctioning

Consensus on Science

Depressed infants born through meconium-stained amniotic fluid are at increased risk of developing meconium aspiration syndrome (LOE 4^{24,25}). Although these infants are at increased risk of developing meconium aspiration syndrome, the use of tracheal suctioning has not been associated with a reduction in the incidence of meconium aspiration syndrome or mortality (LOE 4²⁶; LOE 5²⁷). No randomized controlled studies have compared intubation and tracheal suctioning and no tracheal suctioning in depressed infants.

Treatment Recommendation

The available evidence does not support or refute the routine endotracheal suctioning of depressed infants born through meconium-stained amniotic fluid.

VENTILATION STRATEGIES^{NRP-028A, NRP-028B}

Ventilation strategies were examined from 4 perspectives: (1) characteristics of the initial assisted breaths and the role of positive end-expiratory pressure (PEEP), (2) continuous positive airway pressure (CPAP) during or following resuscitation, (3) devices to assist ventilation, and (4) strategies when resources are limited.

Initial Breaths

Consensus on Science

Both longer and shorter inspiratory times are in clinical use for initial ventilation in term infants, but there are no randomized controlled trials comparing these 2 approaches. In a small case series in term infants, a pro-

longed initial inflation of 5 seconds produced a 2-fold increase in functional residual capacity compared with historic controls (LOE 4²⁸). A single randomized controlled trial in preterm infants of a 10-second sustained inflation followed by nasal CPAP compared with face mask ventilation demonstrated decreased need for intubation in the first 72 hours, shorter duration of ventilatory support, and reduced bronchopulmonary dysplasia (LOE 1²⁹). Two other randomized controlled trials failed to show a benefit from delivery room application of a sustained initial inflation followed by nasal CPAP (LOE 1^{30,31}). Multiple variables among the 3 randomized controlled trials, including mode of intervention (nasopharyngeal tube versus face mask, T-piece versus self-inflating bag), as well as the use of CPAP in the delivery room make it difficult to determine the effect of the initial sustained inflation on establishing a functional residual capacity in very preterm infants.

Pressure

There is no evidence to support the use of inflation pressures higher than those that are necessary to achieve improvement in heart rate or chest expansion. This can usually be achieved in term infants with an inflation pressure of 30 cm H₂O (LOE 4^{28,32}) and in preterm infants with pressures of 20 to 25 cm H₂O (LOE 4³³). Occasionally higher pressures are required (LOE 4³⁴). In immature animals, ventilation at birth with high tidal volumes associated with the generation of high peak inflation pressures for a few minutes causes lung injury, impaired gas exchange, and reduced lung compliance (LOE 5³⁵).

Positive End-Expiratory Pressure

There is no evidence to support or refute the value of PEEP during resuscitation of term infants. In preterm infants 1 small study did not show a

benefit from PEEP during initial stabilization in reducing the number of infants who required intubation in the delivery room (LOE 1³⁶). In studies of intubated immature animals the use of PEEP during initial stabilization after birth improved functional residual capacity, oxygenation, and lung compliance and reduced lung injury (LOE 5^{37,38}), but high levels of PEEP (8 to 12 cm H₂O) may reduce pulmonary blood flow and increase the risk of pneumothorax (LOE 5^{39,40}).

Treatment Recommendation

To establish initial lung inflation in apneic newborn infants, initiation of intermittent positive-pressure ventilation at birth can be accomplished with either shorter or longer inspiratory times. Initial peak inflating pressures necessary to achieve an increase in heart rate or movement of the chest are variable and unpredictable and should be individualized with each breath. If pressure is being monitored, an initial inflation pressure of 20 cm H₂O may be effective in preterm babies, but a pressure of 30 to 40 cm H₂O may be necessary in some term babies. If pressure is not being monitored, the minimal inflation required to achieve an increase in heart rate should be used. Providers should avoid creation of excessive chest wall movement during ventilation of preterm infants immediately after birth.

Although measured peak inflation pressure does not correlate well with volume delivered in the context of changing respiratory mechanics, monitoring of inflation pressure may help provide consistent inflations and avoid unnecessarily high pressures. If positive-pressure ventilation is required, an initial inflation pressure of 20 to 25 cm H₂O is adequate for most preterm infants. If prompt improvement in heart rate or chest movement is not obtained, then higher pressures to achieve effective ventilation may be needed. PEEP is likely to be beneficial during initial stabilization of apneic

preterm infants who require positive-pressure ventilation and should be used if suitable equipment is available.

Continuous Positive Airway Pressure^{NRP-002A, NRP-002B}

Consensus on Science

For spontaneously breathing preterm infants at ≥ 25 weeks' gestation who have signs of respiratory distress, there is no significant difference between starting CPAP or intubation and mechanical ventilation in the delivery room when considering death or oxygen requirement at 36 weeks postmenstrual age. In spontaneously breathing infants at 25 to 28 weeks' gestation, CPAP compared with intubation reduced the rates of mechanical ventilation from 100% to 46% and surfactant use from 77% to 38% (LOE 1⁴¹). In the same trial infants on CPAP had a significantly increased rate of pneumothorax (9% versus 3%) (LOE 1⁴¹). There is no evidence to support or refute the use of CPAP in the term baby.

For very preterm infants, a multifaceted intervention, including PEEP, giving a sustained inflation and starting CPAP in the delivery room reduces the need for intubation and rate of mechanical ventilation within 72 hours and reduces incidence of bronchopulmonary dysplasia compared with positive-pressure ventilation with a self-inflating bag via a face mask (LOE 1²⁹). When compared with historic controls, use of delivery room CPAP for very premature infants was associated with a decrease in the requirement for intubation, days on mechanical ventilation, and use of postnatal steroids (LOE 4³³), although a small underpowered feasibility trial of delivery room CPAP/PEEP versus no CPAP/PEEP did not show a significant difference in immediate outcomes (LOE 1³⁶).

Treatment Recommendation

Spontaneously breathing preterm infants who have respiratory distress may be supported with CPAP or intuba-

tion and mechanical ventilation. The most appropriate choice may be guided by local expertise and preferences.

Assisted Ventilation

Devices^{NRP-015A, NRP-015B, NRP-015C, NRP-017A, NRP-017B}

Consensus on Science

There are no clinical studies in newborns requiring positive pressure during resuscitation to support or refute the superiority of the T-piece resuscitator over bag-mask ventilation in improving outcome. In mechanical models target inflation pressures are delivered more consistently when using T-piece resuscitators than with self-inflating bags or flow-inflating bags (LOE 5^{42,43}). In mechanical models PEEP is maintained more consistently with T-piece resuscitators compared with self-inflating bags or flow-inflating bags (LOE 5⁴⁴). In mechanical models the ability to deliver a sustained inflation is better with either a T-piece resuscitator or flow-inflating bag than with a self-inflating bag (LOE 5^{42,45}).

Treatment Recommendation

Ventilation of the newborn can be performed effectively with a flow-inflating bag, a self-inflating bag, or a pressure-limited T-piece resuscitator.

Laryngeal Mask Airway^{NRP-017A, NRP-017B}

Consensus on Science

In 1 randomized controlled trial (LOE 1⁴⁶) providers had similar success providing effective ventilation with either the laryngeal mask airway or face mask among newborns in the delivery room. In 1 retrospective cohort study (LOE 2⁴⁷) and 3 large case series (LOE 4⁴⁸) effective ventilation was achieved quickly using a laryngeal mask airway in newborns weighing >2000 g or delivered at ≥ 34 weeks' gestation. In 1 randomized controlled trial (LOE 1⁴⁹) and 1 retrospective cohort study (LOE 2⁵⁰) providers had similar success providing effective ventilation using either

the laryngeal mask airway or endotracheal tube among newborns in the delivery room. Although a single cohort study (LOE 2⁵⁰) suggests that newborns resuscitated with a laryngeal mask may require less respiratory support after initial resuscitation, this conclusion is subject to significant selection bias. In multiple small case reports effective ventilation was achieved with a laryngeal mask airway when both face mask ventilation and endotracheal intubation were unsuccessful. There is limited evidence to evaluate the effectiveness of the laryngeal mask airway for newborns weighing <2000 g, delivered at <34 weeks' gestation, in the setting of meconium-stained amniotic fluid, during chest compressions, or for administration of emergency intratracheal medications.

Treatment Recommendation

The laryngeal mask airway should be considered during resuscitation of the newborn if face mask ventilation is unsuccessful and tracheal intubation is unsuccessful or not feasible. The laryngeal mask airway may be considered as an alternative to a face mask for positive-pressure ventilation among newborns weighing >2000 g or delivered at ≥34 weeks' gestation. There is limited evidence, however, to evaluate its use for newborns weighing <2000 g or delivered at <34 weeks' gestation. The laryngeal mask airway may be considered as an alternative to endotracheal intubation as a secondary airway for resuscitation among newborns weighing >2000 g or delivered at ≥34 weeks' gestation. The laryngeal mask airway has not been evaluated in the setting of meconium-stained amniotic fluid, during chest compressions, or for administration of emergency intratracheal medications.

Upper Airway Interface Devices^{NRP-003A, NRP-003B}

Consensus on Science

Within classes of interfaces, reports conflict about the ability to maintain a seal with an anatomically shaped mask compared with a soft round mask (LOE 5^{51,52}). Delivery of positive-pressure ventilation via nasal prongs has been shown to be superior to delivery via a triangular face mask for outcomes of chest compressions and intubation (LOE 2⁵³). It is likely that differences in clinical outcomes that have been reported in several studies may be attributable to the targeted intervention (ie, CPAP versus intermittent positive-pressure ventilation) rather than the interface. Nasal prongs may be a more effective device than face masks for providing respiratory support after birth (LOE 2⁵³). There is insufficient evidence to support or refute the use of one type of mask over another for achieving clinical outcome, except that the Rendell-Baker style mask is suboptimal in achieving an adequate seal when used for newborns (LOE 5⁵⁴).

Treatment Recommendations

Nasal prongs are an alternative way of giving respiratory support. Whichever interface is used, providers should ensure that they are skilled in using the interface devices available at the institution. Different masks must be held in different ways to appropriately reduce leak.

Exhaled Air Ventilation^{NRP-004A, NRP-004B}

Consensus on Science

Mouth-to-mouth ventilation is less effective than a self-inflating bag or tube and mask in improving survival rates in newborns with birth asphyxia (LOE 3⁵⁵). Use of mouth-to-mask ventilation at 30 insufflations per minute is as effective as self-inflating bag-mask ventilation in increasing heart rate in the first 5 minutes after birth (LOE 2⁵⁶). Mask-to-tube ventilation

may cause infection in newborn infants (LOE 5⁵⁷). Two studies (LOE 5^{58,59}) demonstrated that tube-to-mask ventilation can be easily taught and acceptable breaths delivered. However, tube-to-mask ventilation was more difficult to use (LOE 5⁶⁰; LOE 3⁵⁵).

Treatment Recommendation

Bag-mask ventilation is preferable to mouth-to-mask ventilation or tube-to-mask ventilation during neonatal resuscitation, but one of the latter two should be used when bag-mask devices are not available. Precautions must be taken because mouth-to-mask and mouth tube-to-mask ventilation are less comfortable and more tiring than bag-mask ventilation for the newborn at birth and may be associated with increased risk of infection in the infant and healthcare provider.

MONITORING DURING AND AFTER INTUBATION

Gas Monitoring Devices

Measurement of Tidal Volume^{NRP-005A, NRP-005B, NRP-005C}

Consensus of Science

There are no studies that compare clinical outcomes in newborns after resuscitation with or without monitoring of tidal volume. In preterm animal models the tidal volume used during initial ventilation after birth may alter subsequent lung function and induce inflammation, but other factors, including the use and level of PEEP, appear to interact with tidal volume in determining specific effects (LOE 5^{61,62}). It is unclear whether the absolute tidal volumes used affected outcomes. Studies in manikins and animals (LOE 5^{63,64}) suggest that providers cannot maintain constant pressures or assess delivered volume during manual ventilation. The position of the mask and degree of leak may be improved by the use of a volume monitor (LOE 5⁶⁵).

Treatment Recommendations

Ventilation during newborn resuscitation should aim to adequately inflate the lung while avoiding overinflation. There is insufficient evidence to recommend routine use of tidal volume monitoring in neonates receiving positive-pressure ventilation during resuscitation.

Use of Exhaled CO₂ Detectors to Confirm Tracheal Tube Placement^{NRP-016A}

Consensus on Science

Studies (LOE 2^{66–68}) suggest that detection of exhaled CO₂ confirms tracheal intubation in neonates with cardiac output more rapidly and accurately than clinical assessment alone. False-negative readings have been reported during cardiac arrest (LOE 4⁶⁹) despite models suggesting efficacy (LOE 5⁷⁰). False-positive readings may occur with colorimetric devices contaminated with epinephrine (adrenaline), surfactant, and atropine (LOE 5⁷¹). Neonatal studies have excluded infants who need extensive resuscitation. There is no comparative information to recommend any one method for detection of exhaled CO₂ in the neonatal population.

Treatment Recommendation

Detection of exhaled CO₂ in addition to clinical assessment is recommended as the most reliable method to confirm endotracheal placement in neonates with spontaneous circulation.

Colorimetric CO₂ Detection to Assess Ventilation in Nonintubated Patients^{NRP-018A, NRP-018B, NRP-018C}

Consensus on Science

The use of colorimetric exhaled CO₂ detectors during face mask ventilation of small numbers of preterm infants in the intensive care unit (LOE 4⁷²) and the delivery room (LOE 4⁷³) has been reported and may help identify airway obstruction. It is unclear whether the use of exhaled CO₂ detectors during face mask

ventilation confers additional benefit over clinical assessment alone. No risks attributed to the use of exhaled CO₂ detectors have been identified. The use of exhaled CO₂ detectors with other interfaces (eg, nasal airways, laryngeal masks) during positive-pressure ventilation in the delivery room has not been reported.

Treatment Recommendation

There is insufficient evidence to recommend routine use of colorimetric exhaled CO₂ detectors during mask ventilation of newborns in the delivery room.

