



**Queensland
Government**

Insulin Pump Management Checklist

Facility:

(Affix identification label here)

URN:

Family name:

Given name(s):

Address:

Date of birth:

Sex: M F I

This form to be completed by Medical Officer/Nurse Practitioner during patient with diabetes (PWD) admission to hospital.

Medical Officer/Nurse Practitioner (print name):

Date:

Step 1

	Yes	No	Outcomes
Contact Diabetes Educator (if available in your facility)	<input type="checkbox"/>	<input type="checkbox"/>	
Contact the supervising Medical Officer/Nurse Practitioner/Endocrinologist or facility where the pump was initiated to seek advice	<input type="checkbox"/>	<input type="checkbox"/>	

Step 2: Assessing PWD Safety

Does the PWD have any of the following...	Yes	No	Outcomes										
Altered level of consciousness	<input type="checkbox"/>	<input type="checkbox"/>	If Yes to any question: 1. Turn pump off: Ask the PWD or contact the pump manufacturer's helpline for assistance 2. The PWD should be placed on subcutaneous insulin regimen or an IV insulin infusion while they are in hospital 3. Please refer to the <i>Insulin Subcutaneous Order and Blood Glucose Record</i> for guidance										
Critically ill requiring stabilisation in the Intensive Care Unit	<input type="checkbox"/>	<input type="checkbox"/>											
Serious mental health conditions where the individual is at risk of self-injury or suicide	<input type="checkbox"/>	<input type="checkbox"/>											
Diabetic Ketoacidosis (DKA) or 2 consecutive positive ketone levels	<input type="checkbox"/>	<input type="checkbox"/>											
Impaired judgement	<input type="checkbox"/>	<input type="checkbox"/>											
Any other intercurrent illness affecting their ability to use the insulin pump	<input type="checkbox"/>	<input type="checkbox"/>											
PWD or caregiver refuses or is otherwise unable to participate in care	<input type="checkbox"/>	<input type="checkbox"/>											
Lack of insulin pump consumables, such as infusion sets, cartridges and other required equipment	<input type="checkbox"/>	<input type="checkbox"/>	<table border="1" style="width: 100%;"> <thead> <tr> <th colspan="2">Pump Manufacturer Helplines</th> </tr> </thead> <tbody> <tr> <td>Medtronic</td> <td>1800 777 808</td> </tr> <tr> <td>T-slim</td> <td>1300 851 056</td> </tr> <tr> <td>Ypsomed</td> <td>1800 447 042</td> </tr> <tr> <td>Omnipod</td> <td>1800 954 074</td> </tr> </tbody> </table>	Pump Manufacturer Helplines		Medtronic	1800 777 808	T-slim	1300 851 056	Ypsomed	1800 447 042	Omnipod	1800 954 074
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Medtronic	1800 777 808												
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Lengthy and/or complicated surgery	<input type="checkbox"/>	<input type="checkbox"/>											
Upon assessment, they are unable to use the insulin pump	<input type="checkbox"/>	<input type="checkbox"/>											
Any other medical circumstances deemed unsuitable by the supervising Medical Officer/Nurse Practitioner/Endocrinologist	<input type="checkbox"/>	<input type="checkbox"/>											

Step 3: Assessing the PWD/Parent of Guardians Ability to Manage the Insulin Pump

	Yes	No	Outcomes
Ability to navigate the pump menu or phone app	<input type="checkbox"/>	<input type="checkbox"/>	If No to any question: 1. Turn pump off: Ask the PWD or contact the pump manufacturer's helpline for assistance 2. The PWD should be placed on subcutaneous insulin regimen or an IV insulin infusion while they are in hospital 3. Please refer to the <i>Insulin Subcutaneous Order and Blood Glucose Record</i> for guidance
Ability to adjust basal rates and bolus doses	<input type="checkbox"/>	<input type="checkbox"/>	
Demonstration of managing cannula site and infusion line issues	<input type="checkbox"/>	<input type="checkbox"/>	
Are using a CGM or be willing to do a POC BGL 4 times a day	<input type="checkbox"/>	<input type="checkbox"/>	
Have adequate supplies of infusion sets, reservoirs, spare batteries, charging cable, and rapid acting insulin for the anticipated duration of the admission	<input type="checkbox"/>	<input type="checkbox"/>	
In paediatrics, the parent/guardian is to be responsible for the insulin pump and must stay with the child at all times	<input type="checkbox"/>	<input type="checkbox"/>	
Any other clinical concerns identified by the Diabetes Educator or supervising Medical Officer/Nurse Practitioner/Endocrinologist	<input type="checkbox"/>	<input type="checkbox"/>	

DO NOT WRITE IN THIS BINDING MARGIN

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Step 4: Operations and Procedures

Discuss use of insulin pump in operating theatre and procedure room with...	Yes	No	N/A	Outcomes
Anaesthetist	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	
Surgeon	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	
Physician/Endocrinologist/Nurse Practitioner	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	
Diabetes Educator	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	
PWD	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	

Step 5: Documentation

Before the PWD continues on the insulin pump as an inpatient, the following must be documented in the PWD's chart and the *Blood Glucose Monitoring* form:

1. Place a sticker on the inside of the medical record in the alerts section and on the *Insulin Subcutaneous Order and Blood Glucose Record* stating
 - a. brand name and model of insulin pump;
 - b. manual pump or automated insulin delivery (AID) system (a pump that delivers auto corrections);
 - c. type of insulin used;
 - d. current basal and bolus doses;
 - e. target BGLs;
 - f. insulin : carbohydrate ratio;
 - g. CGM brand/model;
 - h. correction factor (insulin sensitivity).
2. Any changes to the insulin regimen recommended by medical staff during this admission are documented in the medical record and confirmed by the PWD at the time of implementation.
3. Complete *Insulin Subcutaneous Order and Blood Glucose Record* stating that the PWD is self-managing and the frequency of BGLs. Initial BGL frequency is standard if BGLs stable. If BGLs unstable frequency is standard plus 2 hours post-meals and 02:00 hours.

Comments:

Clinician (print name):

Designation:

Signature:

Date:

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