



Nursing Guidelines on the Care of a child with a Temporary External Pacemaker

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Nursing Guidelines for the Care of a Child with a Temporary External Pacemaker

1st Edition - 2008

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2nd Edition

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1. Introduction

A pacemaker is a device which uses electrical impulses to increase or regulate the heart rate and /or rhythm, when the patient's own intrinsic function of conduction or impulse generation is impaired. A temporary external pacemaker is one which is located outside the body for the purpose of regulating the heart rate and/ or rhythm for a temporary period of time. A pacemaker box is used to regulate and control the function of pacing and pacing wires are used to conduct and sense the heart's intrinsic electrical activity. Most often in paediatrics, temporary pacing wires are placed on the epicardium or in the myocardium at the conclusion of cardio-thoracic surgery or during an emergency thoracotomy in an intensive care setting. Other modes of temporary pacing are:

Transvenous – Catheter is Inserted during cardiac catheterisation i.e. femoral vein and advanced via a guide wire to right ventricle.

Transcutaneous – Emergency non-invasive pacing which may be used for severe symptomatic bradycardia. Electrode pads are placed on anterior and posterior chest to deliver stimulus through the chest wall. Available on some defibrillators.

Transoesophageal - Paces by impulse transversing tissue between the electrode in the oesophagus and left atrium. Usually short term pacing i.e. atrial pacing without A.V .block.

(Anderson 2000, Hazinski 2013).

Indications for Temporary Pacing

1. Post cardiac surgery.
 - Higher risk of arrhythmias in first 2 -3 days post surgery, especially left ventricular outflow tract, AVSD or VSD surgery.
 - Temporary support to increase cardiac output.
2. As a prelude to permanent pacing.
3. To reverse certain types of atrial or ventricular arrhythmias
4. Severe symptomatic bradycardia (Hickey and Baas 1991, Hazinski 2013).

Types of Cardiac Pacing (commonly used)

- ♦ **Demand** To sense the patient's intrinsic activity and deliver an impulse only if intrinsic electrical activity is NOT sensed within a predetermined time.
- ♦ **Fixed** To deliver an impulse at a predetermined rate regardless of intrinsic myocardial electrical activity. This type of pacing is less seldom used as it is associated with an increased risk of arrhythmias. Sensitivity needs to be turned to lowest level to avoid sensing of patient's own intrinsic activity.
- ♦ **A-V Sequential (Dual)** Delivers atrial and ventricular pacing in sequence, thereby preserving atrial – ventricular synchrony. This has the advantage of atrial kick and increase in cardiac output of appropriately 20%). (Van Orden-Wallace 2001).

Classification of Pacemaker Modes

Pacemakers are classified in a 3 letter generic code devised by the North American Society of Pacing and Electrophysiology (NASPE) and the British Pacing and Electrophysiology Group (BPEG)

(Obias-Manno 2001, Reynolds and Apple 2001, Van Orden-Wallace 2001, Bernstein *et al* 2002).

NASPE / BPEG Generic Code (Revised 2002)

CHAMBER BEING PACED	CHAMBER BEING SENSED	PACEMAKER RESPONSE
A = Atrium	A = Atrium	T = Triggered
V = Ventricle	V = Ventricle	I = Inhibited (Demand Mode)
D = Dual (Atrium & Ventricle)	D = Dual (Atrium & Ventricle)	D = Dual (Triggered / Inhibit)
O = None	O = None	O = None (Asynchrony)

(Bernstein *et al* 2002)

Mode of Response Response to the intrinsic myocardial activity.

I – Inhibit The pacemaker will not pace if it senses depolarisation, thus allowing the patients' own heart beat to maintain cardiac output.

T – Triggered If the pacemaker does not sense depolarisation.

O – No Response

The mode of pacing selected depends on the patient's inherent heart rate and rhythm and the function of the atria and ventricles. The mode used will be the one which will best optimise cardiac output. The most common temporary pacing is:

- ♦ **AAI - Atrial Pacing**
- ♦ **VVI - Ventricular Pacing**
- ♦ **DDD - A.V Sequential (Dual) Pacing**

The Pacing Circuit

Pulse Generator (Pacing Box) This contains the energy source and electrical circuitry to provide an electrical stimulus to maintain the specified rate. It also recognises and evaluates the heart's intrinsic rhythm. The pacing circuit has terminals for pacemaker wire connection of bi-polar leads. Bipolar leads measure electrical potential between 2 lead wires in contact with the heart.

Lead / Wire / Electrode This transmits the patients' rhythm to the pulse generator and also carries an electrical stimulus, between the pulse generator and the chamber being paced. The electrode needs a negative (output) pole (the tip) and a positive (ground) pole (the insulator) which enables a current to flow between the pulse generator and the heart. Epicardial wires may be placed after cardiac surgery on the epicardium or placed transvenously through guided insertion of specialised catheters at cardiac catheter (Reynolds and Apple 2001, Hazinski 2013).

2. Nursing Care of the Child with a Temporary External Pacemaker

Nurses should only care for a child with a temporary external pacemaker having received the necessary theoretical and practical instruction to practice competently, within their scope of All nursing care is given with regard to guidance for good practice practice (An Bord Altranais, 2000, OLCCH, 2002).

NB: All pacemaker settings and subsequent changes are the responsibility of the medical team ONLY and should not be changed by nursing staff.

Section 1: MONITORING	
Action	Rationale and Reference
Nurse child on cardiac monitor: observe the heart rate and rhythm with continuous ECG recordings. Assess for capture and sensing of the pacemaker. Ensure ECG rate alarm is set 10pm below the pacemaker heart rate.	To allow for evaluation of cardiac and pacemaker function. Early detection of arrhythmias or pacemaker malfunction allows for timely intervention, (Van Orden-Wallace 1998, Boyce and Rost 2000, Overbay and Criddle 2004, Fischer 2008, Hockenberry and Wilson 2011, Yorkhill Children's Hospital 2011, Hazinski 2013).
In addition, monitor heart rate from an alternative source (Pulse rate, arterial line, pulse oximetry).	NB: The presence of satisfactory heart rate on a cardiac monitor DOES NOT ensure effective cardiac contraction and cardiac output.
Monitor: <ul style="list-style-type: none"> Blood pressure Temperature. Assess for changes in responsiveness / behaviour i.e. restlessness or irritability.	To establish baseline and detect changes in a timely fashion. NB: Decreased blood pressure is a late sign of low cardiac output (Hazinski,2013). Paradoxical blood pressure changes may indicate cardiac tamponade secondary to perforated ventricle.
NB: Minimum of 4 hourly or as condition indicates	These changes may be early signs of low cardiac output. (Van Orden-Wallace 1998, Boyce and Rost 2000, Hazinski 2013).

<p>Assess tissue perfusion:</p> <ul style="list-style-type: none"> Peripheral pulses (strong or weak) Capillary refill (brisk or sluggish) Warmth of extremities <p>NB: Minimum of 4 hourly or as condition indicates</p>	<p>Tissue perfusion depends on adequate cardiac output. These are early signs of low cardiac output.</p> <p>(Boyce and Rost 2000, Hazinski 2013).</p>
<p>Assess rate and regularity of respirations.</p> <p>Monitor colour and oxygen saturations to establish parameters for same.</p> <p>Administer oxygen if ordered and clinically indicated</p>	<p>To establish baseline and detect changes in a timely fashion.</p> <p>Increased respiratory rate, dyspnoea or cough may be indications of increasing heart failure (Van Orden-Wallace 1998, Hockenberry and Wilson 2011).</p>
<p>Maintain strict fluid balance chart.</p>	<p>To provide information about fluid balance. Large positive balance and diminished urine output may indicate worsening heart failure (Van Orden-Wallace 1998, Hazinski 2013).</p>
Action	Rationale and Reference
<p>Monitor serum electrolytes (as per medical team).</p> <p>Monitor acid-base balance (as per medical team).</p> <p>Inform medical team of changes in patient's condition or laboratory findings. Document same.</p>	<p>Electrolyte imbalance may interfere with electrical activity of the heart (Reiswig-Timothy and Rodeman 2004, Hockenberry and Wilson 2011).</p> <p>Pacing thresholds can be affected by acid-base balance (Hazinski, 2013).</p> <p>To allow for timely interventions by medical team (Van Orden-Wallace 1998, An Bord Altranais 2002).</p>
<p>Assess bowel function daily.</p> <p>Prevent constipation.</p>	<p>To allow timely interventions in preventing constipation. Straining on defaecation may reduce cardiac output (Van Orden-Wallace 1998).</p>