CIRCULATORY SUPPORT

Chest

Compressions^{NRP-006A, NRP-006B, NRP-007A, NRP-007B}

Consensus on Science

In animal studies of asphyxial models of cardiac arrest, piglets resuscitated with a combination of chest compressions and ventilations had better outcomes than those resuscitated with ventilations or compressions alone (LOE 5^{74,75}). A further study in piglets suggested that sustained chest compressions had a deleterious effect on myocardial and cerebral perfusion, especially during prolonged resuscitation.⁷⁶

A physiologic mathematical modeling study suggested that using higher compression-ventilation ratios would result in underventilation of asphyxiated infants (LOE 5⁷⁷). The model predicts that between 3 and 5 compressions to 1 ventilation should be most efficient for newborns.

Manikin studies confirm that the 3:1 compression-ventilation ratio provides more ventilations per minute when compared with higher ratios, but the resuscitation is perceived as being more physically taxing, especially when performed by a lone rescuer (LOE 5^{78,79}). Adult manikin studies using 2 rescuers have shown that a 5:1 ratio provides better-quality chest compressions than a 15:2

ratio (LOE 5⁸⁰) but can result in more missed ventilations per cycle (LOE 5⁸¹). A pediatric manikin study of mouth-to-mouth ventilation by a lone lay rescuer found equivalent minute ventilation for both the 15:2 and 5:1 ratios, but the 15:2 ratio produced more chest compressions per minute (LOE 5⁸²). With 2-rescuer CPR provided by nursing students, however, minute ventilation and compressions per minute were increased with the 5:1 ratio compared with the 10:2 and 15:2 ratios (LOE 5⁸³). When the 15:2 ratio was compared with the 30:2 ratio in a 1-rescuer model of medical personnel using adolescent, child, and infant manikins, more compression cycles could be achieved with the 30:2 ratio on all manikins with no apparent effect on quality of compressions (LOE 5⁸⁴). Effect on ventilation, however, was not assessed. One study in children suggested that CPR with rescue breathing is preferable to CPR alone when the arrest is of noncardiac etiology (LOE 5⁸⁵). There are no data regarding the optimum compression-ventilation ratios in neonates or neonatal models of primary cardiac versus predominantly asphyxial arrest.

Evidence from randomized studies in swine models (LOE 5^{86,87}), manikin studies (LOE 5^{84,88}), small case series (LOE 4⁸⁹), and cadavers (LOE 5⁹⁰) support the current practice of favoring the 2-thumb-encircling hands technique of chest compressions when compared with the 2-finger technique. The former method produces higher blood pressure, can sustain a consistent quality of compressions for a longer time, and is perceived as easier and less tiring for the provider. One manikin study involving a variety of medical or quasimedical personnel (LOE 5⁹¹) found no difference in a number of qualitative measures between the 2 techniques other than significantly fewer compressions were judged as too shallow with the 2-thumb technique. One small case series in new-

borns found higher systolic blood pressure generated with the 2-finger technique when compared with the 2 thumb–encircling hands technique (LOE 4⁹²). Both techniques, however, generated comparable and adequate diastolic pressures, a more important determinant of coronary perfusion. Compressions should be centered over the lower third of the sternum rather than the mid-sternum (LOE 5^{93,94}). Chest compression depth should favor one third the external anterior-posterior diameter of the chest rather than deeper compressions (LOE 5⁹⁵).

Treatment Recommendation

There is no evidence from quality human, animal, manikin, or mathematical modeling studies to warrant a change from the current compression-ventilation ratio of 3:1. Strategies should be considered for optimizing the quality of compressions and ventilations with as few interruptions as possible. Because ventilation is critical to reversal of newborn asphyxial arrest, any higher ratio that decreases minute ventilation should be introduced with caution. If the arrest is known to be of cardiac etiology, a higher compression-ventilation ratio should be considered (eg, 15:2).

Chest compressions in the newborn should be delivered by the 2 thumb–encircling hands method as the preferred option. Compressions should be centered over the lower third of the sternum and should compress the chest one third the anterior-posterior diameter. Any chest compressions should be performed in combination with adequate inflation breaths.

MEDICATIONS AND FLUID ADMINISTRATION

Epinephrine

Route and Dose of

Epinephrine NRP-008A, NRP-008B, NRP-009A, NRP-009B

Consensus on Science

Despite the widespread use of epinephrine during resuscitation, no controlled

clinical trials have directly compared endotracheal and intravenous administration of epinephrine among neonates with a heart rate of <60 beats per minute despite adequate ventilation and chest compressions. Limited evidence from neonatal case series or case reports (LOE 4^{96,97}) indicates that epinephrine administered by the endotracheal route using a wide range of doses (0.003 mg/kg to 0.25 mg/kg) may result in return of spontaneous circulation (ROSC) or an increase in heart rate when intravenous access is not available. These case series are limited by inconsistent standards for epinephrine administration and are subject to both selection and reporting bias.

Evidence from 1 case series using rigorously defined standards for epinephrine administration and outcomes reporting indicates that endotracheal administration of epinephrine (0.01 mg/kg) is likely to be less effective than intravenous administration of the same dose (LOE 4²). This is consistent with evidence extrapolated from neonatal animal models indicating that higher doses (0.05 mg/kg to 0.1 mg/kg) of endotracheal epinephrine may be required to achieve increased blood epinephrine concentrations and a hemodynamic response equivalent to intravenous administration (LOE 5^{98,99}). Evidence extrapolated from adult animal models indicates that blood concentrations of epinephrine are significantly lower following endotracheal administration (LOE 5^{100,101}), and endotracheal doses ranging from 0.05 mg/kg to 0.1 mg/kg may be required to achieve ROSC (LOE 5¹⁰²).

Although it has been widely assumed that epinephrine can be administered faster by the endotracheal route than by the intravenous route, no clinical trials have evaluated this hypothesis. Two studies have reported cases of inappropriate early use of endotracheal epinephrine before airway and breathing are established (LOE 4^{96,97}). One

case series describing in-hospital pediatric cardiac arrest suggested that survival was higher among infants who received their first dose of epinephrine by the endotracheal route; however, the time required for first dose administration using the endotracheal and intravenous routes was not provided (LOE 5¹⁰³).

Despite the widespread use of epinephrine during resuscitation, no controlled clinical trials have evaluated the ideal dose of epinephrine among neonates with a heart rate of <60 beats per minute despite adequate ventilation and chest compressions. Evidence extrapolated from pediatric studies that included infants <1 year of age (LOE 5^{104,105}) indicate no benefit from intravenous epinephrine doses ≥ 0.03 mg/kg. This is in contrast to a single pediatric case series using historic controls that indicated a marked improvement in ROSC using high-dose intravenous epinephrine (0.1 mg/kg) among children who had not responded to 2 doses of standard epinephrine (0.01 mg/kg) (LOE 5¹⁰⁶). Further extrapolative evidence from a meta-analysis of 5 adult clinical trials indicates that high-dose intravenous epinephrine may increase ROSC but offers no benefit in survival to hospital discharge (LOE 5¹⁰⁷). Evidence from a planned secondary analysis of a pediatric randomized controlled trial suggests an increased risk of mortality among children receiving high-dose intravenous epinephrine (0.1 mg/kg) (LOE 5¹⁰⁴). Additional evidence from 2 pediatric animal studies (LOE 5^{108,109}) indicates that intravenous epinephrine ≥ 0.1 mg/kg increased risk of postresuscitation mortality and interfered with cerebral cortical blood flow and cardiac output. There are no published studies comparing standard- and high-dose endotracheal epinephrine in the neonatal population with hypoxic-hypercarbic arrest, and the ideal dose for endotracheal administration is unknown. Data from

neonatal case series and animal models suggest that higher doses (0.05 mg/kg to 0.1 mg/kg) of endotracheal epinephrine may be required to achieve increased blood epinephrine concentrations and a hemodynamic response equivalent to intravenous administration (LOE 4^{2,96}).

Treatment Recommendation

If adequate ventilation and chest compressions have failed to increase the heart rate to >60 beats per minute, then it is reasonable to use epinephrine despite the lack of human neonatal data. If epinephrine is indicated, a dose of 0.01 to 0.03 mg/kg should be administered *intravenously* as soon as possible. If adequate ventilation and chest compressions have failed to increase the heart rate to >60 beats per minute and intravenous access is *not* available, then it is reasonable to administer endotracheal epinephrine. If epinephrine is administered by the endotracheal route, it is likely that a larger dose (0.05 mg/kg to 0.1 mg/kg) will be required to achieve an effect similar to that of the 0.01 mg/kg intravenous dose. Higher intravenous doses cannot be recommended and may be harmful.

Volume Expansion^{NRP-029A, NRP-029B, NRP-029C}

Consensus on Science

Multiple case series support the use of volume expansion in babies with a history of blood loss, including some who are unresponsive to chest compressions (LOE 4¹¹⁰). Many with pallor and tachycardia responded to volume expansion without having received chest compressions. In the absence of a history of blood loss there is limited evidence of benefit from administration of volume during resuscitation unresponsive to chest compressions/epinephrine (LOE 4¹¹¹) and some suggestion of potential harm from animal studies (LOE 5^{112,113}).

Treatment Recommendation

Early volume replacement with crystalloid or red cells is indicated for babies

with blood loss who are not responding to resuscitation. There is insufficient evidence to support the routine use of volume administration in the infant with no blood loss who is refractory to ventilation, chest compressions, and epinephrine. Because blood loss may be occult, a trial of volume administration may be considered in babies who do not respond to resuscitation.

Other Drugs

Very rarely a narcotic antagonist (naloxone), sodium bicarbonate^{NRP-021A, NRP-021B}, or vasopressors may be useful after resuscitation.

Naloxone^{NRP-022A, NRP-022B}

Consensus on Science

There are no data comparing naloxone with positive-pressure ventilation as the main intervention for opioid-exposed newborn infants who are apneic at birth. For newborns who are vigorous in the delivery room despite maternal use of opioids, naloxone subtly increases ventilation parameters (such as increased alveolar ventilation and improved CO₂ response curves) for a short time, but the clinical relevance of these observations is questionable (LOE 4¹¹⁴). Several other studies found no difference between vigorous treatment with naloxone and placebo or no drug treatment for newborns with outcomes of pH, Pco₂, Apgar scores, and neurologic outcomes (LOE 5¹¹⁵). Studies examining naloxone have consistently demonstrated that it is frequently misused (LOE 4¹¹⁶). Naloxone given to a baby born to an opioid-addicted mother has been associated with seizures (LOE 5¹¹⁷). There are concerns about short- and long-term safety of naloxone in neonates (LOE 5¹¹⁸). Naloxone is absorbed more effectively when given intravenously but has a shorter half-life compared with intramuscular administration.

Treatment Recommendation

Naloxone is not recommended as part of the initial resuscitation for newborns with respiratory depression in the delivery room. For the clinical situation of a newborn with respiratory depression after maternal opiate exposure, the focus needs to remain on effective ventilation and airway support for the persistently apneic newborn.

Vascular Access^{NRP-020A}

Consensus on Science

Multiple clinical series and case reports suggest that fluids and medications can be successfully delivered by the intraosseous route during resuscitation of neonates when equipment or personnel skilled in establishing venous access are not available or if other vascular access sites (especially intravenous) cannot be successfully established within several minutes (LOE 4^{119,120}).

Treatment Recommendation

Temporary intraosseous access to provide fluids and medications to resuscitate critically ill neonates may be indicated following unsuccessful attempts to establish intravenous vascular access or when caregivers are more skilled at securing intraosseous access.

SUPPORTIVE THERAPY

Temperature Control

Maintenance of Body Temperature^{NRP-023A}

Consensus on Science

A large body of evidence supports the wrapping of newborn infants of <28 weeks' gestation in polythene wraps or bags at birth without drying to reduce heat loss (LOE 1^{121,122}). Some of these infants were hyperthermic on admission to the neonatal intensive care unit, but it is unclear whether this is because they were born hot or be-

cause they became overheated during stabilization and transfer. In the absence of polythene wrapping, use of exothermic mattresses maintained the temperature of newborn infants weighing <1500 g within the normal range (LOE 2¹²³). A combination of exothermic mattresses and polythene wrapping during resuscitation is the most effective strategy to avoid hypothermia but may increase the risk of hyperthermia (LOE 3¹²⁴). Delivery room temperatures of at least 26°C for newborns at <28 weeks' gestation in combination with polythene wraps or bags maintained temperatures most effectively (LOE 4¹²⁵; LOE 3¹²⁶).

Treatment Recommendation

Newborn infants of <28 weeks' gestation should be completely covered in a polythene wrap or bag up to their necks without drying immediately after birth and then placed under a radiant heater and resuscitated or stabilized in a standard fashion. Infants should be kept wrapped until admission and temperature check. Hyperthermia should be avoided. Delivery room temperatures should be at least 26°C for infants of <28 weeks' gestation.

POSTRESUSCITATION MANAGEMENT

Temperature

Hyperthermia^{NRP-051A, NRP-051B}

Consensus on Science

Infants born to febrile mothers have been reported to have a higher incidence of perinatal respiratory depression, neonatal seizures, cerebral palsy, and increased risk of mortality (LOE 4^{127,128}). There is no evidence to determine whether the fever or the cause of the fever increases the risk to the baby. In 1 study, neonatal fever at birth resolved spontaneously within 60 minutes (LOE 4¹²⁹). Adult animal trials show decreased central nervous system injury with antipyretic therapy for hyperthermia (LOE 5¹³⁰). In a ran-

domized study high-dose corticosteroids lowered maternal temperature but were associated with an increased number of cases of asymptomatic bacteremia in neonates (LOE 2¹³¹).

Treatment Recommendation

There is insufficient evidence to support or refute the routine use of interventions to lower maternal fever to reduce neonatal morbidity and mortality. There should be an increased awareness that the presence of maternal hyperthermia may lead to a need for neonatal resuscitation. The goal is to achieve normothermia and avoid iatrogenic hyperthermia.

