# **Queensland Clinical Guidelines**

Translating evidence into best clinical practice

## Maternity and Neonatal Clinical Guideline

Newborn baby assessment (routine)



Document title: Newborn baby assessment (routine)

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The document supplement details development processes and implementation Document supplement:

activities, and is integral to and should be read in conjunction with this guideline.

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#### Acknowledgement

The Department of Health respectfully acknowledges the Traditional Owners and Cultural Custodians of the lands, waters and seas across Queensland. We pay our respects to Elders past and present, while recognising the role of current and future leaders in shaping a better health system.

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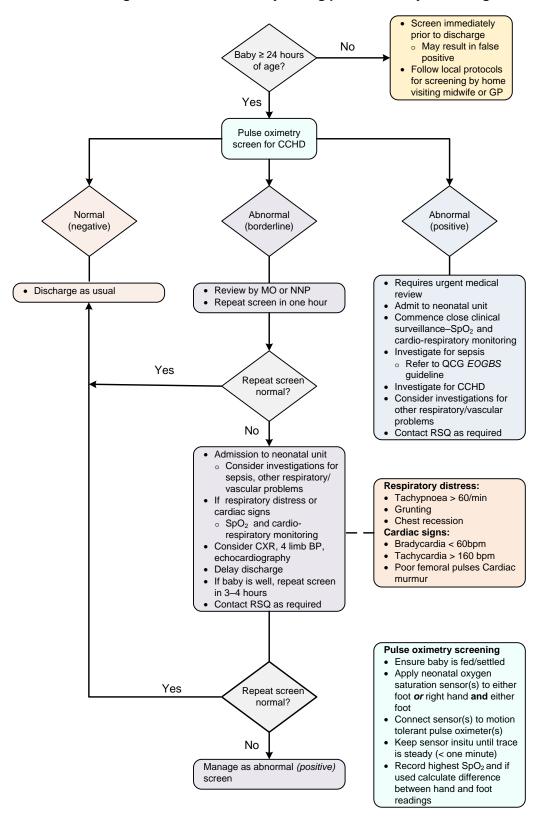
#### Flow Chart: Routine newborn baby assessment

#### Preparation Assessment Further investigation 🗹 Ur Skin colour, integrity, **Growth and appearance** Family centred care perfusion · Consider cultural needs · Dysmorphic features General State of alertness • Discuss with parents: purpose, Excessive weight loss process, timing and limitations of appearance Activity, range of ☑ Jaundice < 24 hours of age spontaneous movement ☑ Central cyanosis assessments Posture, muscle tone · Ask about parental concerns • Petechiae new/unrelated to birth • Pallor, haemangioma · Encourage participation Chart head circumference, Head and neck Timina Growth length, weight on centile ☑ Enlarged/bulging/sunken fontanelle · Initial exam immediately after birth status Macro/microcephaly and any resuscitation ☑ Subgaleal haemorrhage · Full and detailed assessment · Head shape, size Caput, cephalhaematoma within 48 hours and always prior Scalp, fontanelles, sutures Fused sutures to discharge · Facial palsy/asymmetry on crying • Eye size, position structure • Follow-up 5-7 days and 6 weeks • Hazy, dull cornea; congenital cataract • If unwell/premature-stage as Head, face, • Nose, position, structure Absent red eye reflex clinically indicated neck Ear position, structure Mouth, palate, teeth, gums tongue, frenulum Pupils unequal/dilated/constricted **Review history** clinical judgement Purulent conjunctivitis/yellow sclera Maternal medical/obstetric/social ☑ Nasal obstruction Jaw size and family • Dacryocyst; cleft lip/palate · Current pregnancy Unresponsive to noise • Length, proportions, Labour and birth Shoulders, Absent ear canal or microtia • Sex, gestational age, Apgar arms, hands • Structure, number of digits Ear drainage scores and resuscitation Small receding chin/micrognathia exhaustive. Use · Since birth-medications, Neck masses, swelling, webbing observations, feeding • Size, shape, symmetry, Swelling over or fractured clavicle movement Environment-consider: Upper limbs Breast tissue, nipples · Warmth, lighting Chest · Limb hypotonia, contractures, palsy • Heart sounds, rate, pulses · Correct identification Palmar crease pattern · Breath sounds, resp rate ğ Infection control precautions Chest Pulse oximetry Privacy are ☑ Respiratory distress Equipment-prepare: ☑ Apnoeic episodes urgent follow-up Size, shape, symmetry · Abnormal HR, rhythm, regularity · Overhead warmer if required Stethoscope Palpate liver, spleen, · Heart murmurs Abdomen kidneys Ophthalmoscope ☑ Weak or absent pulses Umbilicus Tongue depressor & glove ☑ Positive pulse oximetry Abdomen Pencil torch investigation and/or • Male-penis, foreskin, • Tape measure, infant scales, ☑ Organomegaly testes ☑ Gastrochisis/exomphalos growth charts Female-clitoris, labia, ☑ Bilateral undescended testes Pulse oximeter Genitourinary hymen Documentation **☑** Bilious vomiting Anal position, patency o Infant Personal Health Record • Inguinal hernia · Passage of urine and stool o Medical record Signs of umbilical infection o Neonatal clinical pathway Genitourinary ndications for further ☑ No urine/meconium in 24 hours · Ortolani and Barlow's ☑ Ambiguous genitalia Hips, legs, manoeuvres Discharge ☑ Testicular torsion feet • Leg length, proportions, • Hypospadias, penile chordee symmetry and digits Review discharge criteria micropenis, hydrocele · Observations, feeding, output Hips, legs and feet • Vitamin K · Risk factors for hip dysplasia · Hepatitis B vaccination · Spinal column, skin • Positive/abnormal Barlow's and/or Discuss Back Symmetry of scapulae, Ortolani manoeuvres buttocks • If < 24 hours of age, when to seek · Contractures/hypotonia urgent medical assistance Talipes · Routine screening (e.g. hearing, · Developmental hip dysplasia Behaviour, posture NBST, pulse oximetry) **Back** Muscle tone, spontaneous · Childhood immunisation program Curvature of spine movements Neurological · Support agencies Non-intact spine Crv · Newborn care Tufts of hair/dimple along intact spine • Reflexes-Moro, suck, · Health promotion Neurological grasp Medications as indicated Weak/irritable/absent cry · Personal Health Record (red Absent/exaggerated reflexes Discuss findings with book) No response to consoling **Discuss** parents Referral and follow-up ☑ Seizures Document in health Document o Routine 5-7 days & 6 weeks ☑ Altered state of consciousness Refer record(s)