SECTION 2: DOCUMENTATION

Action	Rationale and Reference
<p>Check pacemaker settings against doctors' order and document same in nursing notes</p> <p>Verify pacemaker settings and record the following information on vital signs flow sheet of Clinical Information management System (CIMS) (Appendix I).</p> <p>NB: Minimum once per shift and following all changes to settings.</p>	<p>To ensure correct settings of mode, rate, sensitivity and output.</p> <p>To have baseline settings in case of alterations.</p> <p>Documentation provides continuity of care when information is shared (Scheider-Hickey and Bass 1991, An Bord Altranais 2002, Fischer 2008, Hazinski 2013).</p> <p>To promote and facilitate continuity of care and good communication through effective documentation. (Schneider-Hickey and Baas 1999, An Bord Altranais 2002, Hazinski 2013).</p>

<input type="checkbox"/> Patient's name and hospital number <input type="checkbox"/> Date and time <input type="checkbox"/> Rate <input type="checkbox"/> Atrial sense <input type="checkbox"/> Ventricular sense <input type="checkbox"/> Battery change (date) <input type="checkbox"/> Pacing wires <ul style="list-style-type: none"> ▪ Secured to patient ▪ Entry site dry ▪ Secured to pacemaker <input type="checkbox"/> Pacing spike(s) on monitor <input type="checkbox"/> Heart rate from arterial line or alternative source <input type="checkbox"/> Ensure lock on Document make and model of external pacemaker and any changes.	(Appendix I, II and III). NB: pacing spikes may not be visible on telemetry (PICU only). Assists in tracking pacemaker malfunction.
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SECTION 3: PACEMAKER AND WIRES

Action	Rationale and Reference
Assess integrity and security of pacing wires, ensuring no loose connections or wire fractures (minimum once per shift). <u>NB:</u> Take extra care when moving patient. Ensure wires are secure and pacemaker box and leads are supported.	To ensure good pacemaker connection and prevent disconnection. To prevent accidental changes to settings. (Appendix IV). (Martin and Aragen 1992, Hazinski 1999, Dwyer 2001, Reynolds and Apple 2001, Dwyer and Bauer 2010).
Ensure the pacemaker box is secure Ensure the cables are secure. NB: Pacemaker should be visible at all times. If pacemaker is dropped or becomes damaged it should be replaced immediately by the medical team and sent to clinical engineers for evaluation.	To prevent strain and accidental disconnection or dislodgement of pacing wires and damage to the pacemaker box. (Keenan 1995, Cottle 1997, Dwyer, 2001, Overbay and Criddle 2004, Reiswig-Timothy and Rodeman 2004). To ensure the pacemaker is functioning correctly.
If alternative pacemaker is required contact PICU 2 nd Floor first and then theatre dept. or clinical engineer for replacement. (Clinical Engineers: Bleep 465 / 008, Ext 6465. Out of Hours via switchboard) Inform cardiothoracic, medical / surgical team via bleep or out of hour's telephone number via switch board. Also contact consultant in charge.	To ensure timely replacement of pacemaker.
Battery Use 9 volt alkaline batteries only. NB: DO NOT USE rechargeable batteries Record battery voltage at beginning of shift and following insertion of new battery.	Risk of low capacity and unstable charge which may cause a pacemaker malfunction (St Jude Medical 2011). To ascertain battery status.

When battery is in use, the battery should be changed when battery depletion symbol displays only one blinking segment and warning message **'Change battery!'** appears. This is repeated every 10 minutes.
Ask cardiothoracic team to change the battery (Appendix III).



Figure 1: Battery Symbol

Cardiothoracic (medical / surgical) team change all temporary external pacemaker batteries

After inserting a new battery, the device Model 3085 needs 30 minutes to recharge its internal power capacitor in order to perform the bridging function again.

Cardiothoracic (medical / surgical) team to change pacemaker battery:

- With each new patient and then minimum of every 3 days

Label rear of pacemaker, with date the battery was last changed, nurses initials and document same in the nursing notes.
Ensure safe disposal of battery.

Have a replacement 9 volt battery available at the bedside at all times.

Critical Battery Depletion

The nurse should avoid this occurring by organising battery change earlier.

When critical battery depletion occurs the battery symbol will be empty and blinking. The warning message;

'Hurry up! Change battery!' will display. This is repeated every 2 minutes.

Battery will need to be replaced immediately.

Ensure manufacturer's user manual is always available for reference, in an area that all staff are aware of and have access to.

To allow cardiothoracic (medical / surgical) team to replace battery in a timely fashion. Battery change level is reached. There is approximately 24 hours reserve of battery life on Model 3085 if pacemaker mode set on standard setting (St Jude Medical 2011).

To minimise risk and create a safe environment should interruption of pacing / complications occur during the procedure.

NB: during battery changeover the pacemaker provides a minimum of 30 seconds additional power for extra safety. Battery change should take place WITHOUT DELAY but avoid undue haste (Jude Medical 2011).

To ensure, fully charged battery in situ.

NB: AV sequential pacing exhausts a battery more quickly than ventricular demand pacing.

(Hazinski 2013).

To have replacement in case of battery failure (Dwyer 2001, Mater Misericordiae University Hospital 2011, Yorkhill Children's Hospital 2011, Hazinski 2013).

To minimise risk and ensure the infant/ child receives continuous and uninterrupted pacing (St Jude Medical 2011).

Critical battery change level has been reached and immediate battery change is required (St Jude Medical 2011).

Readily available for reference.

Increased familiarity with pacemaker (Dwyer 2001, St Jude Medical 2011).

***Electrical safety is especially important when caring for a patient with a pacemaker
(Overbay and Criddle 2004)***

Section 4: ELECTRICAL SAFETY

Action	Rationale and Reference
When not in use cover pacing wires with finger stall (thumb, cut diagonally) of a powder-free non-sterile examination glove i.e. Sempercare.	To prevent micro shocks. Micro shocks are associated with ventricular arrhythmias and are a potential lethal hazard. Temporary epicardial pacing wires provide a direct low resistance pathway to the heart for an electrical current. Rubber is a poor conductor of electricity. (Hickey and Baas 1991, Owen 1991, Baas <i>et al</i> 1997, Norman 1998, Reynolds and Apple 2001, Reiswig-Timothy and Rodeman 2004, Beattie 2005, Mater Misericordiae University Hospital 2011, Hazinski 2013), (Appendix V).
Wrap the wires in finger stall with a gauze square, into a small parcel and secure to chest with Tegaderm. Wrap atrial and ventricular wires separately. NB: DO NOT USE TAPE to secure pacing wires within finger stall Label atrial and/or ventricular wires separately.	To keep dry (avoid baths, showers and unnecessary contact with water) (Hazinski 2013). Water is an excellent conductor of electricity. To prevent child pulling or interfering with wires. To prevent pressure marks to chest. Ensure easy accessibility to pacing wires, if required. Easy identification of wires. (Lynn-Mc Hale <i>et al</i> 1987*, Schneider-Hickey and Baas 1991, Berry <i>et al</i> 1997, Reynolds and Apple 2001, Overbay and Criddle 2004).
Wear non-sterile examination gloves at all times when handling pacing wires, especially terminal ends.	Terminal ends are not insulated. To prevent micro shocks and static electricity being transmitted via the nurses' hands to the patient. (Lynn-Mc Hale <i>et al</i> 1987, Hickey and Baas 1991, Beery <i>et al</i> 1996, Baas <i>et al</i> 1997, Norman 1998, Reynolds and Apple 2001, Overbay and Criddle 2004, Reiswig-Timothy and Rodeman 2004, Beattie 2005).
Appropriate warnings should be issued against the potential serious risk of using mobile communication devices in the vicinity of a patient with a pacemaker. The use of mobile phones / walkie-talkies is PROHIBITED in close vicinity to the patient.	There is a potential risk of electromagnetic interference to external pacemakers by mobile phones and walkie-talkie's (Trigano <i>et al</i> 1999, Medtronic 2001).