Therapeutic

Hypothermia^{NRP-024A, NRP-024B}

Consensus on Science

A large body of evidence from 3 large randomized studies (LOE 1^{132–134}) and 2 small randomized trials (LOE 1^{135,136}) demonstrated that induced hypothermia (33.5° to 34.5°C) implemented within 6 hours of birth in term infants at highest risk for brain injury (as defined by specific protocols) and with further treatment in neonatal intensive care units is associated with significantly fewer deaths and less neurodevelopmental disability at 18-month follow-up. The number needed to treat is 9.¹³⁷ Both cooling methods (systemic versus selective head cooling) were shown to be effective, but none of the studies compared them directly. The randomized trials produced remarkably consistent results despite using different methods of cooling.¹³⁸

Treatment Recommendations

Newly born infants born at or near-term with evolving moderate to severe hypoxic-ischemic encephalopathy should be offered therapeutic hypothermia. Whole body cooling and selective head cooling are both appropriate strategies. Cooling should be initiated and conducted under clearly defined

protocols with treatment in neonatal intensive care facilities and with the capability for multidisciplinary care. Treatment should be consistent with the protocols used in the randomized clinical trials (ie, begin within 6 hours of birth, continue for 72 hours after birth, and rewarm over at least 4 hours). Carefully monitor for known adverse effects of cooling, eg, thrombocytopenia and hypotension. All treated infants should be followed up longitudinally.

General Supportive Care

Glucose^{NRP-019A, NRP-019B}

Consensus on Science

Newborns with lower blood glucose levels have a higher incidence of brain injury and adverse outcomes after a hypoxic-ischemic insult, although no specific level associated with worse outcome has been identified (LOE 4¹³⁹; LOE 3¹⁴⁰). Increased glucose levels after hypoxia-ischemia do not appear to have adverse effects in studies of pediatric patients (LOE 5¹⁴¹) or in animal studies (LOE 5¹⁴²) and may be protective (LOE 5¹⁴³). However, no randomized controlled trials have examined this question. Due to the paucity of data, no specific target glucose concentration range can be identified at present.

Treatment Recommendation

Intravenous glucose infusion should be considered as soon as practical after resuscitation, with the goal of avoiding hypoglycemia.

Timing of Cord

Clamping^{NRP-030A, NRP-030B, NRP-030C, NRP-030D}

Consensus on Science

For the uncomplicated birth at term there is evidence of a benefit to delaying cord clamping for a minimum time ranging from 1 minute until the cord stops pulsating after delivery. Those with delayed clamping had improved iron status through early infancy but were more likely to receive phototherapy (LOE 1¹⁴⁴).

For an otherwise uncomplicated preterm birth, there is evidence of a benefit to delaying cord clamping for a minimum time ranging from 30 seconds to 3 minutes after delivery. Those who experienced delayed clamping in this group had higher blood pressures during stabilization and a lower incidence of intraventricular hemorrhage (LOE 1¹⁴⁵) and received fewer blood transfusions¹⁴⁵ but were more likely to receive phototherapy (LOE 2¹⁴⁴). There are limited data on the hazards or benefits of delayed cord clamping in the nonvigorous infant.^{146,147}

Treatment Recommendation

Delay in umbilical cord clamping for at least 1 minute is recommended for newborn infants not requiring resuscitation. There is insufficient evidence to support or refute a recommendation to delay cord clamping in babies requiring resuscitation.

WITHHOLDING OR DISCONTINUING RESUSCITATIVE EFFORTS^{NRP-025A, NRP-025B, NRP-025C, NRP-026A, NRP-026B, NRP-026C, NRP-027A, NRP-027B}

Noninitiation of Resuscitation

Consensus on Science

For neonates at the margins of viability or those with conditions which predict a high risk of mortality or morbidity, attitudes and practice vary according to region and availability of resources (LOE 4¹⁴⁸). Social science studies indicate that parents would like a larger role in the decisions to start resuscitation and continue life support of severely compromised newborns. Opinions among neonatal providers vary widely regarding the benefits and disadvantages of aggressive therapies in such newborns (LOE 4^{149,150}). Some data are available to help identify conditions associated with high mortality and poor outcome (LOE 4^{151,152}). Such conditions may include extreme prematurity and anomalies that predict extreme morbidity or

early death. Treatment and outcome of infants at the margins of viability may be influenced by factors in addition to gestational age and birthweight.¹⁵³ Noninitiation of resuscitation and withdrawal of cardiorespiratory support are ethically equivalent.¹⁵⁴

Treatment Recommendation

When gestation, birth weight, or congenital anomalies are associated with almost certain early death and an unacceptably high morbidity is likely among the rare survivors, resuscitation is not indicated. In conditions associated with a high rate of survival and acceptable morbidity, resuscitation is nearly always indicated. In conditions associated with uncertain prognosis, when there is borderline survival and a relatively high rate of morbidity and when the burden to the child is high, the parents' views on resuscitation should be supported. There should be a consistent and coordinated approach from the obstetric and neonatal teams in applying these guidelines and in communicating with the parents in developing an agreed-upon management plan when possible. Once resuscitation is initiated it may be appropriate to subsequently decide to discontinue cardiorespiratory support and offer comfort care.

Discontinuation of Resuscitation

Consensus on Science

Available evidence, albeit from relatively small numbers of babies, suggests that babies born without a heart rate that has not returned by 10 minutes of age are likely to either die or have severe neurologic disability (LOE 4^{155,156}). It is not known whether there was significant selection bias in many of these studies, nor indeed that the babies included in these studies did receive "good-quality resuscitation." One study with a large contemporary cohort of infants (some randomized to postresuscitation hypothermia) indicates that in babies born without a detectable heart rate, the lack of ROSC

after 10 minutes of age is associated with survival without severe neurologic deficit in a small number of the survivors (LOE 4¹⁵⁷). Data are not available regarding the number of infants who were deemed too sick for study entry or who died before enrollment. These factors may have resulted in a significant overestimation of the rate of intact survival among infants with an Apgar score of 0 at 10 minutes. In all reported series the cause of the asphyxia and the efficacy of the resuscitation process were not elucidated.

The evidence from 7 LOE 5 studies^{157,158} is insufficient to support or refute any recommendation regarding how much time should elapse with a heart rate of <60 but >0 beats per minute before discontinuing resuscitative efforts.

Treatment Recommendation

In a newly born baby with no detectable heart rate which remains undetectable for 10 minutes, it is appropriate to then consider stopping resuscitation. The decision to continue resuscitation efforts when the infant has a heart rate of 0 for longer than 10 minutes is often complex and may be influenced by issues such as the presumed etiology of the arrest, gestation of the baby, potential reversibility of the situation, and the parents' previously expressed feelings about acceptable risk of morbidity.

The evidence of outcome when the heart rate is <60 beats per minute at birth and persists after 10 or 15 minutes of continuous and adequate resuscitative efforts at delivery is insufficient to guide decisions as to whether to withhold or to continue resuscitation.

Personnel Needs at Elective Cesarean Sections^{NRP-010A, NRP-010B, NRP-010C}

Consensus on Science

Retrospective studies show that delivery by cesarean section at term under regional anesthesia is associated with a

small increase in risk of receiving assisted ventilation during neonatal resuscitation compared with unassisted vaginal birth. The number needed to treat equals 35 (LOE 4^{159,160}). Five retrospective studies showed that delivery by cesarean section at term under regional anesthesia did not increase the risk of requirement for intubation during neonatal resuscitation compared with unassisted vaginal birth (LOE 4^{161,162}). There is no evidence addressing this question in babies born at 34 to 36 weeks' gestation.

Treatment Recommendations

When an infant without antenatally identified risk factors is delivered at term by cesarean section under regional anesthesia, a provider capable of performing assisted ventilation should be present at the delivery. It is not necessary for a provider skilled in neonatal intubation to be present at that delivery.

EDUCATIONAL TECHNIQUES FOR TEACHING, ASSESSING, AND MAINTAINING RESUSCITATION KNOWLEDGE AND SKILLS

Simulation^{NRP-032A, NRP-032B, NRP-032C, EIT-019A, EIT-019B}

Consensus on Science

There is a lack of uniformity in the definition of simulation as a learning

methodology, determination of relevant outcomes, and use of appropriate measurement tools. Use of simulation as an adjunct to traditional education methodologies may enhance performance of healthcare professionals in actual clinical settings (LOE 1¹⁶³; LOE 3¹⁶⁴) and simulated resuscitations (LOE 1¹⁶⁵; LOE 2¹⁶⁶). Some studies did not show any difference in performance between standard training and simulation training in a clinical setting (LOE 1¹⁶⁷) or using other means of evaluation (LOE 1¹⁶⁸). No studies were found that revealed simulation-based training produced inferior results compared with traditional methodologies.

Treatment Recommendations

Simulation should be used as a methodology in resuscitation education. The most effective interventions and evaluation methodologies remain to be defined.

Briefings and

Debriefings^{NRP-033A, NRP-033B, EIT-001A, EIT-001B}

Consensus on Science

Evidence from 1 prospective randomized controlled study (LOE 1¹⁶⁹) and 17 other studies (LOE 3 to 4) of briefings and debriefings document improvement in the acquisition of content

knowledge, technical skills, or behavioral skills required for effective and safe resuscitation. Only a single study (LOE 4¹⁷⁰) revealed no effect of briefing/debriefing on performance, and no studies indicated that the use of briefings and debriefings had any negative effects.

Treatment Recommendations

It is reasonable to recommend the use of briefings and debriefings during learning activities while caring for simulated patients and during clinical activities.

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REFERENCES

1. Perlman JM, Risser R. Cardiopulmonary resuscitation in the delivery room: associated clinical events. *Arch Pediatr Adolesc Med.* 1995;149:20–25
2. Barber CA, Wyckoff MH. Use and efficacy of endotracheal versus intravenous epinephrine during neonatal cardiopulmonary resuscitation in the delivery room. *Pediatrics.* 2006;118:1028–1034
3. 2005 International Consensus on Cardiopulmonary Resuscitation and Emergency Cardiovascular Care Science With Treatment Recommendations. Part 7: Neonatal Resuscitation. *Resuscitation.* 2005;67:293–303
4. 2005 International Consensus on Cardiopulmonary Resuscitation and Emergency Cardiovascular Care Science With Treatment Recommendations. Part 7: Neonatal Resuscitation. *Circulation.* 2005;112:III-91–III-99
5. Dawes GS. Foetal and neonatal physiology. Chicago, IL: Year Book Medical Publishers; 1968:149
6. Owen CJ, Wyllie JP. Determination of heart rate in the baby at birth. *Resuscitation.* 2004;60:213–217
7. Kamlin CO, Dawson JA, O'Donnell CP, Morley CJ, Donath SM, Sekhon J, Davis PG. Accuracy of pulse oximetry measurement of heart rate of newborn infants in the delivery room. *J Pediatr.* 2008;152:756–760
8. Dawson JA, Kamlin CO, Wong C, Te Pas AB, O'Donnell CP, Donath SM, Davis PG, Morley CJ. Oxygen saturation and heart rate during delivery room resuscitation of infants <30 weeks' gestation with air or 100% oxygen. *Arch Dis Child Fetal Neonatal Ed.* 2009;94:F87–F91
9. Altuncu E, Ozek E, Bilgen H, Topuzoglu A, Kavuncuoglu S. Percentiles of oxygen saturations in healthy term newborns in the first minutes of life. *Eur J Pediatr.* 2008;167:687–688
10. O'Donnell CP, Kamlin CO, Davis PG, Morley CJ. Obtaining pulse oximetry data in neonates: a randomised crossover study of sensor application techniques. *Arch Dis Child Fetal Neonatal Ed.* 2005;90:F84–F85
11. Wang CL, Anderson C, Leone TA, Rich W, Govindaswami B, Finer NN. Resuscitation of preterm neonates by using room air or 100% oxygen. *Pediatrics.* 2008;121:1083–1089
12. Vento M, Asensi M, Sastre J, Garcia-Sala F, Pallardo FV, Vina J. Resuscitation with

