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Queensland Maternity and Neonatal Clinical Guideline

**Neonatal resuscitation** 



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#### Flow Chart: Newborn life support

- Apply each set of activities in the flowchart for 30 seconds then assess response
- Progress to the next step if the heart rate, breathing, tone and oxygenation do not improve or the neonate is deteriorating
- Continuing care after resuscitation may include cardiorespiratory management, blood glucose management, antibiotics, induced hypothermia for hypoxic ischaemic encephalopathy, and stabilisation and transfer



Source: Australian Resuscitation Council Online February 2011: Flowcharts Queensland Maternity and Neonatal Clinical Guideline: MN11.5-V2-16 Neonatal resuscitation

## Abbreviations

bpm	Beats per minute
BGL	Blood glucose level
CPAP	Continuous positive airway pressure
CO <sub>2</sub>	Carbon dioxide
CTG	Cardiotocograph
ETT	Endotracheal tube
F	French gauge
HIE	Hypoxic-ischaemic encephalopathy
HR	Heart rate
H <sub>2</sub> O	Water
IM	Intramuscular
IV	Intravenous
LM	Laryngeal mask
PEEP	Positive end expiratory pressure
PIP	Peak inspiratory pressure
PPV	Positive pressure ventilation
QCC	Queensland Emergency Medical Systems Coordination Centre
SpO2	Saturation of peripheral oxygen
UVC	Umbilical venous catheter

## Definition of terms

Endotracheal	Intratracheal
Intratracheal	Endotracheal
Newborn	Refers to the infant in the first few minutes to hours following birth <sup>1</sup>
Neonate	Refers to the infant in the first 28 days of life <sup>1</sup>
Infancy	Infancy includes the neonatal period and extends through the first 12 months of life <sup>1</sup>

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# 1 Introduction

Although the need for neonatal resuscitation can often be anticipated, on many occasions it is unexpected<sup>1</sup>:

- Most newborns are vigorous at birth<sup>1</sup>:
  - Approximately 10% will require some assistance at birth to begin breathing
  - o Less than 1% will require extensive resuscitation

The guideline focus is on the neonatal period and particularly newborns. Paediatric resuscitation techniques may be used in term neonates after the newborn period and specifically in the situation of cardiac aetiology underlying their arrest.

## 1.1 Clinical standards

Clinical standards recommended for maternity facilities providing planned birthing services are outlined in Table 1.

Table 1. Clinical standards recommended for facilities providing pla	anned birthing services
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Aspect	Consideration		
Resources <sup>1</sup>	<ul> <li>Required at all times:         <ul> <li>A suitable place to resuscitate a newborn</li> <li>Suitable neonatal resuscitation equipment</li> <li>Clinicians trained in neonatal resuscitation</li> </ul> </li> <li>Organised programs to develop and maintain the standards, skills and teamwork required for newborn resuscitation:         <ul> <li>Training requires regular reinforcement in clinical practice and/or refresher courses which should be undertaken at least annually</li> </ul> </li> <li>Debriefing services for clinicians:         <ul> <li>Regardless of seniority, resuscitations can be stressful</li> <li>Reflection on practice provides a valuable learning opportunity</li> </ul> </li> </ul>		
Clinical skill requirements <sup>1</sup>	<ul> <li>Clinicians trained in:         <ul> <li>Basic neonatal resuscitation:</li> <li>Airway support, ventilation via face mask and chest compressions</li> <li>Advanced neonatal resuscitation:</li> <li>All the skills of basic neonatal resuscitation</li> <li>Endotracheal intubation</li> <li>Vascular cannulation</li> <li>The use of drugs and fluids</li> </ul> </li> <li>Clinicians responsible for neonatal resuscitation should be familiar with available neonatal resuscitation equipment</li> </ul>		
Clinician attendance at births <sup>1</sup>	<ul> <li>Low risk births: <ul> <li>A clinician trained in basic neonatal resuscitation should be in attendance and responsible only for the care of the newborn</li> <li>The Australian Resuscitation Council recommends a clinician trained in advanced neonatal resuscitation should also be available</li> </ul> </li> <li>High risk births: <ul> <li>A clinician trained in advanced neonatal resuscitation should be in attendance and responsible only for the care of the newborn</li> <li>More than one experienced person should be present to care for the newborn</li> </ul> </li> <li>The Queensland Clinical Services Capability Framework describes the first principle as applying to all services and the second principle as applying to Level 3 services and above.<sup>2</sup> Advanced neonatal resuscitation may not be possible in all Level 1 and 2 services</li> </ul>		
Documentation <sup>1</sup>	<ul> <li>Comprehensive and contemporaneous</li> <li>When possible, one person should be appointed to document, the time, interventions and newborn's response during resuscitation</li> <li>Use of a standardised neonatal resuscitation record is recommended [refer to Appendix A: Sample form - Neonatal resuscitation record]</li> </ul>		

# 2 Anticipating the need for resuscitation

A variety of maternal, fetal and intrapartum circumstances can increase the risk of the newborn needing resuscitation at birth<sup>1</sup> [refer to Table 2]

Risk factors					
	<ul> <li>Prolonged rupture of membranes (greater than 18 hours)</li> </ul>				
	Bleeding in second or third trimester				
	Pregnancy induced hypertension				
	Chronic hypertension				
	Substance abuse				
	<ul> <li>Drug therapy (e.g. lithium, magnesium, adrenergic blocking agents, narcotics)</li> </ul>				
Maternal <sup>1</sup>	Diabetes mellitus				
	Chronic illness (e.g. anaemia, cyanotic congenital heart disease)				
	Maternal pyrexia				
	Maternal infection				
	Chorioamnionitis				
	Heavy sedation				
	Previous fetal or neonatal death				
	No prenatal care				
	Multiple gestation (e.g. twins, triplets)				
	<ul> <li>Preterm gestation (especially less than 35 weeks)</li> </ul>				
	<ul> <li>Post term gestation (greater than 41 weeks)</li> </ul>				
	Large for dates				
	Fetal growth restriction				
Fetal <sup>1</sup>	<ul> <li>Alloimmune haemolytic disease (e.g. anti-D, anti-Kell, especially if fetal anaemia or hydrops fetalis present)</li> </ul>				
	<ul> <li>Polyhydramnios and oligohydramnios</li> </ul>				
	<ul> <li>Reduced fetal movement before onset of labour</li> </ul>				
	<ul> <li>Congenital abnormalities which may effect breathing, cardiovascular function or other aspects of perinatal transition</li> </ul>				
	Intrauterine infection				
	Hydrops fetalis				
	Non reassuring fetal heart rate patterns on cardiotocograph (CTG)				
	Abnormal presentation				
	Prolapsed cord				
	<ul> <li>Prolonged labour (or prolonged second stage of labour)</li> </ul>				
	Precipitate labour				
Intrapartum <sup>1</sup>	• Antepartum haemorrhage (e.g. abruption, placenta praevia, vasa praevia)				
	Meconium in the amniotic fluid				
	<ul> <li>Narcotic administration to mother within 4 hours of birth</li> </ul>				
	Forceps birth				
	Vacuum-assisted (ventouse) birth				
	Maternal general anaesthesia				

