





Transcutaneous Monitoring in Neonates

Sites where Local Guideline and Procedure Neonatal Intensive Care Unit (NICU) JHCH applies

This Local Guideline and Procedure applies to:

1. Adults No 2. Children up to 16 years No 3. Neonates - less than 29 days Yes

Target audience All clinicians caring for infants in NICU

Description Provides guidance to neonatal clinicians for the use and management of transcutaneous monitoring in infants

Go to Guideline

Keywords NICU, SCU, JHCH, neonate, newborn, transcutaneous, TCM,

carbon dioxide, CO2 monitoring

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Replaces existing document? Yes

Transcutaneous Oxygen/Carbon Dioxide Monitoring in Registration number and dates of

Neonates JHCH NICU 12.05 superseded documents

Related Legislation, Australian Standard, NSW Ministry of Health Policy Directive or Guideline, National Safety and Quality Health Service Standard (NSQHSS) and/or other, HNE Health Document, Professional Guideline, Code of Practice or Ethics:

NSW Health Policy Directive PD2017 013 Infection Prevention and Control Policy

NSW Health Policy Directive PD2017_032 Clinical Procedure Safety

NSW Health Policy Directive PD2020 020: Incident Management Policy

NSW Health Policy Directive PD2014_007 Pressure Injury Prevention and Management

Position responsible for and Jason Simpson, General Manager/Director of Nursing, JHCH/CYPFS document authorised by

Natalie Butchard, Manager Newborn Services, NICU JHCH Contact person

Contact details Natalie.Butchard@health.nsw.gov.au

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Version 3 Page 1 Note: Over time links in this document may cease working. Where this occurs please source the document in the PPG Directory at: http://ppg.hne.health.nsw.gov.au/

PURPOSE AND RISKS

This local clinical procedure has been developed to provide instruction to the health clinician and to ensure that the risks of harm to the child associated with application of transcutaneous monitoring (TCM) in NICU are prevented, identified and managed.

The risks are:

- Epidermal damage
- Pressure injury
- Skin burns

The risks are minimised by:

- Clinicians having knowledge of transcutaneous carbon dioxide monitoring implementation and management
- Following the instructions set out in this document
- Recognition of the common clinical signs of the epidermal damage
- Notification and management of the complications/risks to the patient

Any unplanned event resulting in, or with the potential for injury, damage or other loss to infants/staff/family as a result of this procedure must be reported through the Incident Management System and managed in accordance with the NSW Health Policy Directive PD2020_020: Incident Management Policy. This would include unintended injury that results in disability, death or prolonged hospital stay.

It is mandatory for staff to follow relevant: "Five moments of hand hygiene", infection control, moving safely/safe manual handling, documentation practices and to use HAIDET for patient/carer communication: **H**and hygiene **A**cknowledge, **I**ntroduce, **D**uration, **E**xplanation, **T**hank you or closing comment.

Risk Category: Clinical Care & Patient Safety

CLINICAL PROCEDURE SAFETY LEVEL

Every clinician involved in the procedure is responsible for ensuring the processes for clinical procedure safety are followed. The following level applies to this procedure (click on the link for more information):

Level 1 procedure

CONTENT

Considerations for Use

Contraindications for Use

Equipment Required

Transcutaneous Monitor Settings

Sensor Positioning

Sensor Application

Sensor Calibration

Sensor Membrane Replacement

TRANSCUTANEOUS MONITORING SUMMARY

- TCM allows for continuous monitoring of capillary O₂ and CO₂, and adjust respiratory support accordingly
- TCM has the potential to reduce bloodletting
- TCM has skin integrity risks associated with its use (i.e. burns, pressure injury, epidermal stripping), strict observation for risks is required

GUIDELINE

While not requiring mandatory compliance, staff must have sound reasons for not implementing standards or practices set out within guidelines issued by HNE Health, or for measuring consistent variance in practice.

Introduction

Transcutaneous monitoring (TCM) is non-invasive and continuous transcutaneous oxygen and carbon dioxide measurement system. Using TCM has the potential to reduce the number of blood gases needed to manage respiratory support and/or mechanical ventilation. Provides real time alert to clinicians about potential changes in clinical condition of the infant. The sensor heats the skin, effectively increasing local perfusion so that O₂ and CO₂ can diffuse to the skin surface more easily and enabling monitoring transcutaneous partial pressure of oxygen/carbon dioxide (tcPO₂/tcPCO₂).