Refer as indicated

Flowchart: F21.4-1-V6-R26

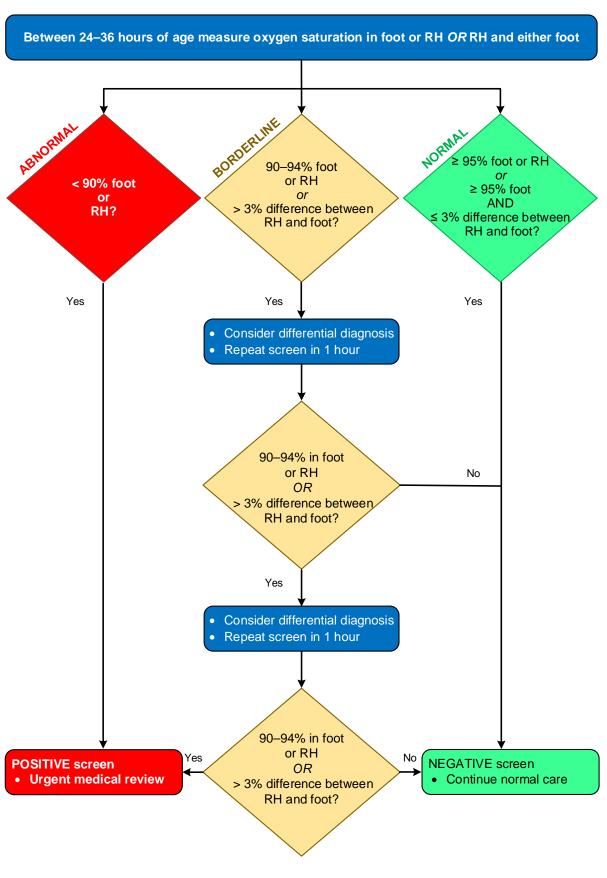
Flow Chart: Management of newborn baby during pulse oximetry screening for CCHD



CCHD: critical congenital heart disease; BP: blood pressure, bpm: beats per minute, CXR: chest X-ray, EOGBS: early onset Group B Streptococcus, GP: general practitioner; HR: heart rate, min: minute, MO: medical officer, NNP: neonatal nurse practitioner, QCG: Queensland Clinical Guidelines, RSQ: Retrieval Services Queensland, SpO₂: oxygen saturation, >: greater than; ≥: greater than or equal to; <: less than; ≤ less than or equal to

Flowchart: F21.4-2-V1-R26

Flow Chart: Pulse oximetry screening of newborn baby for critical congenital heart disease



**!RH:** right hand, >: greater than; ≥:greater than or equal to; <: less than; ≤: less than or equal to

#### **Table of Contents**

Abbreviations	7
Definitions	7
1 Introduction	8
1.1 Clinical standards	8
2 Initial examination and assessment after birth	
3 Full and detailed newborn baby assessment	
3.1 Principles of assessment	
3.2 Preparation for the full and detailed newborn assessment	
3.2.1 History	
3.2.2 Assessment preparation	
3.3 Physical examination	
3.4 Isolated abnormalities	
4 Consultation and follow-up	
5.1 Screening newborn baby	
5.2 Pulse oximetry screening	
5.2.1 Pulse oximetry results	
5.3 Discharge preparation	
5.4 Health promotion	
References	
Acknowledgements	24
List of Tables	
Table 1. Clinical standards	
Table 2. Initial examination and assessment	
Table 3. Principles of assessment	
Table 4. History Table 5. Assessment preparation	
Table 6. Newborn baby examination	
Table 7. Suggested follow-up actions	
Table 8. Screening newborn baby	
Table 9. Pulse oximetry screening	
Table 10. Pulse oximetry results	
Table 11. Discharge preparation	
Table 12. Health promotion	21

#### **Abbreviations**

CCHD	Critical congenital heart disease
GP	General practitioner
NBST	Newborn bloodspot screening test
NEWT	Newborn early waring tool
SpO <sub>2</sub>	Peripheral oxygen saturation
SUDI	Sudden and unexpected death in infancy

#### **Definitions**

Family centred care	Is an approach to the planning, delivery and evaluation of health care that is grounded in mutually beneficial partnerships among health care providers, patients and families; and incorporates the core concepts of respect and dignity, information and sharing, participation and collaboration. <sup>1,2</sup>
Newborn baby	A recently born infant in the first minutes to hours following birth. <sup>3</sup>
Neonatal unit	In this document 'neonatal unit' may be interpreted to mean neonatal observation or stabilisation area, unit or equivalent as per local terminology.
Newborn baby assessment	In this document 'newborn baby assessment' is a broad term referring to the assessment of the newborn occurring at various points in time within the first 6–8 weeks after birth. It includes the brief initial assessment, the full and detailed newborn assessment within 48 hours of birth and the follow-up assessments at 5–7 days and 6 weeks of age.
Urgent follow-up	Immediate and/or life-threatening health concern for the newborn requires urgent (same day) follow-up.
Positive screen	Means the screening test has detected an increased likelihood of having disease, anomaly or condition.