- room air instead of 100% oxygen prevents oxidative stress in moderately asphyxiated term neonates. *Pediatrics*. 2001;107:642–647
13. Saugstad OD, Rootwelt T, Aalen O. Resuscitation of asphyxiated newborn infants with room air or oxygen: an international controlled trial: the RESAIR 2 study. *Pediatrics*. 1998;102:e1
 14. Davis PG, Tan A, O'Donnell CP, Schulze A. Resuscitation of newborn infants with 100% oxygen or air: a systematic review and meta-analysis. *Lancet*. 2004;364:1329–1333
 15. Rabi Y, Rabi D, Yee W. Room air resuscitation of the depressed newborn: a systematic review and meta-analysis. *Resuscitation*. 2007;72:353–363
 16. Lakshminrusimha S, Russell JA, Steinhorn RH, Swartz DD, Ryan RM, Gugino SF, Wynn KA, Kumar VH, Mathew B, Kirmani K, Morin FC III. Pulmonary hemodynamics in neonatal lambs resuscitated with 21%, 50%, and 100% oxygen. *Pediatr Res*. 2007;62:313–318
 17. Solberg R, Andresen JH, Escrig R, Vento M, Saugstad OD. Resuscitation of hypoxic newborn piglets with oxygen induces a dose-dependent increase in markers of oxidation. *Pediatr Res*. 2007;62:559–563
 18. Solas AB, Kutzsche S, Vinje M, Saugstad OD. Cerebral hypoxemia-ischemia and reoxygenation with 21% or 100% oxygen in newborn piglets: effects on extracellular levels of excitatory amino acids and microcirculation. *Pediatr Crit Care Med*. 2001;2:340–345
 19. Presti AL, Kishkurno SV, Slinko SK, Randis TM, Ratner VI, Polin RA, Ten VS. Reoxygenation with 100% oxygen versus room air: late neuroanatomical and neurofunctional outcome in neonatal mice with hypoxic-ischemic brain injury. *Pediatr Res*. 2006;60:55–59
 20. Escrig R, Arruza L, Izquierdo I, Villar G, Saenz P, Gimeno A, Moro M, Vento M. Achievement of targeted saturation values in extremely low gestational age neonates resuscitated with low or high oxygen concentrations: a prospective, randomized trial. *Pediatrics*. 2008;121:875–881
 21. Gungor S, Kurt E, Teksoz E, Goktolga U, Ceyhan T, Baser I. Oronasopharyngeal suction versus no suction in normal and term infants delivered by elective cesarean section: a prospective randomized controlled trial. *Gynecol Obstet Invest*. 2006;61:9–14
 22. Waltman PA, Brewer JM, Rogers BP, May WL. Building evidence for practice: a pilot study of newborn bulb suctioning at birth. *J Midwifery Womens Health*. 2004;49:32–38
 23. Simbruner G, Coradello H, Fodor M, Havelec L, Lubec G, Pollak A. Effect of tracheal suction on oxygenation, circulation, and lung mechanics in newborn infants. *Arch Dis Child*. 1981;56:326–330
 24. Usta IM, Mercer BM, Sibai BM. Risk factors for meconium aspiration syndrome. *Obstet Gynecol*. 1995;86:230–234
 25. Rossi EM, Philipson EH, Williams TG, Kalhan SC. Meconium aspiration syndrome: intrapartum and neonatal attributes. *Am J Obstet Gynecol*. 1989;161:1106–1110
 26. Al Takroni AM, Parvathi CK, Mendis KB, Hassan S, Reddy I, Kudair HA. Selective tracheal suctioning to prevent meconium aspiration syndrome. *Int J Gynaecol Obstet*. 1998;63:259–263
 27. Gupta V, Bhatia BD, Mishra OP. Meconium stained amniotic fluid: antenatal, intrapartum and neonatal attributes. *Indian Pediatr*. 1996;33:293–297
 28. Vyas H, Milner AD, Hopkin IE, Boon AW. Physiologic responses to prolonged and slow-rise inflation in the resuscitation of the asphyxiated newborn infant. *J Pediatr*. 1981;99:635–639
 29. Te Pas AB, Walther FJ. A randomized, controlled trial of delivery-room respiratory management in very preterm infants. *Pediatrics*. 2007;120:322–329
 30. Lindner W, Hogel J, Pohlandt F. Sustained pressure-controlled inflation or intermittent mandatory ventilation in preterm infants in the delivery room? A randomized, controlled trial on initial respiratory support via nasopharyngeal tube. *Acta Paediatr*. 2005;94:303–309
 31. Harling AE, Beresford MW, Vince GS, Bates M, Yoxall CW. Does sustained lung inflation at resuscitation reduce lung injury in the preterm infant? *Arch Dis Child Fetal Neonatal Ed*. 2005;90:F406–F410
 32. Boon AW, Milner AD, Hopkin IE. Lung expansion, tidal exchange, and formation of the functional residual capacity during resuscitation of asphyxiated neonates. *J Pediatr*. 1979;95:1031–1036
 33. Lindner W, Vossbeck S, Hummler H, Pohlandt F. Delivery room management of extremely low birth weight infants: spontaneous breathing or intubation? *Pediatrics*. 1999;103:961–967
 34. Upton CJ, Milner AD. Endotracheal resuscitation of neonates using a rebreathing bag. *Arch Dis Child*. 1991;66:39–42
 35. Hillman NH, Moss TJ, Kallapur SG, Bachurski C, Pillow JJ, Polglase GR, Nitsos I, Kramer BW, Jobe AH. Brief, large tidal volume ventilation initiates lung injury and a systemic response in fetal sheep. *Am J Respir Crit Care Med*. 2007;176:575–581
 36. Finer NN, Carlo WA, Duara S, Fanaroff AA, Donovan EF, Wright LL, Kandefer S, Poole WK. Delivery room continuous positive airway pressure/positive end-expiratory pressure in extremely low birth weight infants: a feasibility trial. *Pediatrics*. 2004;114:651–657
 37. Siew ML, Te Pas AB, Wallace MJ, Kitchen MJ, Lewis RA, Fouras A, Morley CJ, Davis PG, Yagi N, Uesugi K, Hooper SB. Positive end-expiratory pressure enhances development of a functional residual capacity in preterm rabbits ventilated from birth. *J Appl Physiol*. 2009;106:1487–1493
 38. Te Pas AB, Siew M, Wallace MJ, Kitchen MJ, Fouras A, Lewis RA, Yagi N, Uesugi K, Donath S, Davis PG, Morley CJ, Hooper SB. Establishing functional residual capacity at birth: the effect of sustained inflation and positive end-expiratory pressure in a preterm rabbit model. *Pediatr Res*. 2009;65:537–541
 39. Polglase GR, Hooper SB, Gill AW, Allison BJ, McLean CJ, Nitsos I, Pillow JJ, Kluckow M. Cardiovascular and pulmonary consequences of airway recruitment in preterm lambs. *J Appl Physiol*. 2009;106:1347–1355
 40. Probyn ME, Hooper SB, Dargaville PA, McCallion N, Crossley K, Harding R, Morley CJ. Positive end expiratory pressure during resuscitation of premature lambs rapidly improves blood gases without adversely affecting arterial pressure. *Pediatr Res*. 2004;56:198–204
 41. Morley CJ, Davis PG, Doyle LW, Brion LP, Hascoet JM, Carlin JB. Nasal CPAP or intubation at birth for very preterm infants. *N Engl J Med*. 2008;358:700–708
 42. Oddie S, Wyllie J, Scally A. Use of self-inflating bags for neonatal resuscitation. *Resuscitation*. 2005;67:109–112
 43. Hussey SG, Ryan CA, Murphy BP. Comparison of three manual ventilation devices using an intubated mannequin. *Arch Dis Child Fetal Neonatal Ed*. 2004;89:F490–F493
 44. Finer NN, Rich W, Craft A, Henderson C. Comparison of methods of bag and mask ventilation for neonatal resuscitation. *Resuscitation*. 2001;49:299–305
 45. Bennett S, Finer NN, Rich W, Vaucher Y. A comparison of three neonatal resuscitation devices. *Resuscitation*. 2005;67:113–118
 46. Singh R. Controlled trial to evaluate the use of LMA for neonatal resuscitation. *J Anaesth Clin Pharmacol*. 2005;21:303–306
 47. Trevisanuto D, Micaglio M, Pitton M, Maga-

- rotto M, Piva D, Zanardo V. Laryngeal mask airway: is the management of neonates requiring positive pressure ventilation at birth changing? *Resuscitation*. 2004;62:151–157
48. Gandini D, Brimacombe JR. Neonatal resuscitation with the laryngeal mask airway in normal and low birth weight infants. *Anesth Analg*. 1999;89:642–643
 49. Esmail N, Saleh M, et al. Laryngeal mask airway versus endotracheal intubation for Apgar score improvement in neonatal resuscitation. *Egypt J Anesthesiol*. 2002;18:115–121
 50. Zanardo V, Simbi AK, Savio V, Micaglio M, Trevisanuto D. Neonatal resuscitation by laryngeal mask airway after elective cesarean section. *Fetal Diagn Ther*. 2004;19:228–231
 51. Palme C, Nystrom B, Tunell R. An evaluation of the efficiency of face masks in the resuscitation of newborn infants. *Lancet*. 1985;1:207–210
 52. O'Donnell CP, Davis PG, Lau R, Dargaville PA, Doyle LW, Morley CJ. Neonatal resuscitation 2: an evaluation of manual ventilation devices and face masks. *Arch Dis Child Fetal Neonatal Ed*. 2005;90:F392–F396
 53. Capasso L, Capasso A, Raimondi F, Vendemmia M, Araimo G, Paludetto R. A randomized trial comparing oxygen delivery on intermittent positive pressure with nasal cannulae versus facial mask in neonatal primary resuscitation. *Acta Paediatr*. 2005;94:197–200
 54. Wood FE, Morley CJ, Dawson JA, Kamlin CO, Owen LS, Donath S, Davis PG. Assessing the effectiveness of two round neonatal resuscitation masks: study 1. *Arch Dis Child Fetal Neonatal Ed*. 2008;93:F235–F237
 55. Bang AT, Bang RA, Baitule SB, Reddy HM, Deshmukh MD. Management of birth asphyxia in home deliveries in rural Gadchiroli: the effect of two types of birth attendants and of resuscitating with mouth-to-mouth, tube-mask or bag-mask. *J Perinatol*. 2005;25 suppl 1:S82–S91
 56. Massawe A, Kilewo C, Irani S, Verma RJ, Chakrapam AB, Ribbe T, Tunell R, Fischler B. Assessment of mouth-to-mask ventilation in resuscitation of asphyxial newborn babies. A pilot study. *Trop Med Int Health*. 1996;1:865–873
 57. Roberts RB, Day RL. Mouth-to-tube resuscitation of the neonate, II: the transmission of bacteria through endotracheal tubes and its prevention. *Anesth Analg*. 1973;52:242–245
 58. Milner AD, Stokes GM, Tunell R, McKeough M, Martin H. Laboratory assessment of Laerdal mouth tube mask prototype resuscitation device. *Med Biol Eng Comput*. 1992;30:117–119
 59. Terndrup TE, Kanter RK, Cherry RA. A comparison of infant ventilation methods performed by prehospital personnel. *Ann Emerg Med*. 1989;18:607–611
 60. Coffey PS, Kelly K, Tsu V. Preferences and practices: use of neonatal resuscitation devices in low-resource settings. *J Trop Pediatr*. 2007;53:415–419
 61. Polglase GR, Hillman NH, Pillow JJ, Cheah FC, Nitsos I, Moss TJ, Kramer BW, Ikegami M, Kallapur SG, Jobe AH. Positive end-expiratory pressure and tidal volume during initial ventilation of preterm lambs. *Pediatr Res*. 2008;64:517–522
 62. Probyn ME, Hooper SB, Dargaville PA, McCallion N, Harding R, Morley CJ. Effects of tidal volume and positive end-expiratory pressure during resuscitation of very premature lambs. *Acta Paediatr*. 2005;94:1764–1770
 63. Kattwinkel J, Stewart C, Walsh B, Gurka M, Paquet-Brown A. Responding to compliance changes in a lung model during manual ventilation: perhaps volume, rather than pressure, should be displayed. *Pediatrics*. 2009;123:e465–e470
 64. Resende JG, Zaconeta CA, Ferreira AC, Silva CA, Rodrigues MP, Rebello CM, Tavares P. Evaluation of peak inspiratory pressure, tidal volume and respiratory rate during ventilation of premature lambs using a self-inflating bag. *J Pediatr (Rio J)*. 2006;82:279–283
 65. Wood FE, Morley CJ, Dawson JA, Kamlin CO, Owen LS, Donath S, Davis PG. Improved techniques reduce face mask leak during simulated neonatal resuscitation: study 2. *Arch Dis Child Fetal Neonatal Ed*. 2008;93:F230–F234
 66. Hosono S, Inami I, Fujita H, Minato M, Takahashi S, Mugishima H. A role of end-tidal CO₂ monitoring for assessment of tracheal intubations in very low birth weight infants during neonatal resuscitation at birth. *J Perinat Med*. 2009;37:79–84
 67. Repetto JE, Donohue P-CP, Baker SF, Kelly L, Noguee LM. Use of capnography in the delivery room for assessment of endotracheal tube placement. *J Perinatol*. 2001;21:284–287
 68. Roberts WA, Maniscalco WM, Cohen AR, Litman RS, Chhibber A. The use of capnography for recognition of esophageal intubation in the neonatal intensive care unit. *Pediatr Pulmonol*. 1995;19:262–268
 69. Aziz HF, Martin JB, Moore JJ. The pediatric disposable end-tidal carbon dioxide detector role in endotracheal intubation in newborns. *J Perinatol*. 1999;19:110–113
 70. Garey DM, Ward R, Rich W, Heldt G, Leone T, Finer NN. Tidal volume threshold for colorimetric carbon dioxide detectors available for use in neonates. *Pediatrics*. 2008;121:e1524–e1527
 71. Hughes SM, Blake BL, Woods SL, Lehmann CU. False-positive results on colorimetric carbon dioxide analysis in neonatal resuscitation: potential for serious patient harm. *J Perinatol*. 2007;27:800–801
 72. Leone TA, Lange A, Rich W, Finer NN. Disposable colorimetric carbon dioxide detector use as an indicator of a patent airway during noninvasive mask ventilation. *Pediatrics*. 2006;118:e202–e204
 73. Finer NN, Rich W, Wang C, Leone T. Airway obstruction during mask ventilation of very low birth weight infants during neonatal resuscitation. *Pediatrics*. 2009;123:865–869
 74. Berg RA, Hilwig RW, Kern KB, Babar I, Ewy GA. Simulated mouth-to-mouth ventilation and chest compressions (bystander cardiopulmonary resuscitation) improves outcome in a swine model of prehospital pediatric asphyxial cardiac arrest. *Crit Care Med*. 1999;27:1893–1899
 75. Berg RA, Hilwig RW, Kern KB, Ewy GA. “Bystander” chest compressions and assisted ventilation independently improve outcome from piglet asphyxial pulseless “cardiac arrest.” *Circulation*. 2000;101:1743–1748
 76. Dean JM, Koehler RC, Schleiens CL, Atchison D, Gervais H, Berkowitz I, Traystman RJ. Improved blood flow during prolonged cardiopulmonary resuscitation with 30% duty cycle in infant pigs. *Circulation*. 1991;84:896–904
 77. Babbs CF, Nadkarni V. Optimizing chest compression to rescue ventilation ratios during one-rescuer CPR by professionals and lay persons: children are not just little adults. *Resuscitation*. 2004;61:173–181
 78. Srikantan SK, Berg RA, Cox T, Tice L, Nadkarni VM. Effect of one-rescuer compression/ventilation ratios on cardiopulmonary resuscitation in infant, pediatric, and adult manikins. *Pediatr Crit Care Med*. 2005;6:293–297
 79. Whyte SD, Sinha AK, Wyllie JP. Neonatal resuscitation—a practical assessment. *Resuscitation*. 1999;40:21–25
 80. Greingor JL. Quality of cardiac massage with ratio compression-ventilation 5/1 and 15/2. *Resuscitation*. 2002;55:263–267
 81. Wik L, Steen PA. The ventilation/compression ratio influences the effectiveness of two res-