Considerations for Use

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Any baby where continuous CO₂ monitoring could help target respiratory support and/or reduce bloodletting. Examples include:

- Infants with significant respiratory illness/rapidly changing respiratory disease
- Unstable infants requiring mechanical ventilation/CPAP support
- Infants requiring reduction in number of blood gases
- Other infants who require additional monitoring and targeting of respiratory support

Contraindications for Use

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- Infants with compromised skin integrity
- Infants undergoing body cooling
- Infants with poor perfusion (i.e. sepsis or cardiac condition) or inotropic support (may lead to false high tcPCO₂ and falsely low tcPO₂)

Equipment Required

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- Transcutaneous monitor (see Figure 1)
- Fixation rings
- Contact gel/fluid



Figure 1: SenTec Transcutaneous Module (Image from SenTec Quick Reference Guide)

Transcutaneous Monitor Settings

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Ensure the current settings and profile are on the neonatal settings (user profile). Temperature Settings and sensor site timeframes are outlined in Table 1.

SENSOR TEMPERATURE SETTING	RECOMMENDED SENSOR SITE TIMES
41.0 °C	4 hours
43.0 °C	2 hours

Table 1: Recommended sensor temperature settings and site times

Note: Consideration when using TCM on extreme preterm infants (<29 weeks) is required.

Monitoring for any compromise to skin integrity and consider using 41° temperature setting for 2 hour time period only

Sensor Positioning

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Optimum measurement is obtained from a site that has high blood flow, capillary density and thin epidermis (avoid the nipples at all times). Sensor sites should not be used more than a single time, and fixation rings not in use should be removed at all times. Infants must never be positioned where they are laying on the sensor ring or TCM cable as this has potential to cause a pressure injury.

Ideal sensor sites are outlined in Figure 2 (Note; the infant's forehead is not used in JHCH NICU as sensor application site).

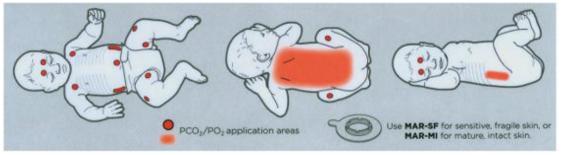


Figure 2: Neonatal Sensor Application Sites (Image from SenTec Quick Reference Guide)

Sensor Application

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- Choose relevant Multi-site Attachment Ring (MAR):
 - o MAR-MI; standard ring for mature/intact skin
 - MAR-SF; ring for more sensitive/fragile skin
- Clean site with alcohol wipe and let it dry (follow steps in Figure 3)
- Attach the ring to the selected sensor site, ensuring that the skin underneath the sensor site is not wrinkled (see Figure 3)
- Apply one small drop of contact liquid to the skin area in the centre of the ring (see Figure 3)
- Holding the sensor at its neck, approach the MAR from the flap side and first insert the nose of the sensor into the ring (see Figure 3)
- Click in the sensor by applying slight downward pressure on its neck (see Figure 3)
- Rotate the sensor in the ring into the best position and press the sensor gently against the skin to spread the contact liquid (see Figure 3)
- Verify that air gaps between the skin and the sensor are eliminated and that the sensor can easily be rotated

• After sensor application, tcPCO₂ readings stabilize within 2 to 10 minutes. After stabilization the displayed values turn from grey to green

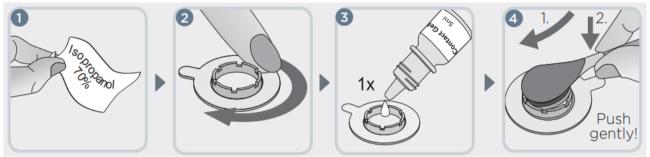


Figure 3: Sensor Application using a MAR (Image from SenTec Quick Reference Guide)

Changing the Sensor Site

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Sensor timer is set to 2 to 4 hours (see Table 1). When a sensor requires changing, a low priority alarm sounds, the message 'Site Time Elapse' will be displayed on the status bar and the 'Remaining Monitoring Time' icon turns red.