* The Nurse Practice Committee acknowledges the age of this reference. However, the article is a seminal piece of work which provides a comprehensive overview of the care of a pacemaker and which has been cited extensively by subsequent authors.

Section 5: INSERTION SITE

There are 2 common insertion sites in paediatrics

- **EPICARDIAL wires via transthoracic site.**
- **TRANSVENOUS (endocardial) wires via CVC insertion site.**

NB: Atrial pacemaker wires traditionally exit the chest to the right of the sternum. Ventricular pacemaker wires exit the chest to the left of the sternum (Hickey and Baas 1991, Owen, 1991, Fisher 2008). Always check the surgical notes to verify the type and location of pacing wires.

An exception to the rule is in cases of dextrocardia or situs inversus.

Aseptic Non-Touch Technique (ANTT) is a mechanism which helps to prevent contamination of susceptible sites by micro-organisms that could cause infection (Hart 2007, Pratt et al. 2007). ANTT is achieved by preventing contamination of external parts of the pacing wires and the insertion site. **Level 2 ANTT** should be used if cleaning or dressing the insertion site of the pacing wires is necessary. **Level 3 ANTT** is appropriate for handling and securing the epicardial pacing wires.

Action	Rationale & Reference
Transvenous Site: Dress as per intravenous clinical guidelines: (ANTT Level 2, Veniguard ® dressing). Assess daily and redress minimum of every 7 days or as clinically indicated.	To prevent infection (OLCHC 2007).
Epicardial Wire Site: Leave uncovered if dry. Dress as necessary with dry dressing, i.e. Mepore, if oozing present. Clean skin with 0.9% Normal Saline as clinically indicated	To prevent infection. (Owen 1991, Overbay and Criddle 2004).
Assess insertion site for bleeding. If present apply pressure dressing. Notify cardiothoracic (medical / surgical) team.	To detect and treat early signs of bleeding.
Assess insertion site for signs of infection i.e. redness, swelling or oozing. If present: Notify cardiothoracic (medical / surgical) team. Clean site and obtain swab for culture and sensitivity.	Early detection of signs of inflammation / infection. Spread of infection along the catheter may cause septicaemia. To ascertain microbiology status. (Lynn-McHale <i>et al</i> 1987, Fischer 2008, Dougherty and Lister 2011). Cleaning site prior to swabbing is required to ensure accurate collection of and reduced contamination of organisms from the wound (Kelly 2003, Kingsley and Winfield-Davies 2003, OLCHC 2008).

Section 6: PSYCHOLOGICAL CARE

Action	Rationale & Reference
Provide explanations, education and emotional support to child and family.	To foster understanding and relieve anxiety.
Involve the multidisciplinary team including: cardiac team; cardiology clinical nurse specialist and play specialist as appropriate.	To provide knowledge and skills as necessary for compliance with treatment (Van Orden-Wallace 1998, Hockenberry and Wilson 2011).

3. Trouble Shooting

Most troubleshooting associated with pacemaker systems is related to changes in the patient's medical condition or misinterpretation of normal pacemaker function. In all instances it is vital to assess the patient and identify the cause.

- ♦ ***It is essential for nurses to contact the cardiothoracic (medical/ surgical) team IMMEDIATELY, for early and timely intervention.***

There are four potential problems which can exist during pacing:

1. **Failure to Fire**
2. **Failure to Capture / Pace**
3. **Under Sensing**
4. **Over Sensing**

1. Failure to Fire

Failure to fire is characterised by the loss of output from the pulse generator, which is identifiable by an abnormally slow heart rate or asystole. Intervention should be specific to the problem found in the pacemaker system. If failure to fire cannot be corrected emergency measures may need to be initiated. Failure to fire related to pacemaker malfunction is rare. It is more likely to be related to settings, connections or changing thresholds.

Contact Cardiothoracic (medical / surgical) Team IMMEDIATELY

Problem	Intervention
Loose connection or disconnection between lead wire, cables and pacemaker	Ensure connections are secure
Fracture / dislodgement of lead wire	Assess integrity of lead wires and replace as necessary NB: Remember the skin can be used as a new or extra positive lead.
Low pacemaker battery	Insert new battery
Failure of pacemaker pulse generator	Replace pacemaker generator. Contact Cardiothoracic Team (medical / surgical).
Over sensing (not common in paediatrics i.e. P wave high, mainly occurs in adults, unless the sense thresholds have been set too low)	Contact Cardiothoracic Team (medical/surgical) to assess sensitivity and decrease if necessary

(Lynn-McHale et al 1987)

2. Failure to Capture / Pace

Capture occurs when the myocardium responds to the pacing stimulus by depolarising i.e. P wave or QRS wave. Failure to capture occurs when the myocardium fails to respond to a pacing stimulus. It will be seen as the pacing spike(s), not been followed by a P wave or QRS complex.

Contact Cardiothoracic Team (medical / surgical) IMMEDIATELY

Possible causes for increased pacing threshold:

- Inflammation or fibrosis at electrode site
- Increased serum Potassium or Calcium
- Acid base imbalances
- Medications i.e. Verapamil or Propanolol
- Fibrillation or flutter

Problem	Intervention
Loose connection between lead wire, cables, and pacemaker.	Ensure connections are secure.
Fracture / insulation break of lead wire. Displacement of lead wire.	Assess integrity. Contact Cardiothoracic Team (medical / surgical) who will replace it if required. NB: The skin can be used as a new positive electrode.
Low pacemaker battery.	Battery replaced by Cardiothoracic Team (medical / surgical).
Failure of pulse generator	Pulse generator replaced by cardiothoracic team.
Increased pacing threshold/ inadequate output (energy) for depolarisation.	Contact Cardiothoracic Team (medical/surgical). Who will reassess pacing threshold and identify and treat the underlying physiological disturbances.

3. Under Sensing

Sensing is the ability of the pulse generator to 'see' the patients' own rhythm. Pacing spikes are present and regular but compete with the patients own inherent rhythm. This can occur when the sensing amplifier fails to detect the intrinsic activity of the heart, the sense threshold has been set too high or when the pacemaker loses the ability for self-inhibition (fires regardless). Mechanical failure of the pacemaker is rare. The pacemaker's response to under sensing is to over pace, with pacing spikes falling randomly in the cardiac cycle. This situation must be corrected as soon as possible because there is a potential for the pacemaker to deliver a stimulus in the refractory period of the cardiac cycle, which corresponds with the T wave when the heart is repolarising (heart vulnerable). It may potentiate lethal arrhythmias: i.e. ventricular tachycardia or ventricular fibrillation.

Possible causes for under sensing (QRS detection):

- Tissue ischaemia / fibrosis
- Electrolyte disturbance
- Poorly positioned lead
- Fibrillation / atrial flutter
- Lead fracture
- Loose connections (Reynolds and Apple 2001)

Problem	Intervention
Inadequate QRS signal	Contact Cardiothoracic Team (medical / surgical) who may increase sensitivity (making the pacemaker more sensitive by decreasing mV to a smaller number) (Slota 2006)
Fracture/ dislodgement of pacing wire	Assess integrity. Contact Cardiothoracic Team (medical / surgical) immediately , who will replace as necessary. NB: Remember the skin can be used as a new positive electrode.
Battery depletion	Contact Cardiothoracic Team (medical / surgical) to replace battery.

4. Over Sensing

Over sensing is when the pacemaker is too sensitive and inappropriately senses internal and external signals and inhibits pacemaker output. The pacemaker generator misinterprets an electrical current as a QRS complex, inhibits itself and therefore does not fire. The Pacemaker may have detected a P wave or T wave, rather than the QRS complex or myopotentials i.e. electrical signals produced by skeletal muscle contraction. The sensing amplifier sees too many signals which the pacemaker interprets as the hearts intrinsic rate and therefore does not fire. In patients with a pacemaker dependent rhythm this will result in a pause in rhythm and reduction in cardiac output. Over sensing may be eliminated by reducing the sensitivity. This is performed by the cardiothoracic team (Fischer 2008).