# 2.1 Communication and information sharing

Preparation for a high risk birth requires effective communication and information sharing and a consistent and coordinated approach between obstetric and neonatal clinicians [refer to Table 3].

Table 3. Con	nmunication and	l information
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Aspect	Consideration			
Obstetric and neonatal clinicians	<ul> <li>Maternal information should include details of<sup>1</sup>:         <ul> <li>Pre-existing or pregnancy related medical condition and treatment that may affect the resuscitation or management of the newborn</li> <li>Antenatal ultrasound diagnoses that may affect immediate postnatal management</li> <li>Assessments of fetal wellbeing (e.g. fetal heart rate monitoring)</li> </ul> </li> <li>Minimum fetal information required by personnel responsible for the baby includes<sup>1</sup>:         <ul> <li>Gestational age</li> <li>Number of expected newborns, if multiple birth</li> <li>Reason this is a high risk birth</li> <li>Presence of meconium in the liquor</li> <li>Assessments of fetal heart rate variability</li> <li>Any known congenital abnormalities</li> <li>Maternal risk factors for infections (including results of screening if known, e.g. Group B Streptococcus)</li> <li>Maternal medication</li> </ul> </li> </ul>			
Parent(s)	<ul> <li>Whenever time permits, the neonatal team should introduce themselves to the parent(s) before the birth and should<sup>1</sup>:         <ul> <li>Outline the proposed plan for the newborn's care</li> <li>Enquire and respond to parent's questions</li> </ul> </li> <li>In cases of potentially lethal fetal malformations or extreme prematurity, whenever possible, parents should be included in decisions about the extent of the resuscitation<sup>1</sup></li> </ul>			
Documentation	Contemporaneous documentation of discussions and care plans			

## 2.2 Equipment

 A complete set of resuscitation equipment and drugs should always be readily available in the areas of hospitals where newborns are born or receive neonatal care<sup>1</sup> [refer to Appendix B: Equipment required for neonatal resuscitation]

## 2.2.1 Equipment checks

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- Facilities should maintain a clear record documenting the checking procedure for each set of resuscitation equipment and drugs<sup>1</sup>
- Each set of resuscitation equipment and drugs should be checked:
  - Before any resuscitation<sup>1</sup> and
    - According to local hospital policy.<sup>1</sup> Consider frequency of checking:
      - At least daily, preferably once per shift
        - Following resuscitation
        - To enable clinical staff familiarisation of layout in the emergency situation

# 3 Assessment of the newborn at birth

Assessing the need to initiate and continue resuscitation should begin immediately after birth and proceed throughout the resuscitation.<sup>1</sup> Evaluation and intervention (if required) are simultaneous processes, especially when more than one resuscitator is present.<sup>1</sup> For clarity this process is described as a sequence of distinct steps<sup>1</sup> [refer to Flowchart: Newborn life support]. Table 4 outlines important aspects of assessment at birth.<sup>1</sup>

Table 4.	Assessment	of the	newborn	at birth
10010 11	,	01 010	1101100111	

Aspect	Consideration
Cord clamping	<ul> <li>In the compromised newborn, the optimal timing of cord clamping remains unknown<sup>1</sup></li> </ul>
	<ul> <li>The more severely compromised the newborn the more likely it is that resuscitation measures need to take priority over delayed cord clamping</li> </ul>
	• Tone
Initial assessment	Breathing
	Heart rate
	Heart rate:
	<ul> <li>A prompt increase in heart rate remains the most sensitive indicator of resuscitation efficacy<sup>1</sup></li> </ul>
Subsequent	Breathing
assessment	• Tone
	Oxygenation:
	<ul> <li>Preferably assessed using pulse oximetry</li> </ul>
	<ul> <li>Hypothermia may increase oxygen consumption impeding effective resuscitation<sup>1</sup></li> </ul>
	<ul> <li>Ensure area is warm and draft free<sup>1</sup></li> </ul>
	<ul> <li>For term and near term newborn – dry newborn and remove wet linen<sup>1</sup></li> </ul>
Temperature	For very premature newborns:
	$\circ$ Increase ambient temperature to 26 <sup>o</sup> C <sup>1</sup>
	<ul> <li>Placing the baby in a polyethylene bag (without drying) or under a polyethylene sheet (food or medical grade, heat resistant) up to the neck, (immediately after birth until temperature is stable)<sup>1</sup></li> </ul>
	<ul> <li>All newborns should be handled gently (premature newborns are especially at risk of damage to skin and internal organs)<sup>1</sup></li> </ul>
Handling and skin	<ul> <li>If emergency vascular access is required, antiseptic solution, particularly alcohol containing, should be applied sparingly<sup>1</sup>:</li> </ul>
protection	<ul> <li>For umbilical catheterisation:</li> </ul>
	<ul> <li>Apply only to the cord and less than 2 cm diameter of surrounding skin</li> </ul>
	<ul> <li>Avoid pooling around the newborn's groin and flanks</li> </ul>
	• Oximetry is recommended when <sup>1</sup> :
	<ul> <li>The need for resuscitation is anticipated</li> </ul>
	<ul> <li>Positive pressure ventilation (PPV) is administered for more than a few breaths</li> </ul>
Pulse oximetry	<ul> <li>Whenever persistent cyanosis is suspected during resuscitation or any time after birth</li> </ul>
	<ul> <li>When supplemental oxygen is administered</li> </ul>
	<ul> <li>The sensor should be placed on the newborn's right hand or wrist and then connect the probe to the instrument.<sup>1</sup></li> </ul>
	<ul> <li>This results in the quickest, accurate monitor display<sup>3</sup></li> </ul>
Supplemental oxygen	Refer to Table 7

