

PATIENT INFORMATION

Name: _____ Date of Birth: _____
Primary Phone: _____ Secondary Phone: _____
Address: _____ City/State/Zip: _____
Primary Insurance: _____ ID# _____
Secondary Insurance: _____ ID# _____
Contact person (if other than patient): _____

REQUIRED INFORMATION FOR ALL PATIENTS

PLEASE COMPLETE THIS SECTION IN FULL

Date of Last Office Visit: _____ Duration of Need (12 months unless other wise noted): _____
Type 1— IDDM Type 2—Pills, Diet, and/or Insulin Treated HbA1c: _____
E10.9 ___ E11.9 **Currently using a pump?** YES NO
E10.65 ___ E11.65 **Currently on CGM Therapy?** YES NO
Other _____ **Other** _____ **Fasting Hyperglycemia:** _____
Testing Frequency: _____ **Fluctuation of blood glucose values:**
Using insulin shots to control? Yes No **Low:** _____ **High:** _____
Number of injections a day: _____

- | | |
|--|---|
| <input type="checkbox"/> Test Strips | <input type="checkbox"/> Transparent Tape |
| Product Name: _____ | <input type="checkbox"/> Prep Wipes |
| <input type="checkbox"/> Lancets | <input type="checkbox"/> Adhesive Remover |
| Product Name: _____ | <input type="checkbox"/> Infusion Sets |
| <input type="checkbox"/> Lancing Device PRN | <input type="checkbox"/> Reservoirs |
| <input type="checkbox"/> Ketostix | <input type="checkbox"/> Omnipod Insulin Pump PDM |
| <input type="checkbox"/> Alcohol Wipes | <input type="checkbox"/> Omnipod Pods |
| <input type="checkbox"/> Meter Battery PRN | <input type="checkbox"/> Other: _____ |
| <input type="checkbox"/> Control Solution PRN | Other: _____ |
| <input type="checkbox"/> Meter PRN | |

CGM Products

DEXCOM G6

Receiver: Dispense 1: DME Only : 1/5 years
Sensors: Quantity 13 boxes (units): DME Only: 9 units/90 days
Transmitter Dexcom G6 (3 month use): 4/365 Days

LIBRE 2

Receiver: Dispense 1: DME Only: 1/5 years
Sensors: Quantity 26.00 units: 6 units/90 days

USE PER MANUFACTURER RECOMMENDATIONS

My signature below denotes to the best of my knowledge the patient/caregiver is able to follow instructions for controlling diabetes and is able to use the ordered items which are designed for home use. The patient/caregiver has successfully completed training or is scheduled to begin training in the use of supplies or equipment ordered. I am a provider who manages patients with diabetes, insulin pump, or CGM therapy and work closely with a team including nurses, diabetic instructors, and dietitians who are knowledgeable in the use of subcutaneous insulin infusion therapy.

Physician Staff Contact: _____

Provider Name (print): _____ **Provider NPI #** _____

Provider Phone Number: _____ **Provider Fax:** _____

Provider Signature: _____ **Date:** _____

CLINICAL DOCUMENTATION CHECKLIST FOR AIM PLUS MEDICAL SUPPLIES, LLC

The following must be completed and signed by the physician with supporting documentation.

CONTINUOUS GLUCOSE MONITOR PREREQUISITE CRITERIA- Please check all that apply and include supporting documentation.

- Medicaid eligible EPSDT recipients less than 21 years of age and recipients of all ages with Type 1 diabetes and **pregnant**
- Patient is diagnosed with Type 1 diabetes mellitus.
- Patient has been using a blood glucose monitor (BGM) and performing frequent (four or more per day) testing. Supporting documentation must be submitted.
- Patient is insulin-treated with multiple (three or more) daily injections of insulin or a Medicaid-covered continuous subcutaneous insulin infusion (CSII) pump.
- Patient's insulin treatment regimen requires frequent adjustment by the patient and/or caregiver based on BGM or CGM testing results.
- Within six (6) months before ordering the CGM, the treating practitioner has an in-person visit with the patient to evaluate their diabetes control (to include HbA1c) and determined that criteria (1-4) above are met.
- Every six (6) months following the initial prescription of the CGM, the treating practitioner has an in-person visit with the patient to assess adherence to their CGM regimen and diabetes treatment plan.

EXTERNAL AMBULATORY INSULIN INFUSION PUMP (E0784) PREREQUISITE CRITERIA

Please check all that apply and include accompanying documentation.

- Patient must be Medicaid eligible, less than 21 years of age, and EPSDT eligible.
- Patient must have a documented diagnosis of insulin-dependent diabetes mellitus (IDDM, also known as Type 1).
- A board-certified endocrinologist must have evaluated the patient and ordered the insulin pump.
- Patient must have been on a program of multiple daily injections (MDI) of insulin (i.e., at least three injections per day) for at least six months prior to initiation of the insulin infusion pump. Supporting documentation must be submitted.
- Patient has documented frequency of glucose self-testing (i.e., patient "logs") an average of at least four times per day during the three months prior to the insulin pump initiation. The patient must include six consecutive weeks' worth of logs within the three months of the prior authorization request.
- Patient and/or caregiver must be capable, physically and intellectually, of operating the pump. Patient/caregiver must demonstrate ability and commitment to comply with the regimen of pump care, diet, exercise, medications, and glucose testing at least four times a day. Supporting documentation must be submitted.
- Education on insulin pump must have been conducted before prior authorization request. The patient (caregiver if a child) and educator must sign to document their understanding.
- Documentation of active and past recipient compliance with medications, diet, appointments, and other treatment recommendations must be provided.

ADDITIONAL CRITERIA- The patient must also meet one or more of the following, supported by documentation:

- Two elevated glycosylated hemoglobin levels (HbA1c>7.0%) within 120 days while on multiple daily injections of insulin.
- History of severe glycemic excursions (commonly associated with brittle diabetes, hypoglycemic unawareness, nocturnal hypoglycemia, extreme insulin sensitivity, and/or very low insulin requirements).
- Widely fluctuating blood glucose levels before mealtime (i.e., preprandial blood glucose level consistently exceeds 140 mg/dL).
- Dawn phenomenon with fasting blood sugars frequently exceeding 200 mg/dL.

I certify that this treatment is indicated and necessary and meets the guidelines for use as outlined by the **Mississippi Division of Medicaid**. I will be supervising the patient's treatment. Required supporting documentation from the patient's medical record is attached.

Prescribing Practitioner Signature (Required)

Date

Stamps/copies of the physician's signature will not be accepted.