Problem	Intervention
P or T wave sensing	Contact Cardiothoracic Team (medical / surgical) who may reduce sensitivity (making the pacemaker less sensitive by increasing mV to a higher number) (Slota 2006).
Skeletal muscle contractions (myopotentials) or shivering	Contact Cardiothoracic Team (medical / surgical) who may decrease sensitivity.
Electromagnetic interference	Identify and remove source. Contact Cardiothoracic Team (medical / surgical) who may decrease pacemaker sensitivity.

(Reynolds and Apple 2001)

POTENTIAL COMPLICATIONS OF TEMPORARY EXTERNAL PACING

Contact Cardiothoracic Team (medical / surgical) IMMEDIATELY

Problem	Cause
Arrhythmias <ul style="list-style-type: none"> PVC's Ventricular Tachycardia / Fibrillation Asystole 	<p>May result from myocardial irritability caused by pacing wires.</p> <p>If pacemaker stimulus occurs during QT interval, when the heart is repolarising. Removal of pacing wires can rarely cause ventricular arrhythmias, e.g. ventricular fibrillation.</p> <p>If pacing is discontinued abruptly or if batteries fail.</p>
Electrical Hazards	Leads provide a direct low resistance pathway to the heart for an electrical current.
Haemorrhage	Can occur during or after epicardial or endocardial lead placement or removal resulting in cardiac tamponade.
Pneumothorax or Pneumomediastinum	Cardiac perforation or air embolism can occur during transvenous pacemaker insertion.
Infection	Insertion sites should be inspected each shift to detect early signs of infection. (See c/o insertion site).
Displacement / Fracture of leads	Lead fracture impairs ability of unit to conduct an impulse (Slota 2006).
Failure to recognise asystole	Monitor may read pacing spikes as a QRS complex (even when no QRS follows the pacing spike). Asystole may therefore be missed (Oslizlok 2007).

4. Nursing Responsibilities in Assisting with the Removal Of Epicardial Pacing Wires

5.

Epicardial pacing wires (atrial x 2 +/- ventricular x 2) are routinely inserted by the cardio-thoracic surgeon following open heart surgery i.e. AVSD repair, Fallots Tetralogy repair and VSD repair. They are used to diagnosis and treat rhythm disturbances. Epicardial pacing wires are traditionally placed: atrial wires on right side of chest and ventricular on the left. Should pacing be required post operatively these wires allow the heart to be temporarily paced by an external pacemaker (Lynn-McHale *et al* 1998, Beattie 2005, Clark 2007, O'Brien 2008).

ACTION	RATIONALE
<p>Timing of Wire Removal Epicardial pacing wires are usually removed a minimum of 3-5 days post operatively and at least 24 hours prior to hospital discharge on instruction from cardio-thoracic team. The child will have a normal heart rate for age and be in sinus rhythm.</p> <p>Pre Procedure Investigations The child will have a 12 lead ECG/ 24 hour Holter ECG and Chest X-ray performed and reviewed by the medical team.</p> <p>The child will have a coagulation screen and platelet count performed and reviewed by the cardiothoracic team (medical / surgical).</p> <p>NB: Therapeutic Heparin infusion is discontinued 4 hours prior to the removal of pacing wires. The heparin infusion is then restarted 2 hours post procedure if there is no bleeding.</p> <p>IV Access Ensure patient has a patent intravenous cannula in situ prior to the procedure</p> <p>Monitoring The child will have observations taken and recorded prior to removal i.e. temperature, pulse, respirations, SaO2 and blood pressure</p> <p>The child will be attached to telemetry / cardiac monitor for the procedure for minimum of 24 hours</p> <p>Safety The nurse will ensure emergency equipment is working and available at the bedside</p> <ul style="list-style-type: none"> • Amubag / rebreathing circuit and appropriate mask • Oxygen and mask • Suction equipment and suction catheters • Antiarrhythmic drugs and defibrillator (available on ward / unit). <p>Location Plan location of procedure. Use treatment room if available</p> <p>Psychological Preparation The child and / or family will receive adequate explanation of the procedure at an appropriate level and emotional support prior to wire removal. Encourage questions and answers. Child should be informed of sensation likely to be experienced during procedure i.e. 'mild to moderate pulling sensation' as clinically indicated. A play therapist may be utilised for preparation and / or distraction if clinically indicated.</p>	<p>To ensure that epicardial pacing wires are removed under safe conditions and observation throughout the day following removal under optimal conditions should emergency intervention be required (Johnson <i>et al</i> 1993, PCNA 2003, Beattie 2005).</p> <p>To ensure patient safety (PCNA 2003).</p> <p>The presence of coagulopathy requires treatment before removal of pacing wires. To minimise the risk of bleeding post removal of wires and development of pericardial tamponade (Wollan 1995, PCNA 2003, Beattie 2005, Jowett <i>et al</i> 2007, O'Brien 2008).</p> <p>Pacing wires should only be removed after therapeutic heparin has been discontinued (Reade 2007, Mater Misericordiae University Hospital 2011, OLCHC 2012a).</p> <p>To provide a route for fluid resuscitation or anti arrhythmic medication should it be required (Johnson <i>et al</i> 1993, Beattie 2005).</p> <p>To establish baseline observations for comparison post procedure and detection of changes in patients condition in a timely fashion (Lynn-McHale <i>et al</i> 1998, Clark 2007).</p> <p>To assess the child for potential arrhythmias or pericardial tamponade (Johnson <i>et al</i> 1993, Wollan 1995, O'Brien 2008).</p> <p>To create a safe environment and maintain patient safety (Wollan 1995, PCNA 2003).</p> <p>Avoid performing the procedure in a child's "safe zone" to minimise stress of hospitalisation (O'Brien 2008).</p> <p>To relieve fear, anxiety and foster understanding and cooperation of the procedure. Information may need to be reinforced if the child is stressed (Van Orden-Wallace 1998, PCNA 2003, Roschkov and Jensen 2004, O'Brien 2008, Mullins <i>et al</i> 2009, Hockenberry and Wilson 2011).</p>

