Continuous Glucose Monitors (CGM)

The treating clinician must complete the following items:

☐ Conduct and document a Face-to-Face Evaluation (FTF)
☐ Complete a 5 Element Order (5EO)
☐ Medical record documentation requirements (see below)

All Related CGM Accessories and Supplies

The treating clinician must complete the following items:

☐ Complete a Dispensing Order
☐ Sign and date a detailed written order (DWO)
☐ Medical record documentation requirements (see below)

Medical Record requirements for CGM

Medical records should indicate:

☐ Beneficiary has diabetes, and
☐ Beneficiary has been using a blood glucose monitor (BGM) and testing four or more times a day; and
☐ Beneficiary is insulin treated with multiple (3 or more) daily injections of insulin, or a Medicare covered continuous subcutaneous infusion pump (CSII), and
☐ Beneficiary's insulin treating regime requires frequent adjustments by the beneficiary on the basis of BGM or CGM testing results, and
☐ An in-person diabetic evaluation within 6-months of ordering the CGM to determine and document that the above criteria are met; and
☐ A subsequent in-person evaluation every 6 months to verify and document daily use of CGM and assess adherence to the diabetes treatment plan.