- Apply a second, new attachment ring to the patient prior to removing the sensor cable
- Remove sensor from current site and wipe clean with alcohol swab
- Insert sensor into docking station (this will calibrate sensor and also reset the 'site time')
- From this point it is ready for use again

Infants with poor skin integrity and/or poor perfusion may need more frequent sensor site changes

Sensor Calibration

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Calibration should occur in the following circumstances:

- A new monitoring period is to commence
- The membrane has been changed on the sensor
- The sensor operating temperature has been changed
- The accuracy of the measurement is in doubt (troubleshooting step)
- The monitoring site has been changed

To calibrate the TCM:

- Wipe away any excess contact gel on sensor and clean with 70% alcohol swab only
- Place sensor in docking station
- Once returned to docking station, calibration is automatic

Sensor Membrane Replacement

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The sensor membrane needs to be changed using the membrane cartridge and insert.

Sensor membrane needs to be changed in the following circumstances:

- Every 28 days
- When damaged/or missing
- There is a loose fitting membrane
- The electrode has air trapped or becomes dry
- When a sensor 12 error message appears

The sensor membrane is changed by the Technical Assistants (or nursing staff after hours when required).

Please see the following link for <u>SenTec How to Videos</u> (for available tutorials sensor application, sensor cleaning and sensor membrane change).

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IMPLEMENTATION PLAN

The clinical guideline will be:

- Circulated to Head of Department and Managers in NICU
- Circulated to the clinicians via the Children Young People and Families Network and the Women's Health and Maternity Network (where applicable)
- Made available on the intranet (PPG) and HNEKids website
- Presented at facility/unit meetings and tabled for staff to action

MONITORING AND AUDITING PLAN

- The person or leadership team approving the clinical guideline is responsible for ensuring timely and effective review of the guideline.
- Evaluation will require a review of the most current evidence as well as consideration of the experience of Neonatal staff at JHCH in the implementation of the clinical guideline.
- Data derived from monitoring and evaluation should inform the review of the clinical guideline either as required or scheduled.
- Implementation, education support and monitoring compliance be completed by local Clinical Educators and Unit Managers.
- Amendments to the guideline will be ratified by the Clinical Director and Manager of Newborn Services prior to final sign off by the JHCH.

CONSULTATION WITH KEY STAKEHOLDERS

AUTHORS: Amy May, Clinical Nurse Specialist, NICU JHCH

Michelle Stubbs, Research Nurse, NICU JHCH Jo Davis, CNC Newborn Services, JHCH

REVIEWERS: Dr Larissa Korostenski, Neonatologist, NICU JHCH

A/prof. Koert De Waal, Neonatologist, NICU JHCH Samantha Hassall, Registered Nurse, NICU JHCH Ruth Wootton, Clinical Nurse Specialist, NICU JHCH

CONSULTATION: Neonatal Team, Neonatal Intensive Care Unit, JHCH

NICU Operational Planning & Management Committee JHCH Clinical Quality & Patient Care Committee

APPROVED BY: Natalie Butchard, Manager Newborn Services, NICU JHCH

Dr Larissa Korostenski, Head of Newborn Services, NICU JHCH Jason Simpson, General Manager/Director of Nursing, CYPFS

APPENDICES

1. Glossary & Abbreviations

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- SenTec Transcutaneous Monitoring System Product Information and Website

FEEDBACK

Any feedback on this document should be sent to the Contact Officer listed on the front page.

APPENDIX 1

GLOSSARY & ABBREVIATIONS

Acronym or Term	Definition
CO ₂	Carbon Dioxide
CPAP	Continuous Positive Airway Pressure
HNELHD	Hunter New England Local Health District
JHCH	John Hunter Children's Hospital
MAR	Multi-Site Attachment Ring
MAR-MI	Multi-Site Attachment Ring – Mature/Intact (skin)
MAR-SF	Multi-Site Attachment Rings – Sensitive/fragile (skin)
NICU	Neonatal Intensive Care Unit
O ₂	Oxygen
SCU	Special Care Unit
ТСМ	Transcutaneous Monitoring
TcPCO ₂	Transcutaneous Partial Pressure of Carbon Dioxide
TcPO ₂	Transcutaneous Partial Pressure of Oxygen Dioxide