<p>Pain Relief Administer analgesia and sedation if required as prescribed by the medical team as per 'Procedural Analgesia and Sedation' Algorithm (Appendix VI). Sedation will always be given in conjunction with analgesia. Assess pain score.</p> <p>Positioning The child will be positioned supine or alternatively at 30-45o angle if not possible in bed for the procedure. Ensure privacy in older child / adolescent.</p> <p>Procedure</p> <p>Responsibility for removal of pacing wires <i>The cardio-thoracic team are responsible for removal of the epicardial pacing wires.</i></p> <p>Equipment</p> <ul style="list-style-type: none"> • Dressing trolley • Dressing pack including sterile gloves and gauze • 0.05% Chlorhexidine solution • Opsite occlusive dressing • Stitch cutter <p>Cardio-thoracic Surgeon will wash hands using a Aseptic Non-touch Technique (ANTT) (level 2) and put on sterile gloves</p> <p>The nurse will decontaminate hands and assist doctor in laying dressing trolley.</p> <p>Nurse will remove dressing around pacing wires to expose pacing wires and then repeat handwashing.</p> <p>The doctor will clean around pacing wire sites with Chlorhexidine 0.05%.</p> <p>The atrial pacing wires are usually removed first if present and ventricular wires last.</p> <p>The holding suture of the pacing wire is released using a stitch cutter.</p> <p>Holding the pacing wire near to the chest it will be pulled with a smooth, continuous, downward, pulling motion, exerting gentle traction until release from the epicardium is felt.</p> <p>The tip of the epicardial pacing wire is inspected for intactness and pieces of myocardial tissue.</p> <p>The procedure is repeated by the doctor for each additional pacing wire(s).</p>	<p>To provide comfort and minimise pain. Patients have reported 'mild to moderate pulling sensation' on epicardial pacing wire removal (Mullins <i>et al</i> 2009, Mater Misericordiae University Hospital 2011, OLCHC 2012b).</p> <p>To ensure correct positioning for removal of epicardial pacing wires. Semi upright position is often preferred in children as it is often associated with less anxiety (Wollan 1995, Clark 2007, O'Brien 2008; Beattie 2008).</p> <p>Procedure only performed by Cardiothoracic Team because of the potential complications that may occur following the procedure (Roschkov and Jensen 2004, O'Brien 2008).</p> <p>To prevent cross infection, universal precautions (PCNA 2003). Epicardial pacing wires provide a direct low resistant pathway to the heart and patient may receive micro shocks due to static electricity (Wollan 1995, Beattie 2005).</p> <p>To minimise transmission of organisms (O'Brien 2008).</p> <p>To allow complete visualisation of pacing wire site and holding suture (O'Brien 2008).</p> <p>Reduces risk of infection (O'Brien 2008).</p> <p>This allows pacing of the ventricle to restore cardiac output in the event of a symptomatic arrhythmia, following removal of atrial pacing wires.</p> <p>To reduce the risk of trauma. Jerking or pulling against resistance may cause bleeding. (Wollan 1995, Sheehan <i>et al</i> 2001, PCNA 2003, Clark 2007). (Clark 2007).</p> <p>To ensure that the entire wire has been removed and determine the risk of infection, migration or haemorrhage (Johnson <i>et al</i> 1993, Wollan 1995, Clark 2007, Beattie 2008).</p>
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<p>Following removal an Opsite occlusive dressing is applied to the site for a minimum of 24 hours</p> <p>Dispose of used supplies and sharps appropriately.</p> <p>Remove gloves and wash hands.</p> <p>Post Procedure</p> <p>Bedrest The child will remain on bedrest for 1- 2 hour following the procedure</p> <p>Monitoring Monitor and record vital signs immediately following the procedure: heart rate; rhythm; respirations and blood pressure. Repeat every 15 minutes x 2 and then every 30 minutes x 2 and then as patients clinical condition dictates. Observe patients SaO₂; colour; perfusion and conscious level.</p> <p>Complications The child will be observed for complications</p> <ul style="list-style-type: none"> • Bleeding If bleeding occurs apply direct pressure with gauze for several minutes until ceases. Persistent bleeding should be reported immediately to the cardiothoracic surgical team. Patients on anticoagulation therapy are at greater risk of bleeding. • Arrhythmias i.e. ventricular ectopic beats, due to mechanical irritation of the myocardium. Be extra vigilant if the child has a history of heart failure or previous cardiac surgery. Report excessive ectopic beats or sustained arrhythmias to the cardio-thoracic team • Pericardial Tamponade Rare but serious complication. (Signs & symptoms include: pallor, collapsed child; tachycardia; tachypnoea; dyspnoea, reduced capillary refill, cool extremities, decreased SaO₂; sweating; decreased conscious level, hypotension). Report immediately to cardio-thoracic surgical team. <p>An echocardiogram may be performed post procedure if clinically indicated or there is deterioration in the patient's condition.</p> <p>Documentation The doctor and nurse will record the procedure in medical / nursing notes / clinical information management system (CIMS) including date, time, who removed epicardial pacing wires, number and type. Also patients' condition and response to the procedure.</p> <p>Discharge Information The parents and if appropriate the child should be aware of signs and symptoms of possible complications and who to phone for advise following discharge</p>	<p>To prevent infection (O'Brien 2008).</p> <p>Standard precautions and to reduce transmission of organisms.</p> <p>Standard precautions.</p> <p>To prevent injury from cardiovascular compromise secondary to bleeding, arrhythmia or tamponade (Johnson <i>et al</i> 1993).</p> <p>To ensure early detection and timely treatment of any potential complication. Pericardial tamponade usually presents within 2 hours of removal. (Johnson <i>et al</i> 1993, Wollan 1995, Carroll 1998, Lynn-McHale <i>et al</i> 1998, PCNA 2003, Beattie 2005).</p> <p>To control bleeding (Beattie 2008).</p> <p>These patients may be at greater risk of arrhythmias. Transient arrhythmias are common and often subside spontaneously (Carroll <i>et al</i> 1998).</p> <p>To ensure prompt and timely treatment (O'Brien 2008, Mahon <i>et al</i> 2012).</p> <p>Echocardiogram may exclude or reveal pericardial tamponade (Leahy 1993, Clark 2007).</p> <p>To ensure satisfactory documentation of the procedure and continuity of patient care (An Bord Altranais 2002, Clark 2007).</p> <p>To ensure patient safety and referral in an appropriate manner (PCNA 2003).</p>
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Ensure retained wire lead or fragments are communicated to ward nursing staff on transfer documentation as clinically indicated.

It should be clearly documented in the patient's medical and nursing notes also.

Advise parent regarding the long term need to inform doctor regarding any possible signs of infection i.e. malaise, chills, fever and signs of infection at epicardial pacing wire exit sites

There is increased risk of infection as they create an open wound through the skin which communicates with the pericardial space.

Complications from retained epicardial wires have been described in the literature i.e. localised abscess / fistula to infective endocarditis. Complications have been reported to occur up to many years later.

Ensure satisfactory communication and continuity of care (An Bord Altranais 2002, Yorkhill Children's Hospital 2011, Shaikhrezai *et al* 2012).

To ensure prompt and timely treatment of any infection at pacing wire sites or due to retained epicardial pacing wire

There is a potential risk of endocarditis and doctor or dentist may decide to administer prophylactic antibiotics prior to any invasive procedure (Johnson *et al* 1993).

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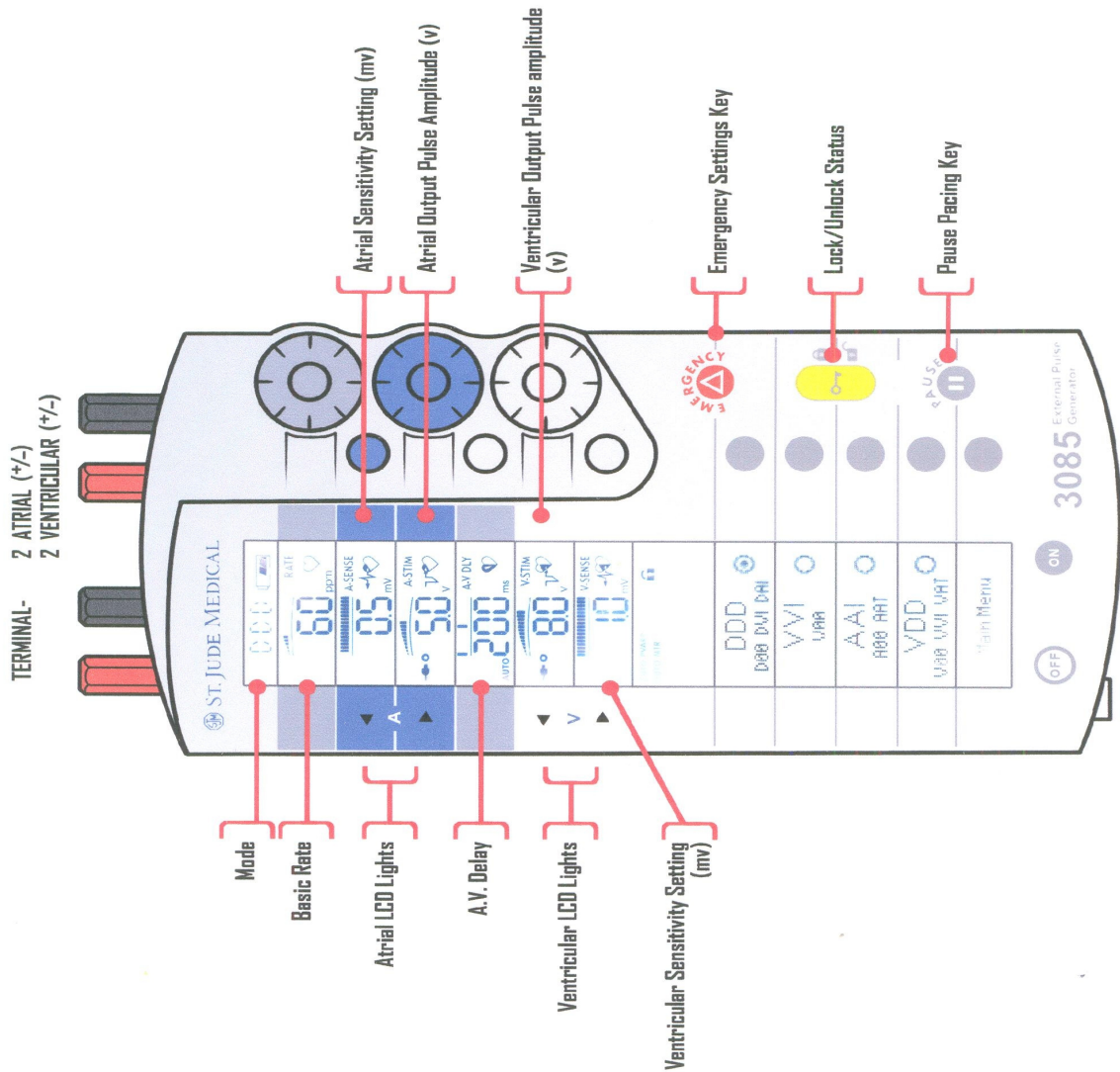
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APPENDIX I: PACEMAKER CONTROLS