### 1 Introduction

Every newborn baby requires a brief physical examination within the first few minutes after birth, and then a full and detailed assessment within the next 48 hours and prior to discharge from hospital.<sup>4</sup> A follow up assessment is required after hospital discharge at five to seven days (by a general practitioner (GP) or midwife), and then at 6 weeks of age. The physical examination is the most important screen for major occult congenital anomalies.

#### 1.1 Clinical standards

Table 1. Clinical standards

Aspect	Consideration	
Model of care	<ul> <li>Adhere to the principles of family centred care when assessing the newborn baby         <ul> <li>Refer to Queensland Clinical Guideline: <u>Standard care</u><sup>5</sup></li> </ul> </li> <li>Perform the examination with at least one parent present<sup>4</sup></li> <li>Provide information to parents in a complete, unbiased and timely manner to support effective participation in their baby's care and any decision making</li> </ul>	
Clinician	<ul> <li>Develop local protocols regarding clinicians responsible for performing the newborn baby examination<sup>4</sup></li> <li>Develop local protocols for clinicians regarding:         <ul> <li>Training and skills maintenance in newborn baby assessment<sup>4,6,7</sup></li> <li>Recognising and managing variances, seeking guidance as required, and referring appropriately<sup>8</sup></li> </ul> </li> </ul>	
Documentation	Complete the baby's Personal Health Record ('red book')     Explain the booklet's purpose to the baby's parents	

### 2 Initial examination and assessment after birth

Complete the initial brief assessment immediately after birth on all babies following any resuscitation (if required). (Refer to Queensland Clinical Guideline: <u>Neonatal resuscitation</u>.<sup>6</sup>

Table 2. Initial examination and assessment

Aspect	Consideration
Brief initial exam	<ul> <li>Assess the baby for<sup>4</sup>:         <ul> <li>Successful transition to extra-uterine life</li> <li>Any obvious dysmorphic features or gross anomalies that will require immediate attention or discussion with the family</li> </ul> </li> <li>Confirm baby's sex<sup>4</sup></li> <li>Be flexible with the timing of this initial examination and assessment</li> <li>Do not restrict skin-to-skin contact and allow the healthy baby to spend time with their parents<sup>4</sup></li> </ul>
Hypoxic ischaemic encephalopathy	<ul> <li>If the baby had a perinatal event and/or acidosis assess baby for therapeutic hypothermia criteria and management as indicated</li> <li>Refer to Queensland Clinical Guideline: <a href="https://doi.org/10.1007/j.chaemic.gov/"><u>Hypoxic-ischaemic.gov/gov/</u></a> encephalopathy<sup>9</sup></li> </ul>
Newborn observations	<ul> <li>Commence newborn vigilance observations and document on appropriate newborn early warning (NEWT) tool<sup>10</sup> (dependent on service level of facility as per Clinical Services Capability Framework)<sup>10,11</sup></li> <li>Assess, record and (if indicated) action baby's vital signs:         <ul> <li>Every 15 minutes for two hours following birth—colour, position (for airway patency), respiration rate and any respiratory distress</li> <li>Within 1 hour of birth—heart rate, temperature and oxygen saturation</li> <li>Every eight (after initial 2 hours following birth)—respiratory rate, respiratory distress, temperature, heart rate, colour</li> </ul> </li> <li>If baby has signs of respiratory distress or has any oxygen requirement, monitor and record oxygen saturations</li> <li>Refer to NEWT<sup>12</sup> and Neonatal clinical pathway<sup>13</sup></li> </ul>
Hypoglycaemia	<ul> <li>If baby has risk factor(s) for hypoglycaemia commence blood glucose monitoring</li> <li>Refer to Queensland Clinical Guideline: <a href="https://doi.org/10.1007/jhp.goglycaemia-newborn"><u>Hypoglycaemia-newborn</u></a><sup>14</sup></li> </ul>
Sub-galeal	<ul> <li>Risk of sub-galeal haemorrhage (SGH) is increased after an instrumental birth<sup>15</sup> or as clinically indicated</li> <li>If attempted or actual instrumental birth or as clinically indicated refer to NEWT for scalp examination including: <sup>16</sup> <ul> <li>Inspection and palpation of the scalp and nape of the neck</li> <li>Head circumference measurements</li> </ul> </li> <li>If ballotable mass or free fluid in the scalp or nape of neck, or ear displacement–escalate care as a medical emergency</li> <li>If asymmetrical head, caput succedaneum, cephalhaematoma, new or increased bruising or abrasion–urgent follow up</li> </ul>

