Dear Physician,

The following information is intended to provide you with guidance on Medicare’s coverage and documentation requirements for blood glucose monitors (BGMs) and testing supplies.

**Coverage**

Glucose monitors and related supplies are covered for patients with diabetes if they or their caregivers can be trained to use the prescribed device appropriately.

The quantity of test strips and lancets that are covered, if the basic criterion above is met, is shown below.

<table>
<thead>
<tr>
<th>Treatment regimen</th>
<th>Basic coverage</th>
</tr>
</thead>
<tbody>
<tr>
<td>Test strips and lancets</td>
<td></td>
</tr>
<tr>
<td>Insulin treated</td>
<td>300 per 3 months</td>
</tr>
<tr>
<td>Non-insulin treated</td>
<td>100 per 3 months</td>
</tr>
</tbody>
</table>

- Additional quantities of test strips can be covered if they are documented to be medically necessary – as outlined below.
- Coverage is also provided for a lancing device, calibration solution, and replacement batteries.

**Medical Necessity Documentation**

CMS expects that physician records will reflect the care provided to the patient including evidence of the medical necessity for the prescribed frequency of testing. You are not required to fill out additional forms from suppliers or to provide additional information to suppliers unless specifically requested by the DME MAC.

It is critical that the patient’s medical record demonstrates the medical necessity for glucose testing supplies, which includes:

- Diagnosis
- Treatment regimen (insulin treated versus non-insulin treated)
- Basic coverage criteria for the BGM and any related supplies; and,
- Evidence of the patient’s use at this frequency

For quantities of supplies that exceed the limits specified in the local coverage determination (LCD), there must be:

- Documentation by the physician in the patient’s medical record of the necessity for the higher frequency of testing, which may include some of the following elements (not all-inclusive):
  - Names, dosages, and frequency of administration of medications used to treat the diabetes;
  - Frequency and severity of symptoms related to hyperglycemia and/or hypoglycemia;
  - Review of patient-maintained log of glucose testing values:
    - Logs of self-testing values including the date, time, and results;
    - Information about medication dosage adjustments related to the results is also helpful;
Changes in the patient’s treatment regimen as a result of glucose testing results review;
- Dosage adjustments that the patient should make on their own based on self-testing results;
- Laboratory tests indicating level of glycemic control (e.g., hemoglobin A1C);
- Other therapeutic interventions and results.

Not every patient’s medical record will contain all of these elements; however, there must be enough information in the patient’s medical record to support the medical necessity for the quantity of item(s) ordered and dispensed.

**ORDERS**

All durable medical equipment, prosthetics, orthotics and supplies (DMEPOS) require a written order/prescription which must be received by the supplier prior to billing Medicare. Someone other than the treating practitioner may complete the standard written order (SWO) for the item unless statute, manual instructions, the contractor's LCD or policy articles specify otherwise; however, the prescribing practitioner must review the content and sign the document.

**Note:** A new order for diabetic testing supplies is required only if:

- there is a change in the frequency of testing/quantity to be dispensed; or
- when replacing a BGM; or
- there is a change in supplier.

**Note:** If the supplier provides you with a prepared SWO for your signature, you should inspect this document carefully. Suppliers must not add unrelated items to the SWO, whether requested by the patient or not, in the absence of your explicit approval.


Sincerely,

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