Key to St. Jude Medical 3085 Pacemaker Controls and LCD Display



NB: For guidance only, pacemakers may vary

OUTPUT CONTROL

(Stimulation Threshold)
Adjust the amount of current deliver to the epicardium or endocardium in order to produce myocardial depolarisation - capture

SENSITIVITY CONTROL

The responsiveness of the pacemaker to the patient's intrinsic electrical activity. The sensitivity threshold refers to the minimum R wave amplitude needed to be detected by the Pacemaker.

A-V-INTERVAL

The interval between atrial and ventricular depolarisation.

VENTRICULAR RATE

The minimum ventricular rate accepted below which the pacemaker will fire

SENSE/PACE INDICATOR

Indicates whether the Pacemaker is sensing intrinsic activity (P or R = atrium or ventricle) or pacing (A or V = atrium or ventricle).

APPENDIX II: ST JUDE MEDICAL 3085 PACEMAKER

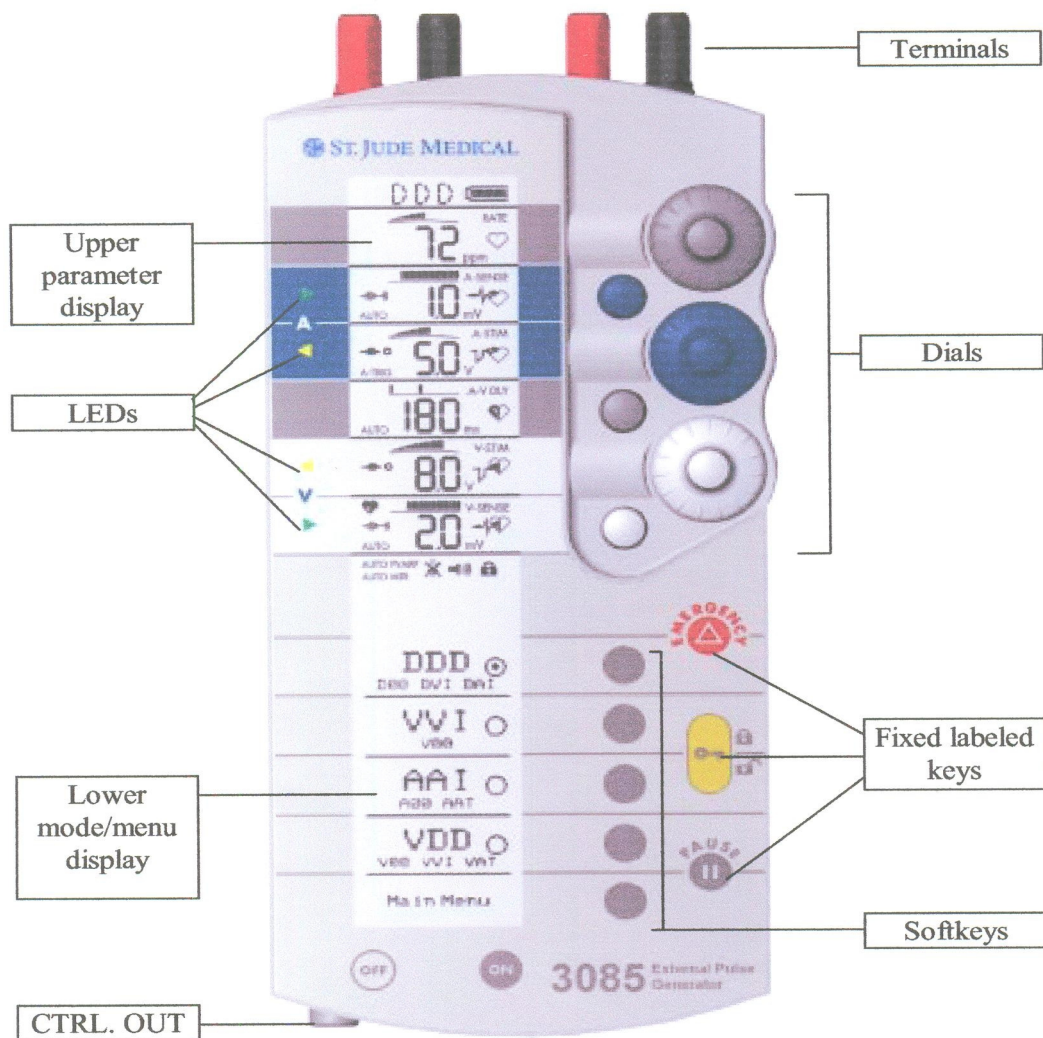


Figure 2: St Jude Medical 3085 Pacemaker Front Display Face.



Figure 3: Ventricular and Atrial Terminals.

