

SUBCUTANEOUS INJECTION SUPPORT PLAN

Identified need for support	Interventions
<input type="checkbox"/> Unable to self-administer injections <input type="checkbox"/> Type 1 Diabetes <input type="checkbox"/> Type 2 Diabetes <input type="checkbox"/> Pain Management <input type="checkbox"/> Palliative Care <input type="checkbox"/> Other: _____ Related for: <input type="checkbox"/> Hypoglycaemia <input type="checkbox"/> Hyperglycaemia <input type="checkbox"/> Pain Goals <input type="checkbox"/> Maintain BGL range <input type="checkbox"/> Prevent and/or manage Hypoglycaemia/ Hyperglycaemia <input type="checkbox"/> Pain Relief Evidenced by <input type="checkbox"/> Medical Directive <input type="checkbox"/> Medical Notes <input type="checkbox"/> Other: _____ Diabetes Association <input type="checkbox"/> Yes <input type="checkbox"/> No Membership No. & details: _____ Palliative Care <input type="checkbox"/> Yes <input type="checkbox"/> No Contact Details: _____	Diabetes Management <input type="checkbox"/> Insulin Dependent <input type="checkbox"/> Oral Medication (refer to medication chart) <input type="checkbox"/> Diet controlled <input type="checkbox"/> Injection (refer to medication chart) <input type="checkbox"/> Medical Directive completed date: _____ Blood Glucose Monitoring <input type="checkbox"/> Yes <input type="checkbox"/> No / BGL frequency _____ Participant's BGL Range (See Medical Directive) <input type="checkbox"/> Consult with Management if BGL <i>outside of accepted range</i> Insulin Administration Order (refer to medication chart) <input type="checkbox"/> Yes – Notify Management of any action taken <input type="checkbox"/> No – Consult with Management before omitting insulin or oral medication Glucagon Administration order <input type="checkbox"/> Yes, refer to medication chart <input type="checkbox"/> No Observe for risks <u>Hypoglycaemia</u> Sweating, dizziness, trembling, anxiety, tingling (hands, feet or tongue), blurred vision, confusion, slurred speech, unconsciousness <u>Hyperglycaemia</u> Excessive urination, excessive thirst, dry mouth, nausea/vomiting, tiredness/fatigue Pain Management/Palliative Care (Refer to Medication Chart) <input type="checkbox"/> Yes – Notify Management of any action taken <input type="checkbox"/> No – Consult with Management before omitting any medication Subcutaneous Injection Sites Observe for:

	<p>Pain, swelling, leakage, bleeding, warmth, infection, redness, tenderness, Haematoma</p> <p><input type="checkbox"/> Site rotation and location (do not inject in bruised areas).</p> <p>Notify health practitioner immediately if any of these complications is observed.</p>
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Other participants specific interventions:

Risk / Response:

- Withdrawal
- Overdose
- Loss of consciousness

Refer to Health Practitioner or transfer to hospital if any of the above risks occur to ensure participants well-being.

Prepared by:		
Position Title:		
Signature:		Date:
Reviewed and Approved by:		
General Practitioner Name:		
General Practitioner Signature:		Date:
Health Professional Name:		
Health Professional Signature:		Date:

Agreement

By signing this Support Plan, I agree that I have been involved in the development of my plan. I agree and consent to the care and interventions of this Subcutaneous Injection Support Plan.

Participant/Representative Name:		
Participant/Representative Signature:		Date:
Company Representative Name:		
Company Representative Signature:		Date:

Communication / Copy of Support Plan

**Copy of Support Plan
given to:**

- Participant
- Health Professional
- Health Practitioner
- Other:

Subcutaneous Injections

Medical Practitioner / Specialist / Consultant Directive

Date:

Diagnosis/Medical History

Specific Care Orders/Treatment Plan

Risks and Complications

Plan Review Frequency

Informed Consent Obtained

Yes No

If NO, state details:

Authorisations

Medical Practitioner Name			
Medical Practitioner Signature		Date	
Client Name			
Client Signature		Date	

Document Control

Version No.	Issue Date	Document Owner
1	08/01/2025	Elizabeth Bradshaw
Version History		
Version No.	Review Date	Revision Description