- cuier advanced cardiac life support on a manikin. *Resuscitation*. 1996;31:113–119
82. Dorph E, Wik L, Steen PA. Effectiveness of ventilation-compression ratios 1:5 and 2:15 in simulated single rescuer paediatric resuscitation. *Resuscitation*. 2002;54:259–264
 83. Kinney SB, Tibballs J. An analysis of the efficacy of bag-valve-mask ventilation and chest compression during different compression-ventilation ratios in manikin-simulated paediatric resuscitation. *Resuscitation*. 2000;43:115–120
 84. Haque IU, Udassi JP, Udassi S, Theriaque DW, Shuster JJ, Zaritsky AL. Chest compression quality and rescuer fatigue with increased compression to ventilation ratio during single rescuer pediatric CPR. *Resuscitation*. 2008;79:82–89
 85. Kitamura T, Iwami T, Kawamura T, Nagao K, Tanaka H, Nadkarni VM, Berg RA, Hiraide A. Conventional and chest-compression-only cardiopulmonary resuscitation by bystanders for children who have out-of-hospital cardiac arrests: a prospective, nationwide, population-based cohort study. *Lancet*. 2010;375:1347–1354
 86. Houry PK, Frank LR, Menegazzi JJ, Taylor R. A randomized, controlled trial of two-thumb vs two-finger chest compression in a swine infant model of cardiac arrest. *Prehosp Emerg Care*. 1997;1:65–67
 87. Menegazzi JJ, Auble TE, Nicklas KA, Hosack GM, Rack L, Goode JS. Two-thumb versus two-finger chest compression during CPR in a swine infant model of cardiac arrest. *Ann Emerg Med*. 1993;22:240–243
 88. Udassi JP, Udassi S, Theriaque DW, Shuster JJ, Zaritsky AL, Haque IU. Effect of alternative chest compression techniques in infant and child on rescuer performance. *Pediatr Crit Care Med*. 2009;10:328–333
 89. David R. Closed chest cardiac massage in the newborn infant. *Pediatrics*. 1988;81:552–554
 90. Thaler MM, Stobie GH. An improved technique of external cardiac compression in infants and young children. *N Engl J Med*. 1963;269:606–610
 91. Whitelaw CC, Slywka B, Goldsmith LJ. Comparison of a two-finger versus two-thumb method for chest compressions by health-care providers in an infant mechanical model. *Resuscitation*. 2000;43:213–216
 92. Moya F, James LS, Burnard ED, Hanks EC. Cardiac massage in the newborn infant through the intact chest. *Am J Obstet Gynecol*. 1962;84:798–803
 93. Orłowski JP. Optimum position for external cardiac compression in infants and young children. *Ann Emerg Med*. 1986;15:667–673
 94. Phillips GW, Zideman DA. Relation of infant heart to sternum: its significance in cardiopulmonary resuscitation. *Lancet*. 1986;1:1024–1025
 95. Braga MS, Dominguez TE, Pollock AN, Niles D, Meyer A, Myklebust H, Nysaether J, Nadkarni V. Estimation of optimal CPR chest compression depth in children by using computer tomography. *Pediatrics*. 2009;124:e69–e74
 96. Jankov RP, Asztalos EV, Skidmore MB. Favourable neurological outcomes following delivery room cardiopulmonary resuscitation of infants < or =750 g at birth. *J Paediatr Child Health*. 2000;36:19–22
 97. O'Donnell AI, Gray PH, Rogers YM. Mortality and neurodevelopmental outcome for infants receiving adrenaline in neonatal resuscitation. *J Paediatr Child Health*. 1998;34:551–556
 98. Crespo SG, Schoffstall JM, Fuhs LR, Spivey WH. Comparison of two doses of endotracheal epinephrine in a cardiac arrest model. *Ann Emerg Med*. 1991;20:230–234
 99. Jasani MS, Nadkarni VM, Finkelstein MS, Mandell GA, Salzman SK, Norman ME. Effects of different techniques of endotracheal epinephrine administration in pediatric porcine hypoxic-hypercarbic cardiopulmonary arrest. *Crit Care Med*. 1994;22:1174–1180
 100. Mielke LL, Frank C, Lanzinger MJ, Wilhelm MG, Entholzner EK, Hargasser SR, Hipp RF. Plasma catecholamine levels following tracheal and intravenous epinephrine administration in swine. *Resuscitation*. 1998;36:187–192
 101. Roberts JR, Greenberg MI, Knaub MA, Kendrick ZV, Baskin SI. Blood levels following intravenous and endotracheal epinephrine administration. *JACEP*. 1979;8:53–56
 102. Hornchen U, Schuttler J, Stoeckel H, Eichelkraut W, Hahn N. Endobronchial instillation of epinephrine during cardiopulmonary resuscitation. *Crit Care Med*. 1987;15:1037–1039
 103. Guay J, Lortie L. An evaluation of pediatric in-hospital advanced life support interventions using the pediatric utstein guidelines: a review of 203 cardiorespiratory arrests. *Can J Anaesth*. 2004;51:373–378
 104. Perondi MB, Reis AG, Paiva EF, Nadkarni VM, Berg RA. A comparison of high-dose and standard-dose epinephrine in children with cardiac arrest. *N Engl J Med*. 2004;350:1722–1730
 105. Patterson MD, Boenning DA, Klein BL, Fuchs S, Smith KM, Hegenbarth MA, Carlson DW, Krug SE, Harris EM. The use of high-dose epinephrine for patients with out-of-hospital cardiopulmonary arrest refractory to prehospital interventions. *Pediatr Emerg Care*. 2005;21:227–237
 106. Goetting MG, Paradis NA. High-dose epinephrine improves outcome from pediatric cardiac arrest. *Ann Emerg Med*. 1991;20:22–26
 107. Vanduycke C, Martens P. High dose versus standard dose epinephrine in cardiac arrest—a meta-analysis. *Resuscitation*. 2000;45:161–166
 108. Berg RA, Otto CW, Kern KB, Hilwig RW, Sanders AB, Henry CP, Ewy GA. A randomized, blinded trial of high-dose epinephrine versus standard-dose epinephrine in a swine model of pediatric asphyxial cardiac arrest. *Crit Care Med*. 1996;24:1695–1700
 109. Burchfield DJ, Preziosi MP, Lucas VW, Fan J. Effects of graded doses of epinephrine during asphyxia-induced bradycardia in newborn lambs. *Resuscitation*. 1993;25:235–244
 110. Kirkman HN, Riley HD Jr. Posthemorrhagic anemia and shock in the newborn due to hemorrhage during delivery: report of 8 cases. *Pediatrics*. 1959;24:92–96
 111. Wyckoff MH, Perlman JM, Laptook AR. Use of volume expansion during delivery room resuscitation in near-term and term infants. *Pediatrics*. 2005;115:950–955
 112. Wyckoff M, Garcia D, Margraf L, Perlman J, Laptook A. Randomized trial of volume infusion during resuscitation of asphyxiated neonatal piglets. *Pediatr Res*. 2007;61:415–420
 113. Mayock DE, Gleason CA. Cerebrovascular effects of rapid volume expansion in preterm fetal sheep. *Pediatr Res*. 2004;55:395–399
 114. Box D, Cochran D. Safe reduction in administration of naloxone to newborn infants: an observational study. *Acta Paediatr*. 2006;95:1083–1086
 115. Bonta BW, Gagliardi JV, Williams V, Warshaw JB. Naloxone reversal of mild neurobehavioral depression in normal newborn infants after routine obstetric analgesia. *J Pediatr*. 1979;94:102–105
 116. Gill AW, Colvin J. Use of naloxone during neonatal resuscitation in Australia: compliance with published guidelines. *J Paediatr Child Health*. 2007;43:795–798
 117. Gibbs J, Newson T, Williams J, Davidson DC. Naloxone hazard in infant of opioid abuser. *Lancet*. 1989;2:159–160
 118. Van Woerkom R, Beharry KD, Modanlou HD, Parker J, Rajan V, Akmal Y, Aranda JV. Influence of morphine and naloxone on en-

- dothelin and its receptors in newborn piglet brain vascular endothelial cells: clinical implications in neonatal care. *Pediatr Res*. 2004;55:147–151
119. Ellemunter H, Simma B, Trawogger R, Maurer H. Intraosseous lines in preterm and full term neonates. *Arch Dis Child Fetal Neonatal Ed*. 1999;80:F74–F75
 120. Glaeser PW, Hellmich TR, Szweczuła D, Losek JD, Smith DS. Five-year experience in prehospital intraosseous infusions in children and adults. *Ann Emerg Med*. 1993;22:1119–1124
 121. Cramer K, Wiebe N, Hartling L, Crumley E, Vohra S. Heat loss prevention: a systematic review of occlusive skin wrap for premature neonates. *J Perinatol*. 2005;25:763–769
 122. Vohra S, Roberts RS, Zhang B, Janes M, Schmidt B. Heat loss prevention (help) in the delivery room: a randomized controlled trial of polyethylene occlusive skin wrapping in very preterm infants. *J Pediatr*. 2004;145:750–753
 123. Almeida PG, Chandley J, Davis J, Harrigan RC. Use of the heated gel mattress and its impact on admission temperature of very low birth-weight infants. *Adv Neonatal Care*. 2009;9:34–39
 124. Singh A, Duckett J, Newton T, Watkinson M. Improving neonatal unit admission temperatures in preterm babies: exothermic mattresses, polythene bags or a traditional approach? *J Perinatol*. 2010;30:45–49
 125. Knobel RB, Wimmer JE Jr, Holbert D. Heat loss prevention for preterm infants in the delivery room. *J Perinatol*. 2005;25:304–308
 126. Kent AL, Williams J. Increasing ambient operating theatre temperature and wrapping in polyethylene improves admission temperature in premature infants. *J Paediatr Child Health*. 2008;44:325–331
 127. Petrova A, Demissie K, Rhoads GG, Smulian JC, Marcella S, Ananth CV. Association of maternal fever during labor with neonatal and infant morbidity and mortality. *Obstet Gynecol*. 2001;98:20–27
 128. Lieberman E, Eichenwald E, Mathur G, Richardson D, Heffner L, Cohen A. Intrapartum fever and unexplained seizures in term infants. *Pediatrics*. 2000;106:983–988
 129. Shalak LF, Perlman JM, Jackson GL, Laptook AR. Depression at birth in term infants exposed to maternal chorioamnionitis: does neonatal fever play a role? *J Perinatol*. 2005;25:447–452
 130. Coimbra C, Boris-Moller F, Drake M, Wielech T. Diminished neuronal damage in the rat brain by late treatment with the antipyretic drug dipyrone or cooling following cerebral ischemia. *Acta Neuropathol*. 1996;92:447–453
 131. Goetzl L, Zigelboim I, Badell M, Rivers J, Mastrangelo MA, Tweardy D, Suresh MS. Maternal corticosteroids to prevent intrauterine exposure to hyperthermia and inflammation: a randomized, double-blind, placebo-controlled trial. *Am J Obstet Gynecol*. 2006;195:1031–1037
 132. Gluckman PD, Wyatt JS, Azzopardi D, Ballard R, Edwards AD, Ferriero DM, Polin RA, Robertson CM, Thoresen M, Whitelaw A, Gunn AJ. Selective head cooling with mild systemic hypothermia after neonatal encephalopathy: multicentre randomised trial. *Lancet*. 2005;365:663–670
 133. Shankaran S, Laptook AR, Ehrenkranz RA, Tyson JE, McDonald SA, Donovan EF, Fanaroff AA, Poole WK, Wright LL, Higgins RD, Finer NN, Carlo WA, Duara S, Oh W, Cotten CM, Stevenson DK, Stoll BJ, Lemons JA, Guillet R, Jobe AH. Whole-body hypothermia for neonates with hypoxic-ischemic encephalopathy. *N Engl J Med*. 2005;353:1574–1584
 134. Azzopardi DV, Strohm B, Edwards AD, Dyet L, Halliday HL, Juszczak E, Kapellou O, Levene M, Marlow N, Porter E, Thoresen M, Whitelaw A, Brocklehurst P. Moderate hypothermia to treat perinatal asphyxial encephalopathy. *N Engl J Med*. 2009;361:1349–1358
 135. Eicher DJ, Wagner CL, Katikaneni LP, Hulsey TC, Bass WT, Kaufman DA, Horgan MJ, Languani S, Bhatia JJ, Givelichian LM, Sankaran K, Yaeger JY. Moderate hypothermia in neonatal encephalopathy: efficacy outcomes. *Pediatr Neurol*. 2005;32:11–17
 136. Lin ZL, Yu HM, Lin J, Chen SQ, Liang ZQ, Zhang ZY. Mild hypothermia via selective head cooling as neuroprotective therapy in term neonates with perinatal asphyxia: an experience from a single neonatal intensive care unit. *J Perinatol*. 2006;26:180–184
 137. Edwards AD, Brocklehurst P, Gunn AJ, Halliday H, Juszczak E, Levene M, Strohm B, Thoresen M, Whitelaw A, Azzopardi D. Neurological outcomes at 18 months of age after moderate hypothermia for perinatal hypoxic ischaemic encephalopathy: synthesis and meta-analysis of trial data. *BMJ*. 2010;340:c363
 138. Jacobs S, Hunt R, Tarnow-Mordi W, Inder T, Davis P. Cooling for newborns with hypoxic ischaemic encephalopathy. *Cochrane Database Syst Rev*. 2007;CD003311
 139. Salhab WA, Wyckoff MH, Laptook AR, Perlman JM. Initial hypoglycemia and neonatal brain injury in term infants with severe fetal acidemia. *Pediatrics*. 2004;114:361–366
 140. Ondo-Onama C, Tumwine JK. Immediate outcome of babies with low Apgar score in Mulago hospital, Uganda. *East Afr Med J*. 2003;80:22–29
 141. Klein GW, Hojsak JM, Schmeidler J, Rapaport R. Hyperglycemia and outcome in the pediatric intensive care unit. *J Pediatr*. 2008;153:379–384
 142. LeBlanc MH, Huang M, Patel D, Smith EE, Devidas M. Glucose given after hypoxic ischemia does not affect brain injury in piglets. *Stroke*. 1994;25:1443–1447; discussion 1448
 143. Hattori H, Wasterlain CG. Posthypoxic glucose supplement reduces hypoxic-ischemic brain damage in the neonatal rat. *Ann Neurol*. 1990;28:122–128
 144. McDonald SJ, Middleton P. Effect of timing of umbilical cord clamping of term infants on maternal and neonatal outcomes. *Cochrane Database Syst Rev*. 2008;CD004074
 145. Rabe H, Reynolds G, Diaz-Rossello J. A systematic review and meta-analysis of a brief delay in clamping the umbilical cord of preterm infants. *Neonatology*. 2008;93:138–144
 146. Aladangady N, McHugh S, Aitchison TC, Wardrop CA, Holland BM. Infants' blood volume in a controlled trial of placental transfusion at preterm delivery. *Pediatrics*. 2006;117:93–98
 147. Mercer JS, Vohr BR, McGrath MM, Padbury JF, Wallach M, Oh W. Delayed cord clamping in very preterm infants reduces the incidence of intraventricular hemorrhage and late-onset sepsis: a randomized, controlled trial. *Pediatrics*. 2006;117:1235–1242
 148. De Leeuw R, Cuttini M, Nadai M, Berbić I, Hansen G, Kucinskas A, Lenoir S, Levin A, Persson J, Rebagliato M, Reid M, Schroell M, de Vonderweid U. Treatment choices for extremely preterm infants: an international perspective. *J Pediatr*. 2000;137:608–616
 149. Sanders MR, Donohue PK, Oberdorf MA, Rosenkrantz TS, Allen MC. Perceptions of the limit of viability: neonatologists' attitudes toward extremely preterm infants. *J Perinatol*. 1995;15:494–502
 150. Kopelman LM, Irons TG, Kopelman AE. Neonatologists judge the 'Baby Doe' regulations. *N Engl J Med*. 1988;318:677–683
 151. Costeloe K, Hennessy E, Gibson AT, Marlow N, Wilkinson AR. The epicure study: outcomes to discharge from hospital for infants born at the threshold of viability. *Pediatrics*. 2000;106:659–671