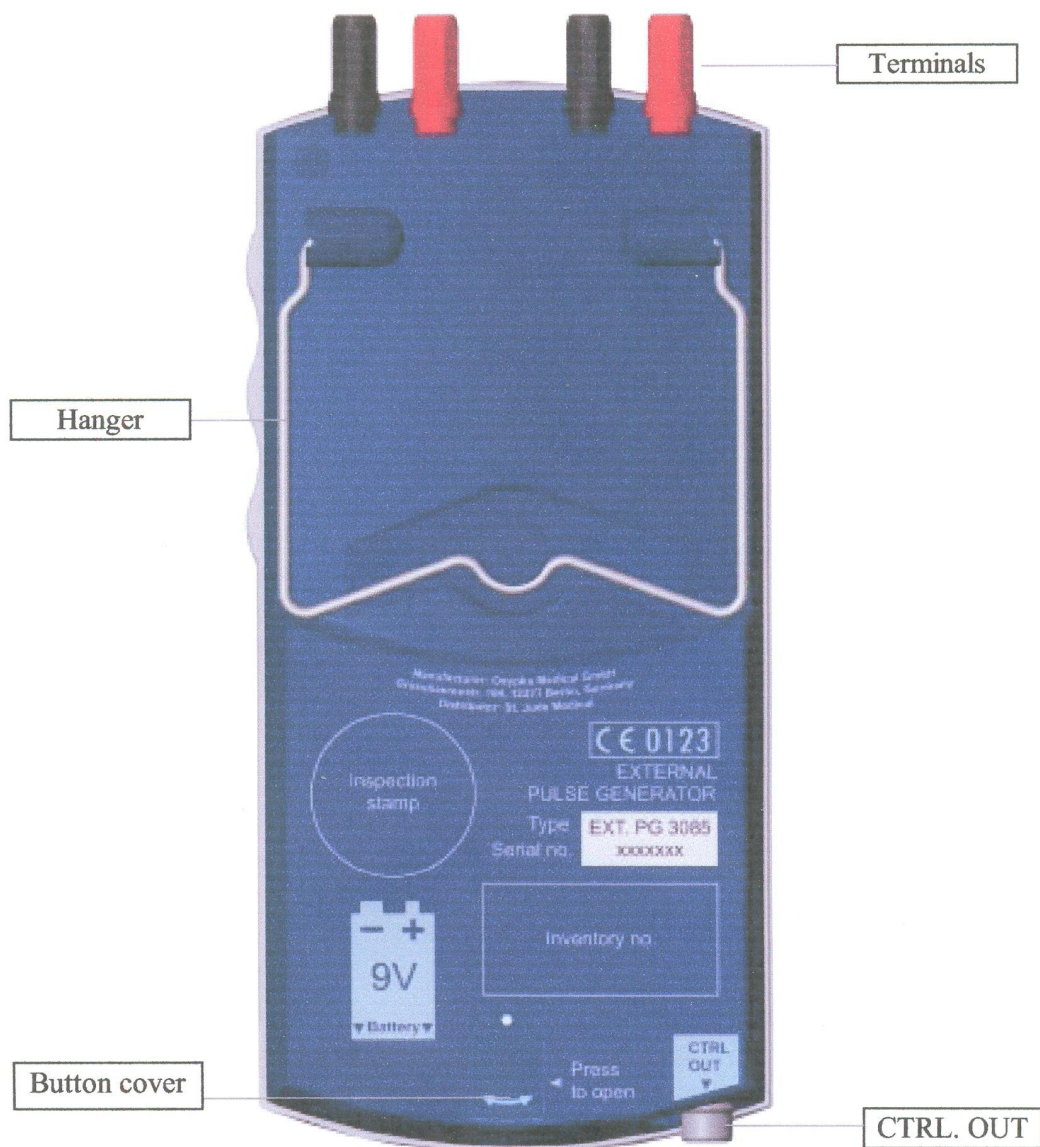


Figure 4: St Jude Medical 3085 Pacemaker Rear View.

APPENDIX III: St Jude Medical Model 3085

Turning on Pacemaker

1. Press key labelled **ON** (pacemaker will run a self test).
2. When the pacemaker was previously in Standby Mode, it will commence functioning at the last saved parameter settings.
3. The key **Lock / Unlock** must be pressed and released to ensure it is functioning properly when the pacemaker was previously OFF.



Figure 6: Lock / Unlock Key

4. When Lock/Unlock key is not pressed and released within 30 seconds, an error message will be displayed ‘**Startup timeout’ (Press unlock)** and pacemaker will switch off.’
5. Turn on programme commences.
6. Soft keys 1-5 will display a Menu Mode
 - Key 1 – Mode DDD
 - Key 2 – Mode VVI
 - Key 3 – Mode AAI

NB: Pacemaker Settings are set by the Medical / Surgical Cardiothoracic Team.

Locking / Unlocking

1. Pacemaker will automatically lock if no key has been pressed for 30 seconds. Prevents accidental
2. To unlock press key Lock / Unlock.
3. Lock symbol will indicate whether the pacemaker is locked or unlocked.



Figure 7: Lock Symbol

4. A warning beep and lock symbol will blink for 2 seconds if keys are pressed / dials turned when pacemaker is locked.

LED lights for Sensing and Stimulation

1. LED lights located at upper left side.
2. They indicate atrial and ventricular sensing and stimulation.
3. Green LED lights flashing indicate sensing.
4. Yellow LED lights flashing indicate stimulation.

NB: Lights flash brightly initially when pacemaker turned on to indicate functioning satisfactorily.

Emergency Key

Pressing the emergency button key will commence pacemaker stimulation at emergency settings.

Pause Pacemaker

Pressing the pause button key will disable pacemaker stimulation as long as it is pressed.

Turning Off the Pacemaker

1. Press lock/ unlock key.
2. Press OFF key.
3. A soft key power-off menu will display.
 - Press key 1 – OFF (with no storage). Actual settings are not saved
 - Press Key 2 – Stand-by with data stored.

NB: No battery power is consumed in the stand-by mode.

(St Jude Medical 2011).

APPENDIX IV: CHANGING THE BATTERY (MODEL 3085)

This procedure is performed by Cardiothoracic Medical/ Surgical Team.

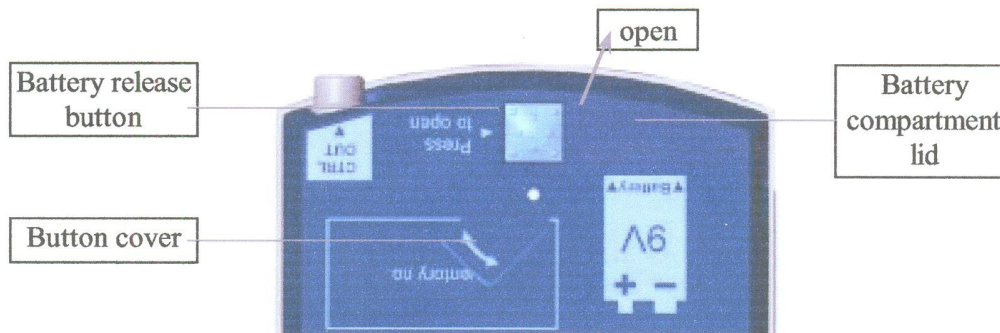


Figure 5: Battery Compartment.

1. Turn protective cover of the battery compartment lid whilst pressing the battery release button.
NB: This button cover prevents the release button from being unintentionally pressed.
2. Open battery lid
3. Battery is removed from the compartment.
4. Replace with new 9 volt battery.
5. Battery compartment lid is closed until audible sound of it latching into place.
6. Protective cover of battery compartment lid is rotated over the battery release button.
7. Dispose of old battery in an environmentally friendly manner

(St Jude Medical 2011).

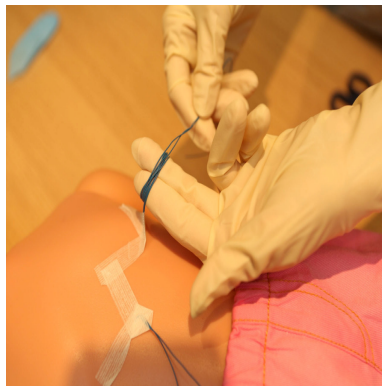
APPENDIX V: Securing Epicardial Pacing Wires when not in use.

Equipment

- Non-disposable gloves
- Gauze
- Tegaderm dressing
- Labels



1. Cut the thumb off a non-disposable glove.
NB: Thumb has a wider opening.



3. Wearing gloves, wrap the two pacing wires around your 2nd and 3rd fingers.



4. and 5. Pacing wires now form a small roll.



6. Insert pacing wire roll into at Bottom of the thumb of the previously cut non-disposable glove.



7. Wires in thumb of glove, now form a small parcel.



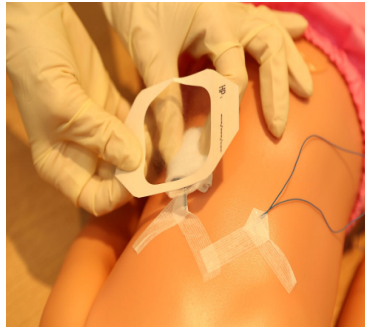
8. Open one sheet of gauze under the wires in the glove.



9. Wrap gauze around wires in the glove.
NB: Gauze protects skin and ensures comfort.



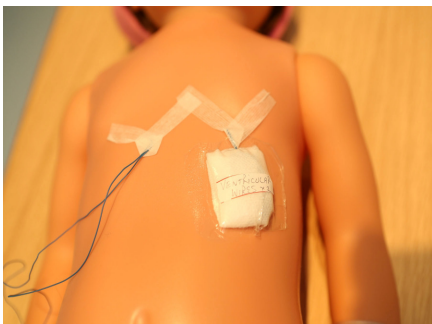
10. The gauze forms a small parcel around wires in the glove.



11 Apply tegaderm dressing over the gauze.



12. Ensure tegaderm dressing Secures gauze to skin at all edges. Apply second dressing PRN.



13. Label wires 'trouser leg' is
NB: Atrial wires are on the right and ventricular wires on the left.



14. Repeat procedure with second set of wires if required.

APPENDIX VI: Procedural Analgesia and Sedation in PICU / HDU (OLCHC 2012).

Our Lady's Children's Hospital, Crumlin, Dublin 12.