# 4 Resuscitation management

## • Effective ventilation is the key to successful neonatal resuscitation<sup>1</sup>

- Drying and stimulation are both assessment and resuscitative interventions.<sup>1</sup> If the newborn fails to establish spontaneous effective respirations and heart rate does not increase to more than 100 beats per minute (bpm) then commence PPV<sup>1</sup>
- Effective teamwork is essential to the success of neonatal resuscitation, particularly advanced resuscitation
- Newborn resuscitation training should include the necessary individual and teamwork skills which are reinforced through regular practice<sup>1</sup>

#### 4.1 **Positive pressure ventilation**

 PPV may be delivered via a face mask, endotracheal tube (ETT) or laryngeal mask<sup>1</sup> (LM) [refer to Table 5 for important aspects of PPV<sup>1</sup>; Table 6 refers to devices for PPV<sup>1</sup>]

Aspect	Consideration			
Indications	<ul> <li>After stimulation commence PPV if:         <ul> <li>Newborn's heart rate is less than 100 bpm <i>and</i></li> <li>The newborn remains apnoeic <i>or</i></li> <li>Breathing is inadequate – gasping is not adequate breathing</li> </ul> </li> </ul>			
Technique	<ul> <li>Ensuring the jaw is supported, keep the neck slightly extended in the 'sniffing position'</li> <li>Two people may provide mask ventilation more effectively than one: <ul> <li>One person supports the jaw holding the mask in place with two hands</li> <li>The other person provides positive pressure breaths</li> </ul> </li> </ul>			
Rate	• 40-60 breaths per minute			
Pressures	<ul> <li>For most newborns, ventilation can be accomplished with progressively lower pressures and rates as resuscitation proceeds</li> </ul>			
Mouth to mouth and nose	<ul> <li>Should be used when neonatal inflation devices are not available</li> <li>Maternal blood and other body fluids should be wiped from the newborn's face to reduce the risk of infection to the resuscitator</li> </ul>			
Assessment of effectiveness	<ul> <li>Ventilation effectiveness is confirmed by observing:         <ul> <li>Heart rate increasing to greater than 100 bpm</li> <li>Slight rise of chest and upper abdomen with each inflation</li> <li>Improving oxygenation (preferably assessed using pulse oximetry)</li> </ul> </li> <li>Reassess technique if the:         <ul> <li>Chest and abdomen do not rise with each inflation</li> <li>Heart rate does not rise above 100 bpm</li> <li>Oxygenation does not improve</li> </ul> </li> <li>Consider tracheal intubation or use of laryngeal mask if face mask ventilation remains ineffective despite corrective intervention</li> </ul>			
PEEP during resuscitation	<ul> <li>If suitable equipment is available, positive end expiratory pressure (PEEP) (at least 5 cm H<sub>2</sub>O) should be used during resuscitation to:         <ul> <li>Assist lung expansion</li> <li>Help establish a functional residual capacity</li> <li>Improve oxygenation</li> </ul> </li> </ul>			
Continuous positive airway pressure (CPAP) <sup>1</sup>	<ul> <li>A trial of CPAP is reasonable for:         <ul> <li>Spontaneously breathing newborns who have laboured breathing / respiratory distress</li> <li>Newborns who are breathing but whose saturations are not meeting targets</li> </ul> </li> </ul>			

Table 5. Positive pressure ventilation

# 4.2 Positive pressure ventilation delivery devices

Table 6.	Positive	pressure	ventilation	deliver	v devices
1 4010 0.	1 001010	probuild	vontilation	aonvor	,

Aspect	Consideration
Acceptable delivery devices <sup>1</sup>	<ul> <li>T-piece device is preferred. Recommended initial pressures: <ul> <li>Peak inspiratory pressure (PIP):</li> <li>Term newborn 30 cm H<sub>2</sub>O (water)</li> <li>Preterm newborn 20 – 25 cm H<sub>2</sub>O</li> <li>PEEP: 5 – 8 cm H<sub>2</sub>O</li> <li>Maximum pressure relief valve: 50 cm H<sub>2</sub>O*</li> </ul> </li> <li>Self inflating bag <ul> <li>Pressure release valve factory set at approximately 40 cm H<sub>2</sub>O</li> </ul> </li> <li>Flow inflating (or anaesthetic) bag</li> <li>Face masks should be: <ul> <li>The appropriate size</li> <li>Cushioned</li> <li>Self-inflating bags cannot effectively deliver CPAP, PEEP or sustained inflation breaths</li> </ul> </li> </ul>
T-piece device*	<ul> <li>Although a maximum pressure relief valve set at 50 cm H<sub>2</sub>O is recommended by the Australian Resuscitation Council<sup>1</sup>, a lower maximum pressure relief limit for e.g. 35 cm H<sub>2</sub>O will be adequate for the majority of babies receiving resuscitation.</li> <li>Within each hospital, neonatal T-piece devices should be set up using standard flow and pressure settings that are appropriate for newborns and well known to all clinicians</li> <li>All clinicians should be trained to safely adjust pressure settings during resuscitation if necessary.</li> <li>Rarely babies will need high peak inspiratory pressures of greater than or equal to 50 cm H<sub>2</sub>O at the commencement of ventilation</li> </ul>

## 4.3 Supplemental oxygen administration

Supplemental oxygen administration should be provided only when oxygen saturation has failed to improve with effective ventilation of the lungs. Points for consideration are provided in Table 7.