152. Field DJ, Dorling JS, Manktelow BN, Draper ES. Survival of extremely premature babies in a geographically defined population: prospective cohort study of 1994–9 compared with 2000–5. *BMJ*. 2008; 336:1221–1223
153. Tyson JE, Parikh NA, Langer J, Green C, Higgins RD. Intensive care for extreme prematurity—moving beyond gestational age. *N Engl J Med*. 2008;358:1672–1681
154. Paris JJ. What standards apply to resuscitation at the borderline of gestational age? *J Perinatol*. 2005;25:683–684
155. Casalaz DM, Marlow N, Speidel BD. Outcome of resuscitation following unexpected apparent stillbirth. *Arch Dis Child Fetal Neonatal Ed*. 1998;78:F112–F115
156. Jain L, Ferre C, Vidyasaagar D, Nath S, Sheftel D. Cardiopulmonary resuscitation of apparently stillborn infants: survival and long-term outcome. *J Pediatr*. 1991;118:778–782
157. Laptook AR, Shankaran S, Ambalavanan N, Carlo WA, McDonald SA, Higgins RD, Das A. Outcome of term infants using Apgar scores at 10 minutes following hypoxic-ischemic encephalopathy. *Pediatrics*. 2009; 124:1619–1626
158. Chamnanvanakij S, Perlman JM. Outcome following cardiopulmonary resuscitation in the neonate requiring ventilatory assistance. *Resuscitation*. 2000;45:173–180
159. Annibale DJ, Hulsey TC, Wagner CL, Southgate WM. Comparative neonatal morbidity of abdominal and vaginal deliveries after uncomplicated pregnancies. *Arch Pediatr Adolesc Med*. 1995;149:862–867
160. Atherton N, Parsons SJ, Mansfield P. Attendance of paediatricians at elective caesarean sections performed under regional anaesthesia: is it warranted? *J Paediatr Child Health*. 2006;42:332–336
161. Gordon A, McKechnie EJ, Jeffery H. Pediatric presence at cesarean section: justified or not? *Am J Obstet Gynecol*. 2005;193:599–605
162. Parsons SJ, Sonneveld S, Nolan T. Is a paediatrician needed at all caesarean sections? *J Paediatr Child Health*. 1998;34:241–244
163. Knudson MM, Khaw L, Bullard MK, Dicker R, Cohen MJ, Staudenmayer K, Sadjadi J, Howard S, Gaba D, Krummel T. Trauma training in simulation: translating skills from SIM time to real time. *J Trauma*. 2008; 64:255–263; discussion 263–254
164. Wayne DB, Didwania A, Feinglass J, Fudala MJ, Barsuk JH, McGaghie WC. Simulation-based education improves quality of care during cardiac arrest team responses at an academic teaching hospital: a case-control study. *Chest*. 2008;133:56–61
165. Schwid HA, Rooke GA, Michalowski P, Ross BK. Screen-based anesthesia simulation with debriefing improves performance in a mannequin-based anesthesia simulator. *Teach Learn Med*. 2001;13:92–96
166. Kory PD, Eisen LA, Adachi M, Ribaldo VA, Rosenthal ME, Mayo PH. Initial airway management skills of senior residents: simulation training compared with traditional training. *Chest*. 2007;132:1927–1931
167. Shapiro MJ, Morey JC, Small SD, Langford V, Kaylor CJ, Jagminas L, Suner S, Salisbury ML, Simon R, Jay GD. Simulation based teamwork training for emergency department staff: does it improve clinical team performance when added to an existing didactic teamwork curriculum? *Qual Saf Health Care*. 2004;13:417–421
168. Cherry RA, Williams J, George J, Ali J. The effectiveness of a human patient simulator in the ATLS shock skills station. *J Surg Res*. 2007;139:229–235
169. Savoldelli GL, Naik VN, Park J, Joo HS, Chow R, Hamstra SJ. Value of debriefing during simulated crisis management: oral versus video-assisted oral feedback. *Anesthesiology*. 2006;105:279–285
170. Blum RH, Raemer DB, Carroll JS, Dufresne RL, Cooper JB. A method for measuring the effectiveness of simulation-based team training for improving communication skills. *Anesth Analg*. 2005;100:1375–1380

DISCLOSURES

CoSTR Part 11: Writing Group Disclosures

Writing Group Member	Employment	Research Grant	Other Research Support	Speakers' Bureau/ Honoraria	Ownership Interest	Consultant/Advisory Board	Other
Jeffrey M. Perlman	Weill Cornell Medical College—Professor of Pediatrics	†NIH funding—Co-Investigator—Improving antimicrobial prescribing practices in the NICU	None	*University of Miami, and Cook County Chicago	None	None	None
Jonathan Wyllie	South Tees Foundation NHS Trust Health Service Provider NHS UK Consultant Neonatologist and Clinical Director of Neonatology	None	None	None	None	*Volunteer ICG Newborn Life Support ERC Volunteer author European Newborn Life Support Guidelines Volunteer author UK Newborn Resuscitation Guidelines Volunteer co-author Advanced Paediatric Life support Guidelines Volunteer member Advanced Life Support Group UK. Volunteer acting chair Newborn Life Support Working Group for RC(UK) Volunteer British Association of Perinatal Medicine Neonatal Services and staffing working group	None
John Kattwinkel	University of Virginia—Professor of Pediatrics	*American Academy of Pediatrics research grant to study resuscitators detection of compliance while administering positive pressure ventilation by resuscitation bag	None	None	None	None	*Curley vs Gordon et al, Boston, MA (still active)
Dianne L. Atkins	University of Iowa; Prof. †I am a compensated worksheet editor for AHA 2010 Guidelines process. The compensation is divided: 2/3 to my institution and 1/3 directly to me. The amount paid to my institution does not alter my salary	None	None	None	None	None	None
Leon Chameides	Emeritus Director, Pediatric Cardiology; Clinical Professor, University of Connecticut	None	None	None	None	None	None
Jay P. Goldsmith	Pediatrics Medical Group: Single specialty multi-site group practice—Neonatologist	None	None	None	None	None	None
Ruth Guinsburg	Federal University of São Paulo—Full Professor of Pediatrics	None	None	None	None	None	None
Mary Fran Hazinski	Vanderbilt University School of Nursing—Professor; AHA ECC Product Development—Senior Science Editor †the significant AHA compensation is designed to provide protected time for editing and writing responsibilities. I have a significant relationship with the AHA to support the mission of the AHA with the production of CoSTR and AHA Guidelines for CPR and ECC	None	None	None	None	None	None
Colin Morley	Retired Professor of Neonatal Medicine	None	None	Honoraria *Japanese Neonatal Society *Neonatal Ventilation and Resuscitation—Zagreb, Croatia *UK Middlesborough Neonatal Symposium	None	*Drager Medical about equip design *Education video on neonatal CPAP	None
Sam Richmond	UK National Health Service—Consultant Neonatologist	None	None	None	None	None	None
Wendy M. Simon	American Academy of Pediatrics—Director of Life Support Programs	None	None	None	None	None	None
Nalini Singhal	University of Calgary—Professor	*AAP grant looking at effect of PEEP with and with oxygen on resuscitation. Developing a International program for resuscitation, Helping Babies Breath	None	None	None	None	None
Edgardo Szylid	Fundasamin, Foundation for Women's and Infant Health—Executive Director of a non profit institution (NGO)	None	None	None	None	None	None

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CoSTR Part 11: Writing Group Disclosures, *Continued*

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This table represents the relationships of writing group members that may be perceived as actual or reasonably perceived conflicts of interest as reported on the Disclosure Questionnaire, which all members of the writing group are required to complete and submit. A relationship is considered to be "significant" if (a) the person receives \$10 000 or more during any 12-month period, or 5% or more of the person's gross income; or (b) the person owns 5% or more of the voting stock or share of the entity, or owns \$10 000 or more of the fair market value of the entity. A relationship is considered to be "modest" if it is less than "significant" under the preceding definition.

*Modest.

†Significant.

CoSTR Part 11: Worksheet Collaborator Disclosures

Worksheet Collaborator	Employment	Research Grant	Other Research Support	Speakers' Bureau/Honoraria	Ownership Interest	Consultant/Advisory Board	Other
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Steven Byrne	South Tees Hospital Foundation NHS Trust; National Health Service Trust	None	None	None	None	None	None
Peter Davis	The Royal Women's Hospital, Melbourne, Australia—Staff Neonatologist	None	None	None	None	None	None
William A. Engle	Indiana University School of Medicine Professor of Pediatrics	None	None	None	None	None	None
Marilyn Escobedo	University of Oklahoma—Professor of Pediatrics	None	None	None	None	None	None
Maria Fernanda de Almeida	Federal University of São Paulo: Full time work (40 h/week) at Neonatal Division—Department of Pediatrics—Assoc. Prof; Brazilian Pediatric Society: Voluntary work at Brazilian Neonatal Resuscitation Program—NRP Steering Committee—Co-chair	None	None	None	None	None	None
David Field	University of Leicester: Higher educational institution—UK Government funded—Professor of Neonatal Medicine	None	None	None	None	None	None
Judith Finn	University of Western Australia—Professor	†Multiple National Health and Medical Research Grants (NH & MRC), National Heart Foundation Australia and State Government grants of >\$10 000 since 1999. A—No money came to me—all came to my University to employ research staff and meet research expenses. No grants were directly related to any topic on which I am undertaking a Worksheet and none involved the trialing of a commercial product	None	*Less than \$1000 from the Japanese Resuscitation Council to speak at their JRC Conference in Osaka in 2009	None	None	None

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CoSTR Part 11: Worksheet Collaborator Disclosures, *Continued*

Worksheet Collaborator	Employment	Research Grant	Other Research Support	Speakers' Bureau/Honoraria	Ownership Interest	Consultant/Advisory Board	Other
Louis P. Halamek	Stanford University—Associate Professor	†Source: Laerdal Foundation I was the Principal Investigator on 3 year grant (2006–2009) funded by the Laerdal Foundation in the amount of \$450 000 USD that was given to the simulation center I direct, the Center for Advanced Pediatric and Perinatal Education at Packard Children's Hospital at Stanford. This grant ended 2009-07-31. This grant was not a source of support for my salary	None	None	None	*I provide consultation services regarding simulation product design and function to Laerdal Medical, Inc., and Advanced Medical Simulation, Inc.	*I provide medicolegal consultation services to trial attorneys in the U.S., advising on questions relating to neonatal intensive care
Jane E. McGowan	St Christopher's Pediatric Associates: Practice Group for Children's Hospital—part of Tenet Healthcare—Attending Neonatologist	None	None	*Received honorarium for giving at talk for the March of Dimes on "NRP and the Preterm Infant"	None	None	None
Douglas D. McMillan	Dalhousie University and Academic Pediatrics Incorporated: University and Department of Pediatrics Financial group—Professor and Head, Division of Neonatal Perinatal medicine	None	None	†Medical Legal consulting fees for different hospitals and the Canadian Medical Protective Association (occasionally for the plaintiff)—presently "remitted" to Academic Pediatrics Incorporated *Consulting fees for program reviews related to newborn care and associated education programs—presently "remitted" to Academic Pediatrics Incorporated	None	*Consulting fees for program reviews related to newborn care and associated education programs—presently "remitted" to Academic Pediatrics Incorporated	*Medical Legal consulting fees for different hospitals and the Canadian Medical Protective Association (occasionally for the plaintiff)—presently "remitted" to Academic Pediatrics Incorporated
Lindsay Mildenhall	Counties Manukau District Health Board Auckland New Zealand: Public Health Care Provider—Consultant Neonatologist	None	None	None	None	None	None
Rintaro Mori	Osaka Medical Center and Research Institute for Maternal and Child Health: a public children's hosp. run by a local government—Division Director of Strategic Planning & Collaboration	None	None	None	None	None	None
Susan Niermeyer	University of Colorado Denver School of Medicine: professor, clinical neonatologist—Professor of Pediatrics	†Editorship, Helping Babies Breathe, American Academy of Pediatrics 2008–2009 Salary support through contract with University of Colorado Denver School of Medicine	None	None	None	None	None
Colm O'Donnell	The National Maternity Hospital, Holles Street, Dublin 2, Ireland—Consultant Neonatologist; Our Lady's Children's Hospital, Crumlin, Dublin 12, Ireland—Consultant Neonatologist; University College Dublin, Ireland: University medical school—Clinical Lecturer	None	None	*I have received honoraria from Chiesi Pharma (makers of Curosurf) for speaking at 2 educational courses and 3 scientific meetings (ie. on 5 occasions) in the last 2 years. The combined total of these honoraria is less than 1000 euros	None	None	None
Yacov Rabi	Alberta Health Services: Provide employment income for my role as a neonatologist—Physician; University of Calgary: Provides income for my role as an Assistant Professor of Medicine	None	*Supply of modified Neopuff circuits for a randomized control trial by Fisher Paykell	None	None	None	None
Steven A. Ringer	Brigham and Women's Hospital: Non-profit Hospital—Chief, Newborn Medicine	None	None	*Vermont Oxford Neonatal Network Annual Meeting	None	*Alere Healthcare Advisory Board Consulting on Clinical Care guidelines. Nothing relevant to topics with which I am involved	†Expert Witness in medical legal proceedings (Malpractice cases). A number of different attorneys/ insurance companies Money comes directly to me. Nothing relevant to questions under consideration
Jasmeet Soar	North Bristol NHS Trust: Government Hospital in UK—Consultant in Anesthetics & Intensive Care Medicine	None	None	None	None	None	None