Procedural Analgesia and Sedation in PICU / HDU

This protocol outlines the procedural analgesia and sedation in use in the PICU / HDU

Indications:

1. Chest drain removal
2. Pacing wire removal
3. PD catheter removal
4. CVC/ART line insertion – if required

All analgesic and sedative agents have the potential to compromise respiratory status and cardiovascular stability. Ensure patient monitoring with appropriate alarm parameters and audible SpO2 pulse tone throughout procedure. Safety equipment, fluid boluses and airway adjuncts in appropriate sizes must be available in case of apnoea or respiratory depression.

All Patients

Review analgesia and sedation regime pt already on.
Ensure intervention such as chest drain removal is timed to coincide with peak analgesic effect of agent given

Paracetamol IV/NG/PR

NSAID (only if appropriate)

Children 6-12kgs	Children >12kgs
Ibuprofen PR	Diclofenac PR or Ibuprofen PO

Topical analgesia – apply AMETOP to site 30 minutes prior to procedure.

Babies <3 months old: Use SUCROSE 24% (0.2ml given PO via syringe 2 mins prior to procedure)

Remember: Avoid NSAID in the following:

- Neonates
- Children with reduced urine output (<1ml/kg/hr)
- Children receiving anti-coagulants, e.g. heparin/firzaparin or who are "auto-anti-coagulated"
- Children with gastritis / coffee-ground NG aspirates

All Ventilated Patients	
COMFORT < 12:	Additional analgesia & sedation may not be required.
COMFORT > 12:	Morphine 40microgram/kg IV.
	Repeat if necessary

Extubated post op patients if already on IV morphine	
<1 year	1-2 years
Morphine 10-20 microgram/kg IV +/- Repeat if necessary	Morphine 20-40 microgram/kg IV +/- repeat if necessary
	Midazolam 0.25-0.5mg/kg PO +/-
	Midazolam 0.5mg/kg PO +/-

Extubated patients not on IV morphine		
<1 year	1-2 years	>2 years
Fentanyl 0.25-1 microgram/kg IV +/- Midazolam 0.25 mg/kg PO or Clonidine 1 microgram/kg PO +/- Midazolam 0.25mg/kg PO or Ketamine – Consultant decision only	Fentanyl 1-1.5microgram/kg intranasally plus Midazolam 0.25-0.5mg/kg PO or Clonidine 1-1.5 microgram/kg PO plus Midazolam 0.25-0.5mg/kg PO or Ketamine – Consultant decision only	Fentanyl 1-1.5 microgram/kg intranasally plus Midazolam 0.5 mg/kg PO or Clonidine 1-1.5 microgram/kg PO plus Midazolam 0.5mg/kg PO or Propofol IV – Consultant decision only

APPENDIX VI: PACEMAKER GLOSSARY

A

Arrhythmia. An abnormal rhythm of the heart (too slow, too fast, or uneven), which can cause the heart to pump less effectively. In pacing, any rhythm disturbance. Examples include bradycardia, tachycardia, any markedly irregular rhythm, block or the presence of premature contractions.

A. V Delay. Atrio-ventricular delay in a dual chamber pacing mode. The AV delay is the period between an atrial event (paced or sensed) and a paced ventricular event. In DDD pacing the AV delay is generally programmed to 120 -150 milliseconds (msec) depending on the patient's age, allowing a heart rate of up to 140-150 / minute. If the heart rate is higher the AV interval needs to be reduced (Oslizlok, 2007; Wood, 2007).

C

Capture. The successful depolarisation and contraction of a cardiac chamber caused by the pacemaker's output pulse. One-to-one capture occurs when each pacemaker output pulse results in a contraction.

Cardiac Cycle. One complete heartbeat. Seen on the ECG as a P wave, a QRS complex and a T wave.

Cardiac Output. The volume of blood, measured in litres, ejected by the heart per minute. Cardiac output is determined by multiplying the heart rate and the stroke volume.

F

Fibrillation. A type of cardiac arrhythmia characterised by rapid, unsynchronised quivering of atria or ventricles. Atrial fibrillation may be asymptomatic, but ventricular fibrillation is typically fatal if not corrected within minutes.

I

Intrinsic. An intrinsic beat is a naturally occurring heartbeat. Intrinsic rate is the patient's own heart rate. Sometimes called native.

Inhibition The effect of pulse suppression when pacemaker in a demand mode and senses a cardiac depolarization.

L

Lead The insulated wire plus electrode(s) and terminal pin used to connect the pulse generator to the cardiac tissue. The lead carries the stimulus from the pulse generator to the heart and in demand modes, relays intrinsic cardiac signals back to the sense amplifier of the pulse generator. A single-chamber pulse generator requires one lead, while a dual-chamber pulse generator usually requires two (one for the atrium, the other for the ventricle).

Lead Dislodgement. The detachment of the pacing lead from the intracardiac location to which it had been positioned.

M

Microshock Low-voltage electrical current or static electricity which can pass from the nurse and into the patient. As little as 0.1mA has the potential to cause ventricular fibrillation.

O

Output. The electrical stimulus delivered by the pulse generator and usually defined in terms of pulse amplitude (V) and pulse width (ms). (In pacing, output used alone usually refers to electrical output of the device, while the term cardiac output is used for blood throughput of the heart.) Maximum 10 volts (Wood, 2007). Output usually set 3 times output (pacing) threshold.

Output (Pacing) Threshold The minimum electrical stimulus needed to consistently elicit a cardiac depolarisation (capture) and expressed in millivolts (mV). Usually 2 mV or less.

Over Sensing Detection by the pulse generator's sense amplifier of inappropriate electrical stimulus. The over sensed signal may or may not be visible on a surface ECG. Over sensing can often be corrected by making the pacemaker less sensitive (increasing the mV value), programming to a triggered mode or by the judicious programming of the refractory period.

P

Premature Ventricular Contractures (PVCs) A ventricular contraction initiated by an ectopic focus which occurs earlier than the next expected normal ventricular contraction. Also known as 'ventricular ectopic beats' or ventricular premature beats (VPBs).

R

Refractory. (1) Inability of tissue to respond to a stimulus.
(2) Inability of a pacemaker to respond to an incoming signal.

Refractory Period.

- (1) The length of time the myocardium is incapable of responding to a stimulus.
- (2) In pacing, an interval or timing cycle following a sensed or paced event during which the sense amplifier will not respond to incoming signals. Dual-chamber pacemakers have separate refractory periods for each chamber (atrial and ventricular). In most modern pacemakers, the refractory periods are programmable values.

S

Sensing. The ability of the pacemaker to recognise and respond to electrical activity in the heart. How the pacemaker responds to sensed signals depends on its programmed mode and parameters.

Sensitivity. A pacemaker parameter which determines the amplitude of signals to which the device's sense amplifiers will respond. Sensitivity is stated in millivolts (mV). Note that the higher the mV value, the lower the sensitivity. Thus the lower the mV value, the more sensitive the device. Average setting is 2, lowest 1mV (Wood, 2007).

Sensitivity Threshold The minimum atrial or ventricular intracardiac signal amplitude required to inhibit or trigger a demand pacemaker, expressed in millivolts. Sensitivity is usually 2-3 times more sensitive than sensitivity threshold (i.e. divide threshold by 3).

Spike. A small but sharply vertical deflection that appears on the surface ECG indicating that a pacemaker output was delivered. It is caused by the brief discharge of electricity produced by the pacemaker to stimulate the heart. In some situations, a pacemaker spike may not appear clearly on an ECG.

T

Telemetry. The transmission of signals or data from one electronic unit to another by radiowaves or other means (Medtronic, 2003).

Temporary Lead. A pacing lead intended for short-term use, usually with an external pacemaker. Temporary leads may be epicardial or transvenous. A temporary lead does not have a fixation mechanism, allowing it to be easily removed when it is no longer required.

U

Under Sensing. Occurs if the pacemaker fails to sense the P or R wave and thus inappropriately timed impulses may result.

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Approval by Cardiac Team

I have read and approve the *Nursing Practice Committee's 'Nursing Guidelines on Care of the Child with an External Temporary Pacemaker'*.



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