## 3 Full and detailed newborn baby assessment

### 3.1 Principles of assessment

Table 3. Principles of assessment

Aspect	Consideration
Purpose	<ul> <li>Identify the baby who is acutely unwell and requires urgent treatment</li> <li>Review any concerns the family have about the baby and attempt to address them</li> <li>Review any problems arising or suspected from antenatal screening, family history or labour (e.g. mental health issues, substance use, child protection issues, genetic conditions)</li> <li>Review weight, length and head circumference measurements</li> <li>Check the baby has passed urine and meconium since birth</li> <li>Recognise common neonatal problems and give advice to parents about management</li> <li>Diagnose congenital malformations and arrange appropriate management</li> <li>Discuss:         <ul> <li>Baby care and feeding</li> <li>Refer to Queensland Clinical Guideline: Establishing breastfeeding<sup>17</sup></li> <li>Vitamin K</li> <li>Childhood immunisation schedule including where indicated Bacille Calmette-Guerin (BCG) vaccination</li> <li>Refer to the National Immunisation Program Schedule<sup>18</sup></li> <li>Reducing the risk of sudden unexpected death in infancy (SUDI)</li> <li>Any other matters relevant to the newborn baby<sup>4</sup></li> <li>Explain problems such as jaundice that might not be observable in the baby but may become significant a few days or weeks later</li> <li>Refer to Queensland Clinical Guideline: Neonatal jaundice<sup>19</sup></li> <li>Convey information about local networks, services and access to members of a primary health care team</li> <li>Refer to Section 5 Discharge planning</li> <li>Inform families how they can request and negotiate additional help, advice, and support as relevant to the circumstances</li> </ul> </li> </ul>
Timing	<ul> <li>The Royal Australian College of Physicians (RACP) recommends an initial full and detailed assessment is performed within the first 48 hours after birth<sup>4</sup></li> <li>If baby is discharged home within the first 8 hours after birth, complete full assessment prior to discharge even though this is not the optimal time to detect all abnormalities</li> <li>Advise parents that certain conditions may only become evident in the first few days after birth following discharge from hospital</li> <li>Provide parents with information about local health support services (e.g. Child Health Services) prior to discharge</li> <li>Recommend a follow-up assessment<sup>4</sup> by general practitioner (GP) or midwife at 5–7 days of age</li> <li>Recommend a further assessment at around 6 weeks of age<sup>4,20</sup></li> </ul>
Unwell and/or preterm baby	<ul> <li>Stage the assessment as clinically indicated</li> <li>Recognise the impact of prematurity on the assessment findings</li> <li>Identify the requirement for additional, condition specific assessments (e.g. ophthalmology review for retinopathy of prematurity)</li> </ul>

### 3.2 Preparation for the full and detailed newborn assessment

### 3.2.1 History

Table 4. History

Aspect	Clinical assessment
Maternal history <sup>20</sup>	<ul> <li>Review medical, obstetric, social and family history, including:         <ul> <li>Maternal age, social background, mental health history, Edinburgh Postnatal Depression Score (EDPS), intimate partner violence, child safety alerts</li> <li>Chronic maternal disease and associated treatments (e.g. vitamin D deficiency, hypothyroidism) during pregnancy</li> <li>Substance use including prescription and non-prescription medications, alcohol and/or tobacco</li> <li>Refer to Queensland Clinical Guideline: Perinatal substance use: neonatal<sup>9</sup> and Perinatal substance use: maternal<sup>21</sup></li> </ul> </li> <li>Previous pregnancies including complications and outcomes (e.g. neonatal jaundice, ABO incompatibility, developmental dysplasia of the hips, genetic conditions, baby Group B Streptococcus positive [refer to Queensland Clinical Guideline: Early onset group B Streptococcal disease<sup>17</sup>)</li> </ul>
Current pregnancy <sup>20</sup>	<ul> <li>Results of pregnancy screening tests (e.g. blood group and type, serology, ultrasound scans)</li> <li>Chorionicity if twin pregnancy</li> <li>Any other diagnostic procedures (e.g. amniocentesis)</li> <li>Maternal ill health with any non-specific illnesses</li> <li>Complications such a gestational diabetes mellitus or hypertension</li> <li>Refer to Queensland Clinical Guideline: Gestational diabetes mellitus<sup>22</sup> and Hypertension and pregnancy<sup>23</sup></li> </ul>
Labour and birth <sup>20</sup>	<ul> <li>Progression of labour (e.g. onset, duration, interventions during labour, duration of rupture of membranes, maternal temperature, third stage)</li> <li>Evidence of abnormal fetal status in labour (e.g. cord blood gases, lactate, meconium liquor)         <ul> <li>Refer to Queensland Clinical Guideline: Intrapartum fetal surveillance<sup>14</sup></li> </ul> </li> <li>Maternal drugs in labour (e.g. antibiotics)</li> <li>Presentation and mode of birth</li> <li>Apgar scores and resuscitation at birth</li> <li>Medication since birth (e.g. vitamin K, hepatitis B immunoglobulin/vaccine, antibiotics)</li> </ul>
Gestational age <sup>24</sup>	If discrepancy with known maternal dates perform a formal gestational age assessment <sup>4</sup>
Observations since birth <sup>20</sup>	<ul> <li>Initial observations and NEWT<sup>12</sup></li> <li>Axillary temperature</li> <li>Weight</li> <li>Urine/meconium</li> <li>Relevant tool to assess substance withdrawal (if relevant)         <ul> <li>Refer to Queensland Clinical Guideline: <u>Perinatal substance use:</u> neonatal<sup>9</sup></li> </ul> </li> </ul>
Feeding since birth <sup>20</sup>	<ul> <li>Mode of feeding</li> <li>Suck behaviour</li> <li>Feeding since birth</li> <li>Refer to Queensland Clinical Guideline: <u>Establishing breastfeeding</u><sup>17</sup></li> </ul>