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CoSTR Part 11: Worksheet Collaborator Disclosures, *Continued*

Worksheet Collaborator	Employment	Research Grant	Other Research Support	Speakers' Bureau/Honoraria	Ownership Interest	Consultant/Advisory Board	Other
Benjamin J. Stenson	United Kingdom Public Health Service—Consultant Neonatologist	None	None	None	None	None	None
Enrique Udaeta	Medica Sur Lomas: Private Maternity Hospital—Director Department of Neonatology	None	None	None	None	None	None
Dharmapuri Vidyasagar	University of Illinois Professor emeritus Professor emeritus	None	None	None	None	None	None
Michael Watkinson	NHS Heart of England Foundation Trust, Birmingham, UK This is an NHS hospital in England Consultant neonatologist	None	None	None	None	None	None
Gary M. Weiner	St. Joseph Mercy Hospital—Attending Neonatologist	None	†Received equipment on-loan (3 resuscitation mannequins, 2 sets of video recording equipment) from Laerdal Medical Corporation to be used to complete a research project evaluating educational methods for teaching neonatal resuscitation. The value of the on-loan equipment is approximately \$35 000	None	None	None	None
Myra H. Wyckoff	UT Southwestern Medical Center at Dallas—Associate Professor of Pediatrics	†PI, American Academy of Pediatrics. Neonatal Resuscitation Program. The ergonomics of neonatal cardiac compressions. \$71 030 January 2008–2009 The funding comes to the institution *Co-Investigator (Mentor), American Academy of Pediatrics Neonatal Resuscitation Program Young Investigator Grant. Effectiveness of Plastic Head Coverings for Hypothermia Prevention in Preterm Newborns. January 2009–January 2010, \$10 000 The funding comes to the institution	None	Feb 5, 2009 Pediatric Grand Rounds. University of Oklahoma Health Sciences Center. OKC, OK	None	None	None

This table represents the relationships of worksheet collaborators members that may be perceived as actual or reasonably perceived conflicts of interest as reported on the Disclosure Questionnaire, which all worksheet collaborators are required to complete and submit. A relationship is considered to be "significant" if (a) the person receives \$10 000 or more during any 12-month period, or 5% or more of the person's gross income; or (b) the person owns 5% or more of the voting stock or share of the entity, or owns \$10 000 or more of the fair market value of the entity. A relationship is considered to be "modest" if it is less than "significant" under the preceding definition.

*Modest.

†Significant.

APPENDIX

CoSTR Part 11: Worksheet Appendix

Task Force	WS ID	PICO Title	Short Title	Authors	URL
EIT	EIT-001A	For resuscitation teams (P), do briefings/debriefings (I), when compared to no briefings/debriefings (C), improve performance or outcomes (O)? (INTERVENTION)	Debriefing of CPR performance	Dana P. Edelson, Trevor Yuen	http://circ.ahajournals.org/site/c2010/eit-001a.pdf
EIT	EIT-001B	For resuscitation teams (P), do briefings/debriefings (I), when compared to no briefings/debriefings (C), improve performance or outcomes (O)? (INTERVENTION)	Debriefing of CPR performance	Jasmeet Soar	http://circ.ahajournals.org/site/c2010/eit-001b.pdf

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CoSTR Part 11: Worksheet Appendix, *Continued*

Task Force	WS ID	PICO Title	Short Title	Authors	URL
EIT	EIT-019A	In participants undergoing BLS/ALS courses (P), does the inclusion of more realistic techniques (eg. high fidelity manikins, in-situ training) (I), as opposed to standard training (eg. low fidelity, education centre) (C), improve outcomes (eg. skills performance on manikins, skills performance in real arrests, willingness to perform etc.) (O)?	High fidelity training	Jordan Duval- Arnould, Elizabeth A. Hunt	http://circ.ahajournals.org/site/c2010/eit-019a.pdf
EIT	EIT-019B	In participants undergoing BLS/ALS courses (P), does the inclusion of more realistic techniques (eg. high fidelity manikins, in-situ training) (I), as opposed to standard training (eg. low fidelity, education centre) (C), improve outcomes (eg. skills performance on manikins, skills performance in real arrests, willingness to perform etc.) (O)?	High fidelity training	Judith Finn	http://circ.ahajournals.org/site/c2010/eit-019b.pdf
NRP	NRP-001A	For neonates requiring resuscitation (P), is any adjunct measure (eg. CO ₂ detection, pulse oximeter) as effective as the usual clinical findings (eg. heart rate, chest movement) effective to improve outcome (O)?	Adjuncts: CO ₂ detection, pulse oximeter	John Kattwinkel	http://circ.ahajournals.org/site/c2010/nrp-001a.pdf
NRP	NRP-001B	For neonates requiring resuscitation (P), is any adjunct measure (eg. CO ₂ detection, pulse oximeter) as effective as the usual clinical findings (eg. heart rate, chest movement) effective to improve outcome (O)?	Adjuncts: CO ₂ detection, pulse oximeter	Yacov Rabi	http://circ.ahajournals.org/site/c2010/nrp-001b.pdf
NRP	NRP-002A	In the neonates infant (preterm and term) receiving respiratory support (P), does the use of CPAP(I) versus no-CPAP or IPPV(C) improve outcome—specify (O)?	CPAP and IPPV	Colm O'Donnell	http://circ.ahajournals.org/site/c2010/nrp-002a.pdf
NRP	NRP-002B	In the neonates infant (preterm and term) receiving respiratory support (P), does the use of CPAP(I) versus no-CPAP or IPPV(C) improve outcome—specify (O)?	CPAP and IPPV	Douglas D. McMillan	http://circ.ahajournals.org/site/c2010/nrp-002b.pdf
NRP	NRP-003A	In neonates receiving respiratory support (P) does the use of face mask interface (I) versus CPAP, NPCPAP, NC (C) (excluding intubation improve outcome) (O)?	Face mask interface vs CPAP etc	Golin Morley	http://circ.ahajournals.org/site/c2010/nrp-003a.pdf
NRP	NRP-003B	In neonates receiving respiratory support (P) does the use of face mask interface (I) versus CPAP, NPCPAP, NC (C) (excluding intubation) improve outcome (O)?	Face mask interface vs CPAP etc	Yacov Rabi	http://circ.ahajournals.org/site/c2010/nrp-003b.pdf
NRP	NRP-004A	In neonates receiving resuscitation (P) does the use of mouth-to-mouth, mouth-to-mask, mouth tube to mask (I) as compared to a self-inflating bag (C) give equivalent outcomes (stable spontaneous breathing) (O), when devices for delivering PPV are not available?	Self-inflating bag vs mouth techniques	Nalini Singhal	http://circ.ahajournals.org/site/c2010/nrp-004a.pdf
NRP	NRP-004B	In neonates receiving resuscitation (P) does the use of mouth-to-mouth, mouth-to-mask, mouth tube to mask (I) as compared to a self-inflating bag (C) give equivalent outcomes (stable spontaneous breathing) (O), when devices for delivering PPV are not available?	Self-inflating bag vs mouth techniques	Maria Fernanda de Almeida	http://circ.ahajournals.org/site/c2010/nrp-004b.pdf
NRP	NRP-005A	In neonates receiving positive pressure ventilation (P) does the use of gas volume monitoring (I) versus clinical assessment with or without pressure monitoring (C) improve clinical outcome (O)?	Ventilation volume monitoring	Steven A. Ringer	http://circ.ahajournals.org/site/c2010/nrp-005a.pdf
NRP	NRP-005B	In neonates receiving positive pressure ventilation (P) does the use of gas volume monitoring (I) versus clinical assessment with or without pressure monitoring (C) improve clinical outcome (O)?	Ventilation volume monitoring	Khalid Aziz	http://circ.ahajournals.org/site/c2010/nrp-005b.pdf
NRP	NRP-005C	In neonates receiving positive pressure ventilation (P) does the use of gas volume monitoring (I) versus clinical assessment with or without pressure monitoring (C) improve clinical outcome (O)?	Ventilation volume monitoring	Jane E. McGowan	http://circ.ahajournals.org/site/c2010/nrp-005c.pdf
NRP	NRP-006A	In neonates receiving chest compressions (P) do other ratios (5:1, 15:2) (I) versus a 3:1 (C) improve outcomes (O)?	Compression ventilation ratio	Lindsay Mildenhall	http://circ.ahajournals.org/site/c2010/nrp-006a.pdf
NRP	NRP-006B	In neonates receiving chest compressions (P) do other ratios (5:1, 15:2) (I) versus a 3:1 (C) improve outcomes (O)?	Compression ventilation ratio	Myra H. Wyckoff	http://circ.ahajournals.org/site/c2010/nrp-006b.pdf

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CoSTR Part 11: Worksheet Appendix, *Continued*

Task Force	WS ID	PICO Title	Short Title	Authors	URL
NRP	NRP-007A	In neonates (P) receiving chest compressions does the two thumb (I) versus two finger (C) method of administration improve outcome (O)?	Two thumb vs two finger	Lindsay Mildenhall	http://circ.ahajournals.org/site/c2010/nrp-007a.pdf
NRP	NRP-007B	In neonates (P) receiving chest compressions does the two thumb (I) versus two finger (C) method of administration improve outcome (O)?	Two thumb vs two finger	Myra H. Wyckoff	http://circ.ahajournals.org/site/c2010/nrp-007b.pdf
NRP	NRP-008A	Among neonates (≤ 28 days) with a HR < 60 bpm despite adequate ventilation and chest compressions, does the IV route compared with the ET route of epinephrine administration: 1. Increase heart rate > 100 bpm faster, 2. Increase ROSC, or 3. Increase survival to discharge?	IV vs ET epinephrine	Jonathan Wyllie	http://circ.ahajournals.org/site/c2010/nrp-008a.pdf
NRP	NRP-008B	Among neonates (≤ 28 days) with a HR < 60 bpm despite adequate ventilation and chest compressions, does the IV route compared with the ET route of epinephrine administration: 1. Increase heart rate > 100 bpm faster, 2. Increase ROSC, or 3. Increase survival to discharge?	IV vs ET epinephrine	Gary M. Weiner	http://circ.ahajournals.org/site/c2010/nrp-008b.pdf
NRP	NRP-009A	Among neonates (≤ 28 days) with HR < 60 bpm does HDE (IV > 0.03 mg/kg or ET > 0.1 mg/kg) compared with SDE: 1. Increase HR > 100 bpm faster 2. Increase ROSC, or 3. Increase survival to discharge?	Epinephrine dose	Jonathan Wyllie	http://circ.ahajournals.org/site/c2010/nrp-009a.pdf
NRP	NRP-009B	Among neonates (≤ 28 days) with HR < 60 bpm does HDE (IV > 0.03 mg/kg or ET > 0.1 mg/kg) compared with SDE: 1. Increase HR > 100 bpm faster 2. Increase ROSC, or 3. Increase survival to discharge?	Epinephrine dose	Gary M. Weiner	http://circ.ahajournals.org/site/c2010/nrp-009b.pdf
NRP	NRP-010A	For infants delivered at ≥ 34 weeks gestation (P), is delivery by elective c-section under regional anesthesia (I) in comparison with unassisted vertex vaginal deliveries (C) associated with an increased risk of requirement for intubation during resuscitation (O)?	Prenatal prediction of respiratory compromise	Marilyn B. Escobedo	http://circ.ahajournals.org/site/c2010/nrp-010a.pdf
NRP	NRP-010B	For infants delivered at ≥ 34 weeks gestation (P), is delivery by elective c-section under regional anesthesia (I) in comparison with unassisted vertex vaginal deliveries (C) associated with an increased risk of requirement for intubation during resuscitation (O)?	Prenatal prediction of respiratory compromise	Benjamin J. Stenson	http://circ.ahajournals.org/site/c2010/nrp-010b.pdf
NRP	NRP-010C	For infants delivered at ≥ 34 weeks gestation (P), is delivery by elective c-section under regional anesthesia (I) in comparison with unassisted vertex vaginal deliveries (C) associated with an increased risk of requirement for intubation during resuscitation (O)?	Prenatal prediction of respiratory compromise	Dianne L. Atkins, Edgardo Szlyd	http://circ.ahajournals.org/site/c2010/nrp-010c.pdf
NRP	NRP-011A	In depressed neonates with clear amniotic fluid (P) does suctioning of the mouth and nose (I) versus none (C) improve outcome (O).	Clear amniotic fluid	Sithembiso Velaphi, Dharmapuri Vidyasagar	http://circ.ahajournals.org/site/c2010/nrp-011a.pdf
NRP	NRP-012A	In depressed neonates born through meconium stained amniotic fluid (P), does endotracheal suctioning (I) versus no suctioning (C) improve outcome (O)?	Stained amniotic fluid	Sithembiso Velaphi, Dharmapuri Vidyasagar	http://circ.ahajournals.org/site/c2010/nrp-012a.pdf
NRP	NRP-013A	When resuscitating or stabilizing newborns at birth (P), is there an oxygen administration strategy (I) that is superior to any other (C) in improving outcome (O)?	Oxygen administration	Jay Goldsmith	http://circ.ahajournals.org/site/c2010/nrp-013a.pdf
NRP	NRP-013B	When resuscitating or stabilizing newborns at birth (P), is there an oxygen administration strategy (I) that is superior to any other (C) in improving outcome (O)?	Oxygen administration	Sam Richmond	http://circ.ahajournals.org/site/c2010/nrp-013b.pdf
NRP	NRP-014A	In neonates receiving resuscitation or stabilization (P), is the saturation demonstrated during normal birth (I) preferable to some other target (C), when considering outcome for premature and term neonates (O)?	Oxygen saturation target	John Kattwinkel	http://circ.ahajournals.org/site/c2010/nrp-014a.pdf
NRP	NRP-014B	In neonates receiving resuscitation or stabilization (P), is the saturation demonstrated during normal birth (I) preferable to some other target (C), when considering outcome for premature and term neonates (O)?	Oxygen saturation target	Colin Morley	http://circ.ahajournals.org/site/c2010/nrp-014b.pdf

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CoSTR Part 11: Worksheet Appendix, *Continued*