Table 7. Supplemental oxygen

Aspect	Consideration	1			
	<ul> <li>Regardless of gestation, the goal of oxygen administration is an oxygen saturation resembling that of a healthy term newborn<sup>1</sup>:</li> </ul>				
	o Assess	the newborn's respons	se to supplemental oxy	/gen	
	<ul> <li>Reduce or cease supplemental oxygen if saturations are above 90% at any time in the first 10 minutes, although it is acceptable and normal for some newborn saturations to rise above 90% in room air</li> </ul>				
	<ul> <li>Table 8 resuscit</li> </ul>	below includes target ation:	saturations for newbor	ns during	
		Table 8. Target oxygen s	saturation		
		Time from birth	Target oxygen saturations		
		1 minute	60 – 70%		
		2 minutes	65 – 85%		
Complemental		3 minutes	70 – 90%		
oxygen		4 minutes	75 – 90%		
administration		5 minutes	80 – 90%		
		10 minutes	85 – 90%		
	<ul> <li>The priority is to ensure adequate lung inflation - only increase inspired oxygen concentration if PPV or CPAP have failed to achieve the target peripheral oxygen saturation (SpO<sub>2</sub>)<sup>1</sup></li> <li>Term and near term newborns<sup>1</sup>:</li> </ul>				
	<ul> <li>Use air initially and only administer oxygen to newborns whose saturations do not meet the lower target range despite respiratory support</li> </ul>				
	<ul> <li>Reduce oxygen concentration if saturations reach 90% while supplemental oxygen is used</li> </ul>				
	<ul> <li>Preterm newborns (less than 32 weeks)<sup>1</sup>:</li> </ul>				
	<ul> <li>Initially use room air or blended air and oxygen (at 30-50%)</li> </ul>				
	<ul> <li>Guided by pulse oximetry, adjust oxygen concentration according to response</li> </ul>				
	o If blende	ed air and oxygen are	unavailable, initiate res	suscitation with air	

## 4.4 Endotracheal intubation

A decision to perform endotracheal intubation will depend on the newborn's gestation, degree of respiratory depression, response to face mask ventilation, and the skill and experience of the resuscitator.<sup>1</sup> [refer to Table 9]

Table 9. Endotracheal intubation

Aspect	Consideration
Clinical standard <sup>1</sup>	<ul> <li>Neonatal endotracheal intubation should only be attempted by clinicians with appropriate training and experience in the procedure</li> <li>If there is no-one skilled at intubation present, continue PPV via a face.</li> </ul>
	mask or LM [refer to Table 10]
	<ul> <li>Tracheal suctioning in a non vigorous newborn exposed to meconium stained liquor<sup>4</sup></li> </ul>
	• Onsuccessful ventilation via a face mask (e.g. heart fate remains low, oxygen saturation falling or failing to rise or prolonged)
Indications <sup>1</sup>	Special circumstances (e.g. diaphragmatic hernia, extremely low birth weight)
	Newborns born without a detectable heart rate
	<ul> <li>Administration of endotracheal medications (e.g. Adrenaline or artificial surfactant)</li> </ul>
	Expected need for continued or prolonged ventilation
ETT size and insertion length <sup>1</sup>	• ETT size and insertion length are based on the newborn's weight and corrected gestation [refer to Appendix C]
Position verification <sup>1</sup>	<ul> <li>ETT insertion depth must always be checked by comparing the markings on the tube with the formula or table [refer to Appendix C]</li> <li>Primarily correct ETT position is confirmed by effective ventilation demonstrated by:         <ul> <li>Chest movement with each inflation</li> <li>Heart rate increasing above 100 bpm</li> <li>Improving oxygenation (oximetry is more accurate than visual assessment)</li> </ul> </li> <li>Other signs to verify correct ETT placement include:         <ul> <li>Visual inspection of the ETT passing through the larynx</li> <li>Misting in the ETT during expiration (usually visible for only the first few exhaled breaths)</li> <li>Colour change visible with an end-tidal carbon dioxide (CO<sub>2</sub>) detector (the most reliable method in newborns who have spontaneous circulation):             <ul> <li>False negatives may occur in newborns if there is very low or absent pulmonary blood flow</li> <li>False positives may occur in devices contaminated by adrenaline or surfactant</li> <li>Hearing symmetrical breath sounds using a stethoscope placed on the upper chest. May be asymmetrical, despite optimal tube position, in some circumstances (e.g. pneumothorax, diaphragmatic hernia)</li> </ul> </li> </ul></li></ul>
Signs that ETT is not in the trachea	<ul> <li>No chest movement with inflations<sup>1</sup></li> <li>A heart rate less than 100 bpm that does not increase soon after intubation and inflation has started<sup>1</sup></li> <li>No detection of expired CO<sub>2</sub><sup>1</sup></li> <li>No improvement in oxygenation<sup>1</sup> or sudden deterioration in oxygenation</li> <li>The absence of breath sounds in the axillae<sup>1</sup></li> </ul>

## 4.4.1 Intubation under specific circumstances

Tracheal intubation may be required under specific circumstances [refer to Table 10]

Circumstance	Consideration
	<ul> <li>Need for endotracheal suctioning is based on the newborn's level of activity at birth<sup>4</sup></li> </ul>
	• Only perform intubation to suction meconium from the trachea <sup>1</sup> :
	<ul> <li>In newborns with decreased muscle tone</li> </ul>
Meconium stained	<ul> <li>Immediately after birth</li> <li>Before the onset of breathing or crying (stimulation should be withheld until suction is completed)</li> <li>If an experienced clinician and all the needed equipment are</li> </ul>
ilquor	<ul> <li>Once and then any subsequent resuscitation that is needed should be commenced quickly</li> </ul>
	<ul> <li>Use a meconium aspirator device attached to the ETT adapter after intubation</li> </ul>
	<ul> <li>Negative pressure used should not exceed 100 mmHg</li> </ul>
Very premature	<ul> <li>Many very premature newborns can be treated with CPAP without intubation and ventilation<sup>1</sup></li> </ul>
than 29 weeks)	<ul> <li>Early administration of artificial surfactant can improve response in premature newborns who require endotracheal intubation<sup>1</sup></li> </ul>
	<ul> <li>A LM should be considered during resuscitation of the term and near term newborn if<sup>1</sup>:</li> </ul>
	<ul> <li>Face mask ventilation is unsuccessful (newborns greater than 2000 g or greater than or equal to 34 weeks gestation)</li> <li>Tracheal intubation is unsuccessful or not feasible</li> </ul>
	<ul> <li>A size 1 LM is suitable up to 5 kgs<sup>1</sup></li> </ul>
	<ul> <li>Following LM insertion, correct position is confirmed by effective ventilation as demonstrated by<sup>1</sup>:</li> </ul>
	<ul> <li>Chest movement with each inflation</li> </ul>
Unsuccessful facemask and/or	<ul> <li>Heart rate increasing above 100 bpm</li> <li>Improving oxygenation (oximetry is more accurate than visual assessment)</li> </ul>
tracheal intubation	Other signs to verify correct LM placement include:
	$\circ$ Colour change visible with an end-tidal CO <sub>2</sub> detector (the most reliable
	<ul><li>method in newborns who have spontaneous circulation):</li><li>False negatives may occur in newborns if there is very low or</li></ul>
	<ul> <li>absent pulmonary blood flow</li> <li>Hearing symmetrical breath sounds using a stethoscope placed on the upper chest. May be asymmetrical, despite optimal tube position,</li> </ul>
	in some circumstances (e.g. pneumothorax, diaphragmatic hernia)
	<ul> <li>There has not been an evaluation of the LM for administration of emergency intra-tracheal medications<sup>1</sup></li> </ul>
	<ul> <li>Refer to Appendix D for practical instructions in use of a LM</li> </ul>