### 3.2.2 Assessment preparation

Table 5. Assessment preparation

Aspect	Clinical assessment
Explanation	<ul> <li>Explain the purpose, procedure and limitations of the assessment</li> <li>Ask the baby's name and confirm sex</li> <li>Ask about any concerns and provide opportunity for questions and answers</li> <li>Discuss feeding choice and progress</li> <li>Explain normal weight loss after birth (1–2% of body weight per day up to maximum 10% weight loss at day 5)</li> <li>Provide further information as required</li> </ul>
Environment <sup>25</sup>	<ul> <li>Provide:</li> <li>Adequate warmth and lighting</li> <li>Privacy for discussion about sensitive family/health issues<sup>26</sup></li> </ul>
Equipment <sup>25</sup>	<ul> <li>Overhead warmer if required</li> <li>Stethoscope</li> <li>Ophthalmoscope</li> <li>Pencil torch</li> <li>Tongue depressor and glove</li> <li>Tape measure</li> <li>Infant scales and growth charts</li> <li>If indicated, bilirubinometer and jaundice nomograms</li> <li>Refer to Queensland Clinical Guideline: Neonatal jaundice</li> <li>Baby's Personal Health Record ('red book')</li> </ul>

### 3.3 Physical examination

Use a systematic approach to examine the baby where possible. A recommended systematic approach is 'head to toe' and 'front to back'. Undress the baby down to the nappy as it is not possible to fully examine a dressed baby for all abnormalities. Assess the baby when they are quiet and settled, and do not interrupt breastfeeding.<sup>20</sup>

Table 6 includes aspects of the clinical assessment and possible indications for further investigation or follow up. Indications for urgent follow-up are identified but the list is not exhaustive. Use clinical judgement when determining the need and the urgency of follow-up for all abnormal or suspicious findings. [Refer to Table 7. Suggested follow-up actions].

Table 6. Newborn baby examination

Aspect	Clinical assessment	Indications for further investigation  ☑ Urgent follow-up
General appearance <sup>20,24,27</sup>	While the baby is quiet, alert, not hungry or crying observe: Skin colour/warmth/perfusion Alert/responsive state Activity Range of spontaneous movement Posture Muscle tone	Dysmorphic features
Growth status and feeding <sup>20,24,28</sup>	Document on the age and sex appropriate percentile charts:     Weight     Length     Head circumference	<ul> <li>Less than 10th percentile or greater than 90th percentile</li> <li>Excessive weight loss (more than 10 % of birth weight)</li> </ul>
<b>Skin</b> <sup>20,27,28</sup>	<ul> <li>Colour</li> <li>Trauma</li> <li>Congenital or subcutaneous skin lesions</li> <li>Oedema</li> </ul>	<ul> <li>✓ Any jaundice at less than 24 hours of age</li> <li>✓ Central cyanosis</li> <li>Petechiae not fitting with mode of birth, or newly appearing or associated with purpura</li> <li>Pallor</li> <li>More than 3 café-au-lait spots in a Caucasian, more than 5 in a black African newborn baby</li> <li>Multiple haemangioma</li> <li>Haemangioma on nose or forehead (in distribution of ophthalmic division of trigeminal nerve)</li> <li>Haemangioma or other midline skin defect over spine</li> <li>Oedema of feet (consider Turner syndrome)</li> </ul>
Head <sup>20,24</sup>	<ul> <li>Shape and symmetry</li> <li>Scalp</li> <li>Anterior and posterior fontanelle</li> <li>Sutures</li> <li>Scalp lesions/swelling/bruising/lacerations</li> </ul>	<ul> <li>Striction</li> <li>Enlarged, bulging or sunken fontanelle</li> <li>Microcephaly (less than 2nd percentile)/macrocephaly (greater than 98th percentile)</li> <li>Sub-galeal haemorrhage</li> <li>Caput/cephalhaematoma (consider potential for developing jaundice)</li> <li>Fused sutures</li> </ul>

Table 7. Newborn baby examination continued

Aspect	Clinical assessment	Indications for further investigation
	Symmetry of structure, features and movement	Urgent follow-up     Asymmetry on crying
	Eyes     Size and structure     Position in relation to the nasal bridge     Red reflex	<ul> <li>Hazy, dull cornea</li> <li>Absent red reflex</li> <li>Unequal, dilated or constricted pupils</li> <li>Purulent conjunctivitis</li> <li>Yellow sclera</li> <li>Congenital cataracts</li> </ul>
	Nose     Position and symmetry of the nares and septum	<ul> <li>Nasal flaring</li> <li>Nasal obstruction especially if bilateral</li> <li>Dacryocyst</li> </ul>
Face <sup>20,24</sup>	<ul> <li>Mouth</li> <li>Size, symmetry and movement</li> <li>Shape and structure</li> <li>Gums and teeth (if present)</li> <li>Lips</li> <li>Palate (hard and soft)</li> <li>Tongue and frenulum</li> </ul>	<ul><li>Cleft lip and/or palate</li><li>Mouth drooping</li></ul>
	Ears     Position     Structure including patency of the external auditory meatus     Well-formed cartilage	<ul> <li>Unresponsive to noise</li> <li>Absent external auditory canal or microtia</li> <li>Drainage from ear</li> </ul>
	Jaw size and shape	Small receding chin or micrognathia (e.g. Pierre Robin syndrome)
Neck <sup>20,24,28</sup>	<ul><li>Structure and symmetry</li><li>Range of movement</li><li>Thyroid gland or other masses</li></ul>	<ul><li>Masses/swelling</li><li>Neck webbing</li></ul>
Shoulders, arms and hands	<ul> <li>Length</li> <li>Proportions</li> <li>Symmetry</li> <li>Structure and number of digits</li> </ul>	<ul> <li>Swelling over clavicle/fractured clavicle</li> <li>Hypotonia</li> <li>Palsy (e.g. Erb's palsy, Klumpke's paralysis)</li> <li>Contractures</li> <li>Palmar crease pattern</li> </ul>
Chest, cardio- respiratory <sup>20,24,27,28</sup>	Chest Chest size, shape and symmetry Breast tissue Number and position of nipples Respiratory Chest movement and effort with respiration Respiratory rate Breath sounds  Cardiac Pulses—brachial and femoral Skin colour/perfusion Heart rate Heart rhythm Heart sounds	<ul> <li>Small, malformed or asymmetry</li> <li>Widely spaced nipples (e.g. Turner's syndrome)</li> <li>✓ Signs of respiratory distress</li> <li>✓ Apnoeic episodes</li> <li>Variations in rate, rhythm or regularity</li> <li>Murmurs</li> <li>Central cyanosis/mottling</li> <li>✓ Weak or absent pulses</li> <li>✓ Positive pulse oximetry</li> </ul>