Task Force	WS ID	PICO Title	Short Title	Authors	URL
NRP	NRP-015A	In neonates (P) receiving positive pressure during resuscitation, is positive pressure ventilation by T-piece resuscitator (I) superior to bag ventilation (C) for improving outcome—specify (O)?	T-piece resuscitator	David Boyle	http://circ.ahajournals.org/site/c2010/nrp-015a.pdf
NRP	NRP-015B	In neonates (P) receiving positive pressure during resuscitation, is positive pressure ventilation by T-piece resuscitator (I) superior to bag ventilation (C) for improving outcome—specify (O)?	T-piece resuscitator	Benjamin J. Stenson	http://circ.ahajournals.org/site/c2010/nrp-015b.pdf
NRP	NRP-015C	In neonates (P) receiving positive pressure during resuscitation, is positive pressure ventilation by T-piece resuscitator (I) superior to bag ventilation (C) for improving outcome—specify (O)?	T-piece resuscitator	David Field	http://circ.ahajournals.org/site/c2010/nrp-015c.pdf
NRP	NRP-016A	For neonates (P) following attempted endotracheal intubation, is CO ₂ detection (I) superior to clinical assessment (C) for confirming endotracheal location (O)?	CO ₂ detection	Jonathan Wyllie	http://circ.ahajournals.org/site/c2010/nrp-016a.pdf
NRP	NRP-017A	For neonates requiring positive pressure ventilation (P), is LMA (I) an effective alternative to mask or endotracheal ventilation (C) for improving outcome (O)? (achieving stable vital signs and reducing the need for subsequent endotracheal intubation)?	LMA	Gary M. Weiner	http://circ.ahajournals.org/site/c2010/nrp-017a.pdf
NRP	NRP-017B	For neonates requiring positive pressure ventilation (P), is LMA (I) an effective alternative to mask or endotracheal ventilation (C) for improving outcome (O)? (achieving stable vital signs and reducing the need for subsequent endotracheal intubation)?	LMA	Enrique Udaeta	http://circ.ahajournals.org/site/c2010/nrp-017b.pdf
NRP	NRP-018A	For non intubated bradycardic neonates (P) requiring positive pressure ventilation, is the CO ₂ monitoring device (I) more effective than chest rise, color (C) for assessing adequate ventilation (O)?	Bradycardia and CO ₂ monitoring	Colm O'Donnell	http://circ.ahajournals.org/site/c2010/nrp-018a.pdf
NRP	NRP-018B	For non intubated bradycardic neonates (P) requiring positive pressure ventilation, is the CO ₂ monitoring device (I) more effective than chest rise, color (C) for assessing adequate ventilation (O)?	Bradycardia and CO ₂ monitoring	Masanori Tamura	http://circ.ahajournals.org/site/c2010/nrp-018b.pdf
NRP	NRP-018C	For non intubated bradycardic neonates (P) requiring positive pressure ventilation, is the CO ₂ monitoring device (I) more effective than chest rise, color (C) for assessing adequate ventilation (O)?	Bradycardia and CO ₂ monitoring	Steven A. Ringer	http://circ.ahajournals.org/site/c2010/nrp-018c.pdf
NRP	NRP-019A	In neonates requiring resuscitation, (P) will the early use of supplemental glucose (I) during and/or following delivery room resuscitation, versus none (C) improve outcome (i.e. avoidance of hypoglycemia, reduced long-term neurologic morbidity) (O)?	Supplemental glucose	Jane E. McGowan	http://circ.ahajournals.org/site/c2010/nrp-019a.pdf
NRP	NRP-019B	In neonates requiring resuscitation, (P) will the early use of supplemental glucose (I) during and/or following delivery room resuscitation, versus none (C) improve outcome (i.e. avoidance of hypoglycemia, reduced long-term neurologic morbidity) (O)?	Supplemental glucose	Jeffrey Perlman	http://circ.ahajournals.org/site/c2010/nrp-019b.pdf
NRP	NRP-020A	In neonates requiring resuscitation, does the administration of emergency medications (P) by intraosseous infusion (I) versus the intravenous route improve outcome (O)?	I0 vs IV	William A. Engle	http://circ.ahajournals.org/site/c2010/nrp-020a.pdf
NRP	NRP-021A	In neonates requiring resuscitation and not responding to CPR (P), does the administration of sodium bicarbonate (I) versus no bicarbonate (C) improve outcome (O)?	Sodium bicarbonate	Jeffrey Perlman	http://circ.ahajournals.org/site/c2010/nrp-021a.pdf
NRP	NRP-021B	In neonates requiring resuscitation and not responding to CPR (P), does the administration of sodium bicarbonate (I) versus no bicarbonate (C) improve outcome (O)?	Sodium bicarbonate	Dianne L. Atkins, Sam Richmond	http://circ.ahajournals.org/site/c2010/nrp-021b.pdf
NRP	NRP-022A	In apneic neonates suspected of narcotic depression (P), does naloxone (I) when compared to effective ventilation without naloxone (C) improve outcome (O)?	Nalaxone	Ruth Guinsburg	http://circ.ahajournals.org/site/c2010/nrp-022a.pdf
NRP	NRP-022B	In apneic neonates suspected of narcotic depression (P), does naloxone (I) when compared to effective ventilation without naloxone (C) improve outcome (O)?	Nalaxone	Myra H. Wyckoff	http://circ.ahajournals.org/site/c2010/nrp-022b.pdf

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CoSTR Part 11: Worksheet Appendix, *Continued*

Task Force	WS ID	PICO Title	Short Title	Authors	URL
NRP	NRP-023A	In preterm neonates under radiant warmers (P), does increased room temperature, thermal mattress, or other intervention (I) as compared to plastic wraps alone (C) improve outcome (O)?	Warming adjuncts	Marilyn B. Escobedo, Michael Watkinson	http://circ.ahajournals.org/site/c2010/nrp-023a.pdf
NRP	NRP-024A	In term neonates at risk for hypoxic-ischemic encephalopathy secondary to intra-partum hypoxia (P) does selective/whole body cooling (I) versus standard therapy (C), result in improved outcome (O)?	Hypothermia (induced)	Jeffrey Perlman	http://circ.ahajournals.org/site/c2010/nrp-024a.pdf
NRP	NRP-024B	In term neonates at risk for hypoxic-ischemic encephalopathy secondary to intra-partum hypoxia (P) does selective/whole body cooling (I) versus standard therapy (C), result in improved outcome (O)?	Hypothermia (induced)	Peter Davis	http://circ.ahajournals.org/site/c2010/nrp-024b.pdf
NRP	NRP-025A	In term neonates without a detectable heart rate and no other signs of life (P) is ten minutes (I) as opposed to 15 minutes or longer (C) of effective resuscitation a reliable measure of outcome (abnormal neurologic examination and/or death) (O)?	Duration of CPR with asystole and outcome	Steve Byrne	http://circ.ahajournals.org/site/c2010/nrp-025a.pdf
NRP	NRP-025B	In term neonates without a detectable heart rate and no other signs of life (P) is ten minutes (I) as opposed to 15 minutes or longer (C) of effective resuscitation a reliable measure of outcome (abnormal neurologic examination and/or death) (O)?	Duration of CPR with asystole and outcome	Jay Goldsmith	http://circ.ahajournals.org/site/c2010/nrp-025b.pdf
NRP	NRP-025C	In term neonates without a detectable heart rate and no other signs of life (P) is ten minutes (I) as opposed to 15 minutes or longer (C) of effective resuscitation a reliable measure of outcome (abnormal neurologic examination and/or death) (O)?	Duration of CPR with asystole and outcome	Ruth Guinsburg	http://circ.ahajournals.org/site/c2010/nrp-025c.pdf
NRP	NRP-026A	In term neonates with a heart rate <60 and no other signs of life (P), is ten minutes (I) as opposed to 15 minutes or longer (C) of effective resuscitation a reliable measure of outcome (abnormal neurologic examination and/or death) (O)?	Duration of CPR with bradycardia and outcome	Steve Byrne	http://circ.ahajournals.org/site/c2010/nrp-026a.pdf
NRP	NRP-026B	In term neonates with a heart rate <60 and no other signs of life (P), is ten minutes (I) as opposed to 15 minutes or longer (C) of effective resuscitation a reliable measure of outcome (abnormal neurologic examination and/or death) (O)?	Duration of CPR with bradycardia and outcome	Jay Goldsmith	http://circ.ahajournals.org/site/c2010/nrp-026b.pdf
NRP	NRP-026C	In term neonates with a heart rate <60 and no other signs of life (P), is ten minutes (I) as opposed to 15 minutes or longer (C) of effective resuscitation a reliable measure of outcome (abnormal neurologic examination and/or death) (O)?	Duration of CPR with bradycardia and outcome	Ruth Guinsburg	http://circ.ahajournals.org/site/c2010/nrp-026c.pdf
NRP	NRP-027A	In neonates at the limits of viability or anomalies associated with lethal outcomes (P) does the non initiation (I) versus initiation (C) of resuscitation result in an outcome that is ethically justified (O).	Futile resuscitation rules	Steve Byrne	http://circ.ahajournals.org/site/c2010/nrp-027a.pdf
NRP	NRP-027B	In neonates at the limits of viability or anomalies associated with lethal outcomes (P) does the non initiation (I) versus initiation (C) of resuscitation result in an outcome that is ethically justified (O).	Futile resuscitation rules	Jay Goldsmith	http://circ.ahajournals.org/site/c2010/nrp-027b.pdf
NRP	NRP-028A	In depressed neonates requiring positive pressure ventilation (P) does the administration of longer inspiratory times, higher inflation pressures, use of PEEP (I) as compared to standard management (C) improve outcome (O)?	Ventilation times and pressures	David Boyle	http://circ.ahajournals.org/site/c2010/nrp-028a.pdf
NRP	NRP-028B	In depressed neonates requiring positive pressure ventilation (P) does the administration of longer inspiratory times, higher inflation pressures, use of PEEP (I) as compared to standard management (C) improve outcome (O)?	Ventilation times and pressures	Benjamin J. Stenson	http://circ.ahajournals.org/site/c2010/nrp-028b.pdf
NRP	NRP-029A	In neonates requiring resuscitation and unresponsive to chest compressions/epinephrine (P) does the administration of volume (I) versus no volume (C) improve outcome (O).	Volume resuscitation with CPR	Susan Niermeyer	http://circ.ahajournals.org/site/c2010/nrp-029a.pdf
NRP	NRP-029B	In neonates requiring resuscitation and unresponsive to chest compressions/epinephrine (P) does the administration of volume (I) versus no volume (C) improve outcome (O).	Volume resuscitation with CPR	Douglas D. McMillan	http://circ.ahajournals.org/site/c2010/nrp-029b.pdf

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CoSTR Part 11: Worksheet Appendix, *Continued*

Task Force	WS ID	PICO Title	Short Title	Authors	URL
NRP	NRP-029C	In neonates requiring resuscitation and unresponsive to chest compressions/epinephrine (P) does the administration of volume (I) versus no volume (C) improve outcome (O).	Volume resuscitation with CPR	Masanori Tamura	http://circ.ahajournals.org/site/c2010/nrp-029c.pdf
NRP	NRP-030A	In neonates (P), does delayed cord clamping cord or milking of the cord (I) versus standard management (C), improve outcome (O).	Umbilical cord clamping and milking	Susan Niermeyer	http://circ.ahajournals.org/site/c2010/nrp-030a.pdf
NRP	NRP-030B	In neonates (P), does delayed cord clamping cord or milking of the cord (I) versus standard management (C), improve outcome (O).	Umbilical cord clamping and milking	Dianne L. Atkins, Nalini Singhal	http://circ.ahajournals.org/site/c2010/nrp-030b.pdf
NRP	NRP-030C	In neonates (P), does delayed cord clamping cord or milking of the cord (I) versus standard management (C), improve outcome (O) (milking of the cord).	Umbilical cord clamping and milking	Gary M. Weiner	http://circ.ahajournals.org/site/c2010/nrp-030c.pdf
NRP	NRP-030D	In neonates (P), does delayed cord clamping cord or milking of the cord (I) versus standard management (C), improve outcome (O)?	Umbilical cord clamping and milking	Rintaro Mori	http://circ.ahajournals.org/site/c2010/nrp-030d.pdf
NRP	NRP-031A	In neonates born to febrile mothers (P) does intervention to normalize temperature (I), compared to standard care (C) improve outcome (O)?	Maternal fever	Jeffrey Perlman	http://circ.ahajournals.org/site/c2010/nrp-031a.pdf
NRP	NRP-031B	In neonates born to febrile mothers (P) does intervention to normalize temperature (I), compared to standard care (C) improve outcome (O).	Maternal fever	Steven A. Ringer	http://circ.ahajournals.org/site/c2010/nrp-031b.pdf
NRP	NRP-032A	In participants undergoing resuscitation courses (P), does the inclusion of more realistic techniques (eg. high fidelity manikins, in-situ training) (I), as opposed to standard training (eg. low fidelity, education centre) (C), improve outcomes (eg. skills performance) (O).	Impact of realistic training on skills performance	Jane E. McGowan	http://circ.ahajournals.org/site/c2010/nrp-032a.pdf
NRP	NRP-032B	In participants undergoing resuscitation courses (P), does the inclusion of more realistic techniques (eg. high fidelity manikins, in-situ training) (I), as opposed to standard training (eg. low fidelity, education centre) (C), improve outcomes (eg. skills performance) (O).	Impact of realistic training on skills performance	Louis P. Halamek	http://circ.ahajournals.org/site/c2010/nrp-032b.pdf
NRP	NRP-032C	In participants undergoing resuscitation courses (P), does the inclusion of more realistic techniques (eg. high fidelity manikins, in-situ training) (I), as opposed to standard training (eg. low fidelity, education centre) (C), improve outcomes (eg. skills performance) (O).	Impact of realistic training on skills performance	Khalid Aziz	http://circ.ahajournals.org/site/c2010/nrp-032c.pdf
NRP	NRP-033A	For hospital resuscitation teams (P), do team briefings/debriefings (I), when compared to no briefings/debriefings (C), improve team performance (O)? (INTERVENTION)	Impact of debriefing on team performance	Dianne L. Atkins, Nalini Singhal	http://circ.ahajournals.org/site/c2010/nrp-033a.pdf
NRP	NRP-033B	For hospital resuscitation teams (P), do team briefings/debriefings (I), when compared to no briefings/debriefings (C), improve team performance (O)? (INTERVENTION)	Impact of debriefing on team performance	Louis P. Halamek	http://circ.ahajournals.org/site/c2010/nrp-033b.pdf

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