Table 10. Intubation under specific circumstances

# 4.5 Chest compressions

When adequate airway and breathing support are provided, cardiac compressions are rarely indicated for resuscitation of neonates. In neonates, cardiac output is heart rate dependent. If the heart rate is too slow, circulation will be inadequate to support tissue oxygenation.<sup>1</sup> [refer to Table 11]

Aspect	Consideration
Indication	• Heart rate remains less than 60 bpm despite 30 seconds of effective PPV <sup>4</sup>
Supplemental oxygen	Administer 100% oxygen
Technique	<ul> <li>Perform a chest compression each half second. In a half second pause after each 3<sup>rd</sup> compression, deliver a breath. This results in a 3:1 ratio with a total of 90 compressions and 30 breaths in each minute<sup>1</sup></li> <li>Coordinate compressions and inflations to avoid simultaneous delivery of a compression and a breath</li> <li>Methods: <ul> <li>2 thumb technique:</li> <li>Strongly recommended when two clinicians are available. (Usually the resuscitator performing chest compressions faces the newborn's head. The position can be reversed if access to the abdomen is required) <i>or</i></li> <li>2 finger technique – acceptable as an interim measure or where access is required to the lower chest and abdomen</li> </ul> </li> </ul>
	<ul> <li>administrator's hands should not leave the chest</li> <li>Effectively delivered chest compressions will be evident on a pulse oximeter</li> </ul>
Duration	<ul> <li>Continue uninterrupted compressions for at least 30 seconds between each pause to assess improvement in spontaneous heart rate and cardiac output</li> <li>Continue chest compressions with as little interruption as possible until it is obvious that the heart rate is greater than 60 bpm</li> </ul>
Assessment	<ul> <li>Improvement is evidenced by:         <ul> <li>Audible heart rate on auscultation</li> <li>Spontaneous pulsations on oximetry</li> <li>Note: Chest compressions will also cause pulsations on the pulse oximeter</li> <li>Rise in oxygen saturation</li> <li>Spontaneous movement or breaths</li> </ul> </li> </ul>

Table 11. Combined	chest compr	ession and	positive p	oressure	ventilation

## 4.6 Drugs and fluids

Drugs and fluids are rarely indicated for resuscitation of newborns<sup>1</sup>:

- Bradycardia is usually caused by hypoxia and inadequate ventilation and apnoea by insufficient oxygenation of the brainstem. Establishing adequate ventilation is the most important step to improve heart rate, however if heart rate remains less than 60 bpm despite adequate ventilation, Adrenaline may be needed<sup>1</sup>
- Administration of drugs should not detract from the efficiency and continuity of ventilation and chest compressions
- For administration of Adrenaline and volume expanding fluids, refer to Table 13 and Table 14 below
- Refer to Appendix E for the rarely indicated use of a narcotic antagonist

#### 4.6.1 Administration routes

Preferred vascular access is via the umbilical vein.<sup>1,4</sup> Alternative access is via an endotracheal tube, peripheral vein, or intraosseous lines. [refer to Table 12]

Table 12. Administration routes

Route	Considerations <sup>1</sup>
Umbilical vein	<ul> <li>The umbilical vein is the most rapidly accessible intravascular route for Adrenaline and an umbilical venous catheter (UVC) can also be used for fluid administration</li> <li>A 3-way tap should be attached to the UVC and both primed with normal saline before use</li> </ul>
	<ul> <li>A UVC can provide continued vascular access until an alternative route is established</li> </ul>
	<ul> <li>Blood gases from the UVC may be useful in guiding treatment</li> </ul>
Endotracheal tube	<ul> <li>Give only Adrenaline via the ETT:         <ul> <li>ETT administration of Adrenaline is acceptable when the heart rate is less than 60 bpm despite adequate ventilation and chest compressions and when there is no intravenous (IV) access<sup>1</sup></li> <li>Administration of ETT Adrenaline should not delay attempts to obtain vascular access for administration of IV Adrenaline</li> </ul> </li> </ul>
Peripheral vein	<ul><li>Access can be difficult</li><li>Access may take too long</li></ul>
Intraosseous lines	<ul> <li>Not commonly used in newborns</li> <li>Can be used if umbilical or venous access is not available</li> <li>Consider if the resuscitator has greater experience with inserting intraosseous lines</li> </ul>

#### 4.6.2 Adrenaline

Table 13. Adrenaline administration

Adrenaline 1:10,000		
Indication	<ul> <li>If the heart has failed to increase to greater than 60 bpm within one minute of adequate ventilation and chest compressions</li> </ul>	
Route	<ul> <li>Intravascular is the preferred route:         <ul> <li>Give as close to the heart as possible:                 <ul> <li>UVC is preferable to peripheral IV or intraosseous<sup>1</sup></li> </ul> </li> <li>ETT:                 <ul> <li>Do not allow ETT dose to delay attempts at vascular access<sup>1</sup></li> </ul> </li> </ul> </li> </ul>	
IV dose and administration	<ul> <li>0.01-0.03 mg/kg (equates to 0.1-0.3 mL/kg)<sup>1</sup></li> <li>Refer to Appendix C for suitable rounded doses</li> <li>Give as a rapid bolus followed by 0.9% Sodium Chloride flush<sup>1</sup></li> <li>Practice tip:</li> <li>If resuscitating a term baby, draw up 1 mL of Adrenaline and administer it all</li> </ul>	
ETT dose and administration	<ul> <li>0.05-0.1 mg/kg (equates to 0.5-1.0 mL/kg)<sup>1</sup></li> <li>Follow with PPV – Flush not recommended<sup>1</sup></li> <li>Ensure the Adrenaline dose is ventilated into the lungs and is not left in the ETT or the adaptor</li> </ul>	
Frequency	<ul> <li>Repeat every few minutes, if the heart rate remains less than 60 bpm despite effective ventilation and cardiac compressions<sup>1</sup></li> <li>If ETT dose administered, with inadequate effect, give IV dose as soon as vascular access is obtained</li> </ul>	
Dilution	Not required	