Table 7. Newborn baby examination continued

Aspect	Clinical assessment	Indications for further investigation
, topoot	Shape and symmetry	<ul><li>☑ Urgent follow-up</li><li>☑ Organomegaly</li></ul>
Abdomen <sup>20,27,28</sup>	<ul> <li>Palpate for enlargement of liver, spleen, kidneys and bladder</li> <li>Bowel sounds</li> <li>Umbilicus including number of arteries</li> <li>Tenderness</li> </ul>	<ul> <li>✓ Gastroschisis/exomphalos</li> <li>✓ Bilious vomiting</li> <li>Inguinal hernia</li> <li>Erythema or swelling at base of umbilicus onto anterior abdominal wall</li> </ul>
Genitourinary <sup>20,27,28</sup>	Urine passed including colour and amount  Male genitalia Penis including foreskin Testes (confirm present bilaterally and position of testes) including any discolouration Scrotal size and colour Other masses such as hydrocele Female genitalia (discuss pseudomenses) Clitoris Labia Hymen	<ul> <li>✓ No urine passed within 24 hours</li> <li>✓ Ambiguous genitalia</li> <li>✓ Bilateral undescended testes</li> <li>✓ Testicular torsion</li> <li>Hypospadias, penile chordee</li> <li>Penile torsion greater than 60%</li> <li>Micropenis (stretched length less than 2.5 cm)</li> <li>Unequal scrotal size or scrotal discolouration</li> <li>Testes palpable in inguinal canal</li> </ul>
Anus <sup>20,27,28</sup>	<ul><li>Meconium passed</li><li>Anal position</li><li>Anal patency</li></ul>	✓ No meconium passed within 24 hours
Hips, legs and feet <sup>20,27,28</sup>	<ul> <li>Use Barlow and Ortolani manoeuvres to examine hips</li> <li>A firm surface is necessary</li> <li>Assess legs and feet for <ul> <li>Length</li> <li>Proportions</li> <li>Symmetry</li> <li>Structure and number of digits</li> </ul> </li> </ul>	<ul> <li>Risk factors for hip dysplasia: breech presentation, fixed talipes, fixed flexion deformity, asymmetrical buttock creases, severe oligohydramnios, first degree relative with developmental hip dysplasia</li> <li>Positive Barlow and/or Ortolani test</li> <li>Hypotonia/contractures</li> <li>Positional talipes</li> </ul>
Back <sup>20,27,28</sup>	Spinal column     Scapulae and buttocks for symmetry     Skin	<ul> <li>Curvature of spine</li> <li>Non-intact spine</li> <li>Tufts of hair or dimple along intact spine</li> <li>Sacral pit without visible intact base</li> </ul>
Neurologic <sup>20,27,28</sup>	Observe throughout:     Behaviour     Posture     Muscle tone     Movements     Cry     Examine reflexes     Moro     Suck     Grasp	<ul> <li>Weak, irritable, high pitched cry</li> <li>No cry</li> <li>Does not respond to consoling</li> <li>Absent/exaggerated reflexes</li> <li>Seizures</li> <li>Altered state of consciousness</li> </ul>

#### 3.4 Isolated abnormalities

The following anomalies are usually of no concern when isolated (3 or more such abnormalities are of concern)

- Folded-over ears
- Hyperextensibility of thumbs
- Syndactyly of second and third toes
- · Single palmar crease on one hand
- Polydactyly, especially if familial
- Single umbilical artery
- Hydrocele
- Fifth finger clinodactyly
- Simple sacral dimple just above the natal cleft (less than 2.5 cm from anus and less than 5 mm wide)
- Single café-au-lait spot
- Slate grey naevi/congenital dermal melanocytosis (Mongolian spot)
- Single ash leaf macule
- · Third fontanelle
- · Capillary haemangioma apart from those described in table above
- Accessory nipples

### 4 Consultation and follow-up

Use clinical judgement to determine the appropriate urgency of follow-up in the context of abnormal or suspicious findings arising from a newborn baby assessment. If there is uncertainty about the urgency of follow-up seek advice from a more senior clinician.

Table 7. Suggested follow-up actions

Category	Follow-up action	
✓ Urgent Immediate and/or life-threatening health concern for the newborn baby	Arrange same day (as soon as possible) medical review     If baby is already discharged from hospital arrange review by either:	
Follow-up Existing and/or potential health concern for the newborn baby	transport requirements)  Determine the urgency of the follow-up required  Consider the need for: Consultation with senior practitioners (e.g. review of baby, telephone consultation about findings, telehealth consultation) Additional immediate investigation(s) (e.g. blood test) Referral for specialist review (e.g. cardiology) Re-assessment or recheck at 6 week assessment (or sooner as indicated) Distribution of written summary information (e.g. GP, referring hospital) Advise parents/family of clinical concerns and the importance of review and follow-up arrangements Provide verbal/written information as appropriate Consider parental support needs (e.g. social work involvement, transport requirements)	

### 5 Discharge planning

Evaluate each mother-baby dyad individually and involve the family when determining optimal time of discharge. Criteria for newborn baby discharge include physiological stability, family preparedness to provide care for the baby at home, availability of social support, and access to the health care system and community resources.<sup>29</sup>