# 4.6.3 Volume expanding fluids

Table 14. Fluid administration

0.9% Sodium chloride or O Rhesus negative blood <sup>1</sup>		
Indication	<ul> <li>Initially 0.9% Sodium Chloride for:         <ul> <li>Suspected or proven blood loss</li> <li>Neonate who appears to be in shock (e.g. pale, poor perfusion and weak pulse) and has not responded adequately to other resuscitative measures</li> </ul> </li> <li>Subsequently blood suitable for emergency transfusion:         <ul> <li>In the setting of massive blood loss</li> <li>In babies not responding to resuscitation</li> </ul> </li> </ul>	
Caution	<ul> <li>Routine fluid boluses during or after resuscitation may do more harm than good</li> </ul>	
Route	<ul><li>Intravascular e.g. IV, UVC, intraosseous</li><li>Refer to Table 12</li></ul>	
Method of administration	Infusion over several minutes	
Dose	• 10 mL/kg	
Frequency	May be repeated after observation of response if only minimal improvement	
Dilution	Not required	

# 5 Care of the neonate after resuscitation

The neonate who has required resuscitation remains at risk and requires ongoing assessment.<sup>1</sup> [refer to Table  $15^1$  and Guideline: Neonatal stabilisation<sup>5</sup>

Care	Comments					
Location	Assess the need for admission to an intensive or special care nursery					
Monitoring	<ul> <li>Monitoring required may include:         <ul> <li>Oxygen saturation</li> <li>Heart rate</li> <li>Respiratory rate and pattern</li> <li>Blood glucose measurement</li> <li>Blood gas analysis</li> <li>Fluid balance and nutrition</li> <li>Blood pressure</li> <li>Temperature</li> <li>Neurological</li> </ul> </li> </ul>					
Blood glucose management	<ul> <li>Neonates who require resuscitation are susceptible to hypoglycaemia</li> <li>Check the blood glucose level (BGL) as soon as possible after resuscitation</li> <li>Maintain the BGL greater than 2.5 mmol/L</li> <li>Avoid large bolus doses of glucose (greater than 100 – 200 mg/kg) <ul> <li>1 mL 10% Glucose contains 100 mg Glucose</li> </ul> </li> <li>Refer to Guideline: Neonatal hypoglycaemia and blood glucose level monitoring<sup>6</sup></li> </ul>					
Antibiotics	<ul> <li>The need for resuscitation can be a consequence of sepsis. Soon after resuscitation consider:         <ul> <li>Relevant investigations</li> <li>The need for antibiotics</li> </ul> </li> </ul>					
Induced hypothermia for hypoxic-ischaemic encephalopathy	<ul> <li>Discuss any neonate with hypoxic-ischaemic encephalopathy (HIE) who is considered for induced hypothermia early with a Neonatologist</li> <li>Plans should be made for transfer (if required) and admission to a neonatal intensive care unit</li> <li>Refer to Guideline: Hypoxic-ischaemic encephalopathy<sup>6</sup></li> </ul>					
Inter-hospital transfer	<ul> <li>If the birth facility is unable to provide an appropriate level of post resuscitation monitoring and support, transfer the neonate to an appropriate higher level facility</li> <li>To discuss the decision to transfer and to arrange transfer of the neonate to a higher level facility, contact Queensland Emergency Medical Systems Coordination Centre (QCC) by phoning 1300 799 127</li> </ul>					
Continuing care of the family	<ul> <li>Parent(s) find witnessing the resuscitation of their baby distressing regardless of outcome. Whenever possible:         <ul> <li>Prepare parents for resuscitation if it is anticipated</li> <li>Keep parent(s) informed during and after resuscitation. Ideally information should be provided by a senior clinician</li> <li>Facilitate early parental contact with their baby</li> </ul> </li> </ul>					

Table 15. Continuing care of the neonate after resuscitation

# 6 Ethical issues

The birth of extremely premature neonates and those with severe congenital anomalies raise questions with parent(s) and among clinicians about initiation of resuscitation.<sup>1</sup> [refer to Table 16<sup>1</sup>].

Table 16. Ethical issues	able 16. Ethical	issues
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Issue	Consideration						
Initiating resuscitation	<ul> <li>Where unexpected anomalies are present offer full resuscitation, ongoing care can be discussed with parent(s) after assessment of the neonate's condition.<sup>4</sup> Examples of exceptions include neonates with<sup>4</sup>:         <ul> <li>Anencephaly</li> <li>Extreme prematurity for whom there is very little possibility of intact survival</li> </ul> </li> </ul>						
	<ul> <li>Consider discontinuing resuscitation if the heart rate is undetectable and remains so for 10 minutes in a newly born baby<sup>1</sup></li> </ul>						
	<ul> <li>The decision to continue resuscitation beyond 10 minutes when there is no or very low heart rate may be influenced by<sup>1</sup>:</li> </ul>						
Discontinuing	<ul> <li>Presumed diagnosis</li> <li>Gestation of neonate</li> </ul>						
resuscitation	<ul> <li>Presence or absence of complications</li> <li>Parent(s) previously expressed views regarding acceptable risk of morbidity</li> </ul>						
	• When withdrawing or withholding resuscitation, care should be focused on the neonate's comfort and dignity <sup>1</sup>						
	<ul> <li>Autopsy may be appropriate after failed resuscitation.[refer to Guideline: Stillbirth care<sup>7</sup>]</li> </ul>						

# References

1. Australian Resuscitation Council. Section 13 - Neonatal Guidelines. 2010 Dec [cited 2011 March 15]. Available from: <u>http://www.resus.org.au/</u>.

2. Queensland Health. Clinical Services Capability Framework for Public and Licensed Private Health Facilities v3.0. Brisbane: Queensland Government Department of Health. 2011.

3. O'Donnell CPF, Kamlin COF, Davis PG, Morley CJ. Obtaining pulse oximetry data in neonates: a randomised crossover study of sensor application techniques. Archives of disease in childhood. Fetal and neonatal edition. 2005; 90:F84-5.

4. Kattwinkel J, Perlman JM, Aziz K, Colby C, Fairchild K, Gallagher J, et al. Part 15: neonatal resuscitation: 2010 American Heart Association Guidelines for Cardiopulmonary Resuscitation and Emergency Cardiovascular Care. Circulation. 2010; 122(suppl3):S909-19.

5. Queensland Maternity and Neonatal Clinical Guidelines Program. Neonatal stabilisation. Guideline No. MN11.12-V1-R16. Queensland Health. 2011.

6. Queensland Maternity and Neonatal Clinical Guidelines Program. Neonatal hypoglycaemia and blood glucose level monitoring. Guideline No. MN10.8-V2-R12. Queensland Health. 2010.

7. Queensland Maternity and Neonatal Clinical Guidelines Program. Stillbirth care. Guideline No. MN11.24-V3-R16. Queensland Health. 2011.

# Appendix A: Sample form - Neonatal resuscitation record

Hospital Emergency Telephone Number:

Surname							
Given Names							
DOBSex							
Affix patient identification label here	Affix patient identification label here						
Date Time of Birth/Event Time resuscitation commenced	Time resuscitation commenced						
Est Time code called Weight (if applicable) Time resuscitation ceased							
Resuscitation Team         Name         Signature         Arrival Time         APGAR         1st min	5	10	15				
Doctor in charge Colour							
Doctor Respiratory Rate							
Doctor Heart Rate							
Doctor Tone							
Nurse/Midwife Reflex							
Nurse/Midwife Total							
Nurse/Midwife							
Nurse/Midwife							
Scribe							
Other							
Summary of Events Leading up to Need for Resuscitation (Continue or expand in clinical notes as needed)							
in form							
a mnle ivit							
Samp							
Observations Intervention Note: Follow resuscitation flow diagram	Intervention						
Time         Intubation         External         Adrenaline           HR         Resp         O2 Sat         PPV         O2%         ETT size         Cardiac         1:10,000	tions &	medicat	ions				
CO2 detector Massage dose & route							

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Logo								a and Ma				
								Sumame				
							Given N	Given Names				
							DOB				Sex	
N	Neonatal Resuscitation Record											
	Ob	oservatio	ons	Intervention				diagra	diagram All		other observations,	
Time			0.04			Intub	ation External			Adrenaline	re: interve	sponses, events,
	HR	Resp	O <sub>2</sub> Sat	PPV	02%	CO <sub>2</sub> d	size etector	Massa Massa	iac age	1:10,000 dose & route	interve	intions & medications
												•
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								E		-		
				5/								
						Outcom	e of Rev	suscitati	ion			
					(Continu	e or expa	ind in clini	ical notes	as nee	eded)		
									Sign	ature		
Cord das results:							lf die	d. Time of de	eath			
oord ga	Who Informed Parents											
Print Na	Print Name Signature Date Time											
	Pre-Transfer Check											
ETT see	cured at	:	cm				Lines se	ecured?			Yes	
Transfe	r arrand	ed to			ICN			SCN			Postnat	al ward
Docume	entation	complet	te?		Yes		Identific	ation co	mplete	e? □	Yes	
Names	& Siana	ture con	nplete?		Yes		Medica	tions pre	scribe	d? □	Yes	

Use this form when Neonatal Resuscitation is required and/or Neonatal Code Blue is called

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# Appendix B: Equipment required for neonatal resuscitation

Equipment	Comment				
Standard kits <sup>1</sup>	• Prior preparation of standardised kits containing equipment for procedures (e.g. umbilical catheterisation) can save considerable time in emergencies				
General <sup>1</sup>	<ul> <li>Firm padded resuscitation surface</li> <li>Overhead warmer</li> <li>Light for the area</li> <li>Clock with timer in seconds</li> <li>Warm towel or similar covering</li> <li>Polyethylene bag or sheet, big enough for neonates less than 1500 g birth weight</li> <li>Stethoscope (neonatal preferred)</li> <li>Pulse oximeter plus neonatal probe</li> </ul>				
Airway management <sup>1</sup>	<ul> <li>Suction apparatus and suction catheters (sizes 6F, 8F, and either 10F or 12F)</li> <li>Oropharyngeal airways (sizes 00 and 0)</li> <li>Intubation equipment: <ul> <li>Laryngoscopes with neonatal blades (sizes 00, 0, 1)</li> <li>Spare batteries and bulbs</li> <li>Compatible laryngoscope handles and blades</li> <li>Endotracheal tubes (ETT) (uncuffed, no eye, sizes 2.5, 3.0, 3.5 and 4.0 mm internal diameter)</li> <li>ETT introducer/stylet</li> <li>Supplies for fixing ETT (e.g. sterile scissors, tape)</li> </ul> </li> <li>End-tidal carbon dioxide (CO<sub>2</sub>) detector (to confirm intubation)</li> <li>Meconium suction device (to apply suction directly to ETT)</li> <li>Magill forceps (neonatal size - optional)</li> <li>Laryngeal mask (size 1, suitable for neonates up to 5 kg)</li> </ul>				
Breathing support <sup>1</sup>	<ul> <li>Face masks (range of sizes suitable for premature and term neonates not Rendell Baker type)</li> <li>Positive pressure ventilation device either: <ul> <li>T-Piece resuscitation device (preferred), or</li> <li>Flow-inflating bag with pressure safety valve and manometer, and</li> <li>Self-inflating bag (240 mL) with a pressure release valve and a removable oxygen reservoir</li> </ul> </li> <li>Medical gases: <ul> <li>Source of medical oxygen (reticulated and/or cylinder allowing flow rate of up to 10 L/min) with flow meter and tubing</li> <li>Source of medical air plus air/oxygen blender</li> </ul> </li> <li>Feeding tubes for gastric decompression (sizes 6F, 8F)</li> </ul>				
Circulation support <sup>1</sup>	<ul> <li>Umbilical venous catheter (UVC) kit (including UVC sizes 3.5F and 5F)</li> <li>Peripheral IV cannulation kit</li> <li>Skin preparation solution suitable for neonate skin</li> <li>Tapes/devices to secure UVC/IV cannula</li> <li>Syringes and needles (assorted sizes)</li> <li>Intraosseous needles (50 mm length)</li> </ul>				
Drugs and fluid <sup>1</sup>	<ul> <li>Adrenaline 1:10 000 concentration (0.1 mg/mL)</li> <li>Volume expander:         <ul> <li>Normal saline (0.9% Sodium Chloride)</li> <li>Blood suitable for emergency neonatal transfusion should be readily available</li> </ul> </li> </ul>				
<b>Documentation</b> <sup>1</sup>	<ul><li>Resuscitation record sheet</li><li>Equipment check list</li></ul>				

# Appendix C: Endotracheal tube length and size and Adrenaline dosage by corrected gestation

The following ETT references are sourced from the Australian Resuscitation Council, Section 13 - Neonatal Guidelines, 2010 [accessed 2011 March 15, <u>http://www.resus.org.au/]</u>.

#### Endotracheal tube (ETT):

- Corrected gestation gestation at birth plus postnatal age
- ETT internal diameter in millimetres can be calculated as (gestational age in weeks divided by 10):
  - o 2.5 mm ETT for neonates less than 1 kg
  - 3.0 mm ETT for neonates 1 − 2 kg
  - o 3.5 mm ETT for neonates 2 3 kg
  - o 3.5 4.0 mm ETT for neonates greater than 3 kg
- Approximate ETT insertion depth from middle of upper lip can be calculated as (weight in kg + 6 cm)

Corrected gestation (weeks)	Actual weight (kg)	ETT mark at upper lip (cm)	ETT size - internal diameter (mm)	ETT suction catheter size (F)
23 – 24	0.5 - 0.6	5.5	2.5	5 or 6
25 – 26	0.7 – 0.8	6.0	2.5	5 or 6
27 – 29	0.9 - 1.0	6.5	2.5	5 or 6
30 - 32	1.1 – 1.4	7.0	3.0	6 or 8
33 – 34	1.5 – 1.8	7.5	3.0	6 or 8
35 – 37	1.9 – 2.4	8.0	3.5	8
38 - 40	2.5 – 3.1	8.5	3.5	8
41 – 43	3.2 - 4.2	9.0	3.5 - 4.0	8 or 10

#### Adrenaline:

Corrected gestation (weeks)	IV Adrenaline 1:10,000 (mLs)
23 – 26	0.1
27 – 32	0.25
33 – 37	0.5
38 - 43	1

- Rounded IV Adrenaline doses are provided as a guide only
- The Australian Resuscitation Council, Section 13 -Neonatal Guidelines, 2010 [accessed 2011 September 5, <u>http://www.resus.org.au/]</u> recommends IV Adrenaline 0.01-0.03 mg/kg (equates to 0.1-0.3 mL/kg)
- Adrenaline can be repeated every 2-3 minutes if HR is less than 60 beats per minute
- If given via the less preferred ETT route, Adrenaline dosages should be three times higher

# Appendix D: Laryngeal mask

Figure 1. Laryngeal mask



Source: Australian Resuscitation Council, Guideline 13.5, Tracheal intubation and ventilation of the newborn infant, 2010 Republished with permission of the Australian Resuscitation Council

The following laryngeal mask (LM) instructions are quoted (with the exception of the replacement of abbreviation LMA with LM) from the Australian Resuscitation Council, Guideline 13.5 – Tracheal intubation and ventilation of the newborn infant [accessed 2011 August 30, <a href="http://www.resus.org.au/policy/guidelines/section\_13/13\_5.htm">http://www.resus.org.au/policy/guidelines/section\_13/13\_5.htm</a>].

- 1. Test inflation of the cuff (using a syringe and 4 mL of air), and then slowly and fully deflate the LM cuff before insertion.
- 2. Lubricate the back and sides of the LM with water-soluble lubricant or baby's saliva, avoiding excess lubricant round the anterior surface of the cuff or in the bowl of the mask
- 3. Holding the LM like a pen, insert it with the open side of the cushioned mask facing forwards (towards the tongue, away from the palate). The index finger, placed inside the bowl of the mask, is used to prevent the tip from curling and to guide the mask, sliding the back of it against the hard palate and into the pharynx until resistance is felt.
- 4. The tube is then held firmly and with slight downward pressure with the other hand while the index finger is removed.
- 5. The cuff is then inflated with 4 mL of air. The tube may rise up slightly out of the hypopharynx as the mask is inflated.
- 6. A resuscitation device (bag or T-piece device) is then connected to the adaptor.

Effectiveness of ventilation should be checked using signs indicated above for endotracheal ventilation (chest wall movement, improvement in heart rate, improvement in oxygenation). In addition, the chest should be auscultated. For newborns receiving ventilation via an [*sic*] LM, the accuracy of colorimetric  $CO_2$  detectors to confirm position and seal has not been reported.

# Appendix E: Use of a narcotic antagonist

The following quotes are from the Neonatal Resuscitation Textbook 6th Edition 2011, American Heart Association/American Academy of Pediatrics, Lesson 7, page 247-248.

"Narcotics given to the laboring mother to relieve pain may inhibit respiratory drive and activity in the newborn. In such cases, administration of naloxone (a narcotic antagonist) to the newborn will reverse the effects of narcotics on the baby."

Giving a narcotic antagonist is not the correct first therapy for a baby who is not breathing. The first corrective action is positive-pressure ventilation.

The indications for giving naloxone to the baby require both of the following to be present:

- Continued respiratory depression after positive pressure ventilation has restored a normal heart rate **and**
- A history of maternal narcotic administration during labour

"After naloxone administration, continue to administer positive pressure ventilation until the baby is breathing normally. The duration of action of the narcotic often exceeds naloxone. Therefore, observe the baby closely for recurrent respiratory depression, which may necessitate ongoing respiratory support.

**Caution**: Do not give naloxone to the newborn of a mother who is suspected of being addicted to narcotics or is on methadone maintenance. This may result in the newborn having seizures."

Other drugs given to the mother, such as magnesium sulphate or non-narcotic analgesics or general anesthetics, also can depress respirations in the newborn; the effects of these drugs are not reversed by naloxone. If narcotics were not given to the mother or if naloxone does not result in restoring spontaneous respirations, transport the baby to the nursery for further evaluation and management while continuing to administer PPV and monitoring heart rate and pulse oximetry."

Naloxone hydrochloride 0.4 mg/mL			
Indication	Persistent respiratory depression (after positive pressure ventilation has been provided) in a newborn whose mother has received an opiate analgesic in the 4 hours before birth		
Route	IV (including UVC) preferred Intramuscular (IM) acceptable but delayed onset of action		
Method of administration	IV bolus followed by 0.9% Sodium Chloride flush		
Dose	0.1 mg/kg = 0.25 mL/kg of 0.4 mg/mL stock solution		
Frequency	Duration of action of the narcotic can exceed that of naloxone, necessitating repeated doses of naloxone Recurrent respiratory depression may necessitate repeated doses of naloxone, or consideration of alternative diagnoses		

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