### 5.1 Screening newborn baby

Table 8. Screening newborn baby

Aspect	Consideration
Context	<ul> <li>Explain to baby's parents the importance, and how to access screening of the newborn baby</li> <li>If possible complete all screening before discharge</li> <li>Organise repeat screening and/or follow up as required</li> </ul>
Hearing <sup>30</sup>	<ul> <li>Healthy hearing screening is performed in the birthing hospital prior to discharge</li> <li>May be provided as outpatient up to 3 months of age<sup>30</sup> <ul> <li>Refer to local protocols</li> </ul> </li> </ul>
Jaundice	<ul> <li>Consider checking TcB (transcutaneous bilirubin) prior to discharge</li> <li>Follow local protocols</li> <li>Refer to Queensland Clinical Guideline: Neonatal jaundice<sup>19</sup></li> </ul>
CCHD	Refer to Table 9. Pulse oximetry screening
Newborn bloodspot screening test (NBST)	<ul> <li>Conditions screened are added as screening technology and treatments develop, and in line with national recommendations to maintain interjurisdictional consistency</li> <li>Screens for         <ul> <li>Galactosaemia</li> <li>Phenylketonuria</li> <li>Primary hypothyroidism</li> <li>Congenital adrenal hyperplasia</li> <li>Cystic fibrosis</li> <li>Spinal muscular atrophy (SMA)</li> <li>Severe combined immune deficiency (SCID)</li> <li>A range of rare disorders of amino acid, organic acid and fatty acid metabolism</li> </ul> </li> <li>Refer to Queensland Clinical Guideline: Newborn bloodspot screening<sup>31</sup></li> </ul>

### 5.2 Pulse oximetry screening

The Queensland Maternal and Perinatal Quality Council recommends all newborn babies are offered pulse oximetry screening for critical congenital heart disease (CCHD).<sup>32</sup> Optimal detection of CCHD occurs when pulse oximetry screening is used in conjunction with an antenatal ultrasound and a physical examination of the baby.<sup>33</sup>

Table 9. Pulse oximetry screening

Aspect	Consideration
Context	<ul> <li>Congenital heart disease (CHD) occurs in nearly 1% of live births, approximately one quarter of these will be CCHD<sup>34</sup></li> <li>If CCHD is undetected newborn babies are at risk for death in the first few days or weeks of life<sup>34</sup></li> <li>Pulse oximetry:         <ul> <li>Non-invasive screening technology to assist in detecting hypoxemia, a clinical sign of critical congenital heart disease (CCHD)<sup>20,29</sup></li> <li>Can detect some CCHD that would otherwise be missed on routine examination or antenatal ultrasound</li> <li>Can also identify non-cardiac problems such as sepsis and respiratory problems, and these are common causes of a positive screen</li> <li>Cochrane Review<sup>35</sup> found pulse oximetry:</li></ul></li></ul>
Target population	All healthy term and late preterm newborn babies <sup>33</sup>
Equipment	<ul><li>Motion tolerant pulse oximeter</li><li>Neonatal oxygen saturation sensor</li></ul>
Timing	<ul> <li>Screen between 24 and 36 hours of age as<sup>33</sup>:         <ul> <li>Prior to 24 hours of age is likely to result in increased false positive result<sup>34</sup> or</li> <li>After 48 hours delays early intervention if required</li> </ul> </li> <li>If baby is less than 24 hours of age at time of discharge, screen immediately prior</li> <li>Consider screening by home visiting midwife<sup>36</sup></li> </ul>
Protocol	<ul> <li>Baby has been fed and is settled</li> <li>Site the sensor and then connect to motion-tolerant monitor(s)<sup>37</sup>:</li> <li>Site either: <ul> <li>On either foot<sup>34,35</sup> (postductal) or</li> <li>Right hand (pre-ductal) and either foot (postductal)</li> </ul> </li> <li>Keep sensor(s) insitu until a steady trace is obtained<sup>37</sup> then remove (usually less than one minute)</li> <li>Document-highest oxygen saturation result during the screen and if used difference between sites</li> </ul>

### 5.2.1 Pulse oximetry results

Table 10. Pulse oximetry results

Aspect	Consideration
Saturation greater than or equal to 95% (normal)	<ul> <li>Negative pulse oximetry screen—maximum oxygen saturation:         <ul> <li>Greater than or equal to 95% on either foot<sup>24,37</sup> or</li> <li>Greater than or equal to 95% on either foot or right hand, and a difference between right hand and either foot less than or equal to 3%<sup>33</sup></li> </ul> </li> <li>Baby suitable for discharge (in accordance with other discharge criteria)</li> </ul>
Saturation 90–94%	<ul> <li>Borderline pulse oximetry screen-maximum oxygen saturation:         <ul> <li>90–94% on either foot<sup>37</sup> or</li> <li>90–94% right hand and/or a difference between right hand and either foot greater than 3%<sup>33</sup></li> </ul> </li> <li>Medical review indicated<sup>24,37</sup></li> <li>Consider investigation of other causes including respiratory/vascular problems (e.g. sepsis, respiratory distress syndrome, lung malformations, persistent pulmonary hypertension of the newborn)-monitor as indicated</li> <li>Repeat screen in one hour<sup>37</sup> <ul> <li>Consider using different oximeter)</li> </ul> </li> <li>If oxygen saturations remain borderline repeat the screen in in one hour Consider right hand and either foot simultaneously</li> <li>If repeat screen abnormal, specialist medical review indicated:         <ul> <li>Delay discharge and admit to neonatal unit</li> <li>Consider chest X-ray and four limb blood pressure</li> <li>Consider referral for echocardiography<sup>24</sup></li> </ul> </li> <li>Contact RSQ for advice from a neonatologist or paediatric cardiologist</li> </ul>
Saturation less than 90% (abnormal)	<ul> <li>Positive pulse oximetry screen: maximum oxygen saturation during recording is less than 90%<sup>37</sup> on either foot or right hand<sup>33</sup></li> <li>Admit to neonatal unit–requires urgent specialist medical review</li> <li>Consider chest X-ray and four limb blood pressure</li> <li>Commence close clinical surveillance (e.g. continuous monitoring including oximetry)</li> <li>Investigate for neonatal sepsis         <ul> <li>Refer to Queensland Clinical Guideline: Early onset Group B Streptococcal disease<sup>38</sup></li> </ul> </li> <li>Consider investigations for CCHD including an echocardiogram<sup>34,37</sup> Consider investigation of other causes including respiratory/vascular problems (e.g. respiratory distress syndrome, lung malformations, persistent pulmonary hypertension of the newborn)<sup>34,39</sup></li> <li>Contact RSQ for advice from a neonatologist or paediatric cardiologist</li> </ul>

### 5.3 Discharge preparation

Table 11. Discharge preparation

Aspect	Considerations
Discharge criteria	Review the baby's status prior to discharge including <sup>13</sup> :     Feeding–suck feeding adequately     Newborn baby observations–temperature maintenance, respiratory rate, BGL (if measured), weight     Urine and stool passage     Completion of newborn baby assessment     Vitamin K status–if required give script and education for further oral vitamin K     Hepatitis B vaccination (and if indicated, hepatitis B immunoglobulin) administered–advise parents about next dose due
Discharge at less than 24 hours of age	If discharged at less than 24 hours of age, advise parents to seek urgent medical assistance if baby:  Has not passed meconium within 24 hours of birth  Appears jaundiced within first 24 hours after birth  Has elevated temperature  Vomiting  Difficulty feeding  Lethargic  Has decreased urine output or stools  Respiratory signs  Rash  Provide relevant Queensland Clinical Guidelines consumer information sheets (e.g. breastfeeding, jaundice)  Advise parents about the importance of follow-up assessments at:
Referral and follow-up	<ul> <li>Five to seven days of age</li> <li>Six weeks of age</li> <li>Arrange referral for a newborn baby and/or family with identified problems</li> <li>Document arrangements and inform family</li> <li>If required provide scripts or medication (e.g. vitamin D) for use after discharge</li> <li>Provide discharge information to the GP</li> </ul>
Documentation	<ul> <li>Plot anthropometric parameters on growth charts</li> <li>Complete the baby's Personal Health Record booklet</li> <li>Relevant sections complete before discharge</li> <li>Explain parental use and completion after discharge</li> <li>Document completion of the newborn assessment and associated discussions, findings and follow-up requirements in the medical record</li> </ul>
Support agencies	<ul> <li>Provide information on the role of and how to access relevant support agencies (including but not limited to):         <ul> <li>Child health service</li> <li>GP</li> <li>Child/community Health/health worker</li> <li>Midwife (e.g. home visiting, drop-in clinics, group practice, eligible or private)</li> <li>Lactation consultant</li> <li>Australian Breastfeeding Association</li> <li>13HEALTH (13 43 25 84) telephone help line</li> </ul> </li> <li>Psychological support agencies (e.g. perinatal mental health services; PANDA; groups for fathers)</li> </ul>

### 5.4 Health promotion

Discuss relevant parenting and health education topics with parents/carers prior to discharge, 8,29 and provide relevant Queensland Clinical Guidelines parent information sheets.

Table 12. Health promotion

Newborn baby care	<ul> <li>Hygiene <ul> <li>Bathing</li> <li>Eye and mouth care</li> <li>Umbilical cord care</li> </ul> </li> <li>Nappy changing including skin care</li> <li>Feeding <ul> <li>Feeding cues and behaviour</li> <li>Promote and support breastfeeding</li> <li>Refer to Queensland Clinical Guideline: Establishing breastfeeding<sup>17</sup></li> </ul> </li> <li>Growth and weight gain</li> <li>Sleep patterns</li> <li>Normal bowel and urine patterns</li> <li>Detection and management of jaundice <ul> <li>Refer to Queensland Clinical Guideline: Neonatal jaundice<sup>19</sup></li> </ul> </li> <li>Sun protection<sup>40</sup></li> <li>Prevention of hypothermia and hyperthermia</li> </ul>
Warning signs of illness <sup>41</sup>	Discuss and advise when to seek medical help, e.g.:     Raised temperature     Poor feeding     Vomiting     Irritability, lethargy     Decreased urine or stools     Jaundice     Respiratory signs     Rash
SUDI <sup>42</sup>	<ul> <li>Provide written information about safe infant care to reduce the risk of sudden unexpected death in pregnancy (SUDI)</li> <li>Discuss safe positioning of baby during breastfeeding</li> <li>Discuss safer infant sleeping:         <ul> <li>Positioning and settling</li> <li>Swaddling</li> <li>Sharing sleep surfaces</li> <li>Appropriate clothing</li> </ul> </li> <li>Risks from smoking–provide advice about smoking cessation</li> <li>Refer to Queensland Clinical Guideline: <u>Safer infant sleep</u><sup>42</sup></li> </ul>
Injury prevention	<ul> <li>Car capsule to transport baby</li> <li>Pram safety</li> <li>Reducing home hazards</li> <li>Hip healthy swaddling and use of slings<sup>43</sup></li> </ul>
schedule	<ul> <li>Provide information about childhood immunisation program</li> <li>Refer parents to baby's Personal Health Record ('red book')</